MOLECULAR EVOLUTION AND PHYLOGEOGRAPHY OF THE EASTERN MEDITERRANEAN WATER FROG (*PELOPHYLAX*) COMPLEX

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ABSTRACT

MOLECULAR EVOLUTION AND PHYLOGEOGRAPHY OF THE EASTERN MEDITERRANEAN WATER FROG (*PELOPHYLAX*) COMPLEX

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Water frogs (genus *Pelophylax*) in the eastern Mediterranean region represent a genetically and phylogenetically diverse group. Their dependence on freshwater habitats makes them highly sensitive to geological and climatic changes. Thus they are an ideal group to study the effect of past geological processes on molecular evolution of protein-coding and non-coding genes on the genome and on patterns of phylogeography in the eastern Mediterranean region.

In the first study, the complete sequence of water frog Pelophylax lessonae' serum albumin (SA) gene including all exons, all introns, upstream sequence of the 5' UTR and downstream sequence of the 3' UTR were reported. By aligning P. lessonae' albumin gene with albumin genes of closely and distantly related amphibian species, potential regulatory elements and functional motifs in the 5' UTR (TATA box, HP element, ADEI, ADEII) and 3' UTR (PLE, USE1 and 2, PAS, DSE), and in the exon-intron junctions were documented. Comparison of exon and intron lengths of the water frog with other vertebrate species revealed a strong length conservation in protein coding albumin exons among distinct vertebrate species, but high variation in intron lenghts. Moreover, the search of the entire scaffold for retroelements indicated several distinct classes of truncated retroelement families upstream of the 5' UTR, downstream of the 3' UTR and within individual albumin introns could be one of the important drivers of the albumin gene evolution. Finally, phylogenetic analysis based on nucleotide sequences of protein-coding albumin exons supported previous mitochondrial DNA (mtDNA) and the first intron of the SA gene findings which were the sister group relationship of *P. lessonae* and *P. bergeri*, a close relationships of *P.* ridibundus and P. cf. bedriagae from Anatolia. Moreover, P. epeiroticus is closely related to P. ridibundus/P. cf. bedriagae. In contrast to former mtDNA findings, the SA exon data revealed that P. shqipericus represent a distinct lineage, which was supported by previous protein electrophoretic investigations.

In the second study, phylogeographic patterns between mitochondrial (ND2 and ND3) and nuclear markers (SAI-1+*Rana*CR1) for water frog complex of the eastern Mediterranean region were compared. On the basis of analysis using extensive data set from mtDNA and nuDNA markers, these markers revealed discordances in the tree topology, the number of genetic stocks, the level of genetic differentiation, the times of divergence and the patterns of geographic distribution among distinct water frog lineages. These discordances between two markers were discussed by distinct processes such as incomplete lineage sorting, retention of ancestral polymorphism, introgression after secondary contact

reacting with sex-biased or adaptive processes, and distinct selective constraints on genes. The major water frog lineages in the eastern Mediterranean region, *P. cretensis*, *P. cypriensis*, the Levant (*P. bedriagae*), Cilician groups, Central Asia groups, and Anatolian main allele groups (MAGs) could be affected in distinct ways from these processes that were discussed in this study.

Keywords: molecular evolution, the serum albumin gene, *Pelophylax lessonae*, retroelements, discordant phylogeography, eastern Mediterranean, water frog (*Pelophylax*), mtDNA, ND2, ND3, nuDNA, *SAI-1+RanaCR1*.

ÖΖ

DOĞU AKDENİZ SU KURBAĞALARI (*PELOPYLAX*) KOMPLEKSİNİN MOLEKÜLER EVRİMİ VE FİLOCOĞRAFYASI

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Doğu Akdeniz'deki su kurbağaları (genus *Pelophylax*) hem genetik hemde filogenetik olarak çeşitlilik gösteren bir grubu oluşturmaktadır. Bu hayvanların tatlı su ortamlarına olan bağımlılıkları onları hem jeolojik hem de iklim değişikliklerinin etkisine oldukça hassas duruma bir getirmektedir. Bu sebeplerden dolayı bu grup geçmişteki jeolojik olayların genomdaki protein kodlayan ve kodlamayan genlerin moleküler evrimine ve Doğu Akdeniz'deki filocoğrafik oluşumlarına olan etkileri çalışmak için ideal bir gruptur.

İlk çalışmada su kurbağası *Pelophylax lessonae* türünün serum albumin (SA) geninin tüm exonları, intronları, 5' UTR'ın yukarı, ve 3' UTR'ın aşağı kısmı olmak

üzere tüm gen sekansı bildirilmiştir. P. lessoanae albumin geni hem yakın hemde uzak diğer amfibi türleri ile hizalanarak genin 5' UTR (TATA box, HP element, CAT box ADEI, ADEII) ve 3' UTR (PLE, USE1 ve 2, PAS, DSE) bölgeleri ve exon-intronların birleşme noktalarındaki muhtemel düzenleyici ve fonksiyonel motifler belirlenmiştir. Su kurbağasının exon ve intron uzunluklarının diğer omurgalı türleri ile karşılaştırılması protein kodlayan albumin geninin exon uzunluklarının oldukça güçlü şekilde korunduğu buna karşın intron uzunluklarının oldukça değişkenlik gösterdiğini ortaya koymuştur. Bundan başka tüm scaffoltun retroelementler için taranması sonucu albumin geninin 5' UTR'ın yukarısında, 3' UTR'ın aşağısında ve intronların içinde çok sayıda farklı ailelere ait parçalanmış retroelementler olduğunu göstermiş ve bunların albumin geninin evrimindeki önemli faktörlerden biri olduğunu göstermektedir. Son olarak serum albuminin protein kodlayan exonlarının nükleotid sekansına dayalı olarak oluşturulan filogenetik analiz sonuçları daha önceki mitokondriyal DNA (mtDNA) ve SA'nin birinci intron sonuçları ile desteklenmektedir, buna göre bir tarafta P. lessonae ve P. bergeri kardeş grup oluştururken, diğer tarafta P. ridibundus ve Anadolu'dan P. cf. bedriagae ise diğer yakın grubu oluşturmakta ve P. epeiroticus'un son gruba yakın olduğu görülmektedir. Daha önceki mtDNA bulgularına karşın, SA exon verileri P. shqipericus'un farklı bir soyu gösterdiği ve bu bulgunun daha önceki protein elekroforetik araştırmaları tarafından desteklenmektedir.

İkinci çalışmada, Doğu Akdeniz Bölgesi'ndeki su kurbağalarının mitokondriyal (ND2 ve ND3) ve nükleer marker (SAI-1+RanaCR1)larının filocoğrafik oluşumları karşılaştırılmıştır. Mitokondriyal ve nuklear DNA markerlarının yoğun verilerinin analizlerine dayalı olarak elde edilen sonuçlar bu iki marker arasında ağaç topolojisinde, genetik stokların sayısında, genetik faklılaşmanın düzeyinde, ayrılma zamanlarında ve farklı su kurbağası soylarının coğrafik dağılımlarında uyumsuzluklar olduğunu göstermiştir. Bu iki marker arasındaki uyumsuzluklar olduğunu göstermiştir. Bu iki marker arasındaki dağılımlarında farklı süreçler ile açıklanabilir ki bunlar genlerin soylardaki dağılımının eşit olmaması, atasal polimorfizmin tutulması, ikinci bağlantıdan sonra meydana gelen gen akışı ve bunun eşeye bağlı ve adaptive süreçler ile birlikte hareket etmesi ve

genler üzerindeki farklı selektif baskılar olabilir. Doğu Akdeniz'deki başlıca su kurbağası soyları olan *P. cretensis, P. cypriensis,* the Levant (*P. bedriagae*), Kilikya grupları, Orta Asya grupları, ve Anadolu MAGs bu süreçlerden farklı şekilde etkilendiği düşünülmekte olup bu çalışmada bunlar tartışılmıştır.

Anahtar kelimeler: moleküler evrim, serum albumin geni, *Pelophylax lessonae*, retroelements, uyumsuz filocoğrafya, Doğu Akdeniz, su kurbağası (*Pelophylax*), mtDNA, ND2, ND3, nuDNA, SAI-1+*Rana*CR1

To My Wonderful Family and My Partner, Ozan

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TABLE OF CONTENTS

ABSTRACTV
ÖZVİİİ
ACKNOWLEDGEMENTS
TABLE OF CONTENTSXV
LIST OF TABLESXVİİİ
LIST OF FIGURESXX
LIST OF ABBREVIATIONS
CHAPTERS
1. GENERAL INTRODUCTION
1.1. Analysis of Molecular Evolution
1.2. Mechanisms of Genome Evolution
1.2.1. Gene and Genome Duplications
1.2.2. Transposable Elements
1.2.3. Exon Shuffling
1.2.4. Horizontal Gene Transfer
1.3. Molecular Phylogeography
1.3.1. Phylogenetics and Phylogeography of the Eastern Mediterranean Water
Frogs (genus <i>Pelophylax</i>)
1.4. The Present Study 12
1.5. References
2. MANUSCRIPT 1: MOLECULAR EVOLUTION AND
CHARACTERIZATION OF THE SERUM ALBUMIN GENE OF THE POOL
FROG, PELOPHYLAX LESSONAE
2.1. Introduction
2.2. Material and Methods
2.2.1. Sources of Albumin Sequences

2.2.2. Alignment and Motif Searches	.27
2.2.3. Search For Transposable Elements and Repetitive Sequences	.27
2.2.4. Phylogenetic Analyses Based on Protein Coding Exon Sequences	.28
2.3. Results and Discussion	.29
2.3.1. Structure of the Albumin Gene	.29
2.3.1.1. The 5' Region of the Water Frog Albumin Gene	.30
2.3.1.2. The 3' Region of the Water Frog Albumin Gene	.34
2.3.1.3. 5' and 3' Splice Junctions of the Water Frog's Albumin Gene	.38
2.3.1.4. Exons and Introns of the Pool Frog's Albumin Gene	.42
2.3.2. Drivers of Serum Albumin Gene Evolution	43
2.3.2.1. DNA Transposons	48
2.3.2.1.1. Tc1Mar-Tc1 Family	.48
2.3.2.1.1.1. PL Tc1-1Ory Element	48
2.3.2.1.1.2. PL Tc1-10Xt Element	51
2.3.2.1.1.3. PL Tc1-RT Element	54
2.3.2.1.1.4. PL Tc1-PP Element	57
2.3.2.1.1.5. PL Tc1-DR1 Element	60
2.3.2.1.1.6. PL Tc1-FR3 Element	.63
2.3.2.1.1.7 Structural Characteristics of ITR and Transposase Genes of	of
P. lessonae consensus TLEs	65
2.3.2.1.2. hAT/Charlie Family	66
2.3.2.1.2.1. PL hAT-2-TS Element	.67
2.3.2.1.2.2. PL hAT-2-ET Element	71
2.3.2.1.2.3. PL hAT-2-OG Element	75
2.3.3. Sequence Evolution and Phylogenetic Implications	78
2.4. References	81
2.5. Appendices	.92
3. MANUSCRIPT 2: DISCORDANCE BETWEEN MITOCHONDRIAL AN	ND
NUCLEAR PHYLOGEOGRAPHY IN WATER FROG (PELOPHYLA	4X)
COMPLEX OF THE EASTERN MEDITERRANEAN REGION	49
3.1. Introduction	49

3.2. Material and Methods
3.2.1. Field Trips, Locality Selection and Sampling 153
3.2.2. PCR and Sequencing of Mitochondrial and Nuclear Genes 154
3.2.3. Sequence Alignment and Grouping 155
3.2.4. Phylogenetic Analysis
3.3. Results
3.3.1. Molecular Diversity
3.3.2. Phylogenetic Relationships, Genetic Divergence, and Geographical
Pattern of Mitochondrial Haplogroups159
3.3.3. Phylogenetic Relationships, Genetic Divergence, Geographical Pattern of
Nuclear Allele Groups and Their Relationships with Mitochondrial
Haplogroups
3.3.4. Comparison of mtDNA and nuDNA Tree Topologies and Estimation of
Times of Divergence 174
3.4. Discussion
3.4.1. Potential Mechanisms Causing Phylogeographic Discordance among
mtDNA and nuDNA markers 179
3.4.1.1. P. cypriensis versus P. cretensis
3.4.1.1.1. Isolation of Cyprus
3.4.1.1.2 Isolation of Crete
3.4.1.1.3 Possible Explanations for the Discrepancy
3.4.1.2. The Levant (<i>P. bedriagae</i>)
3.4.1.3. Cilician versus Central Asia Group 186
3.4.1.4. Discordance within the Cilician Groups
3.4.1.5. Ridibundus versus Anatolia groups 187
3.4.1.6. Discordance Patterns within the Anatolia MAG 190
3.4.2. Systematic Implications
3.5. References
3.6. Appendices
4. CONCLUSIONS
CURRICULUM VITAE

LIST OF TABLES

TABLES

Table 2.1. Classification of truncated repetetive elements obtained from Repeat
Masker (RM) and TRF programs' searches45
Table 2.2. Main features of P. lessonae consensus TLEs
Table 2.3. Comparison of SA exon and intron lengths among distinct vertebrate
species92
Table 2.4. Alignment of <i>P. lessonae</i> scaffold 10.7 with the serum albumin cDNA
of P. shqipericus
Table 2.5. Calculation of the number of nucleotide differences and uncorrected p
distance among western Palearctic water frog species on the basis of nucleotide
sequence of the serum albumin protein coding exons147
Table 2.6. Calculation of the number of aa differences and uncorrected p distance
among western Palearctic water frog species on the basis of aa sequence of the
serum albumin protein coding exons
Table 3.1. Estimates of divergence times for mtDNA and nuDNA markers178
Table3.2.Locality informationformitochondrialND2andND3
haplotypes
Table3.3.LocalityinformationofnuclearmarkerSAI-1+RanaCR1
alleles
Table 3.4. Calculation of genetic distance among mitohondrial subgroups, main
haplogroups, and outgroup
Table 3.5. Calculation of genetic distance among nuclear subgroups, main allele
groups and outgroup species

Table	3.6.	Group	specific	unique	molecular	variations	for	nuclear	SAI-
1+Rand	aCR1	gene					•••••		333

LIST OF FIGURES

FIGURES

Figure 1.1. Geographic distribution of water frogs' main haplogroups (MHGs)
and subgroups (a-d) of water frogs in Anatolia and neighbouring regions11
Figure 2.1. Schematic structure of pool frog'serum albumin gene from the Cap
site of the first exon to the first polyadenylation site of the last exon
Figure 2.2. 5' regions of the serum albumin gene P. lessonae, P. ridibundus, P.
chensinensis, P. kukunoris, X. laevis, and S. tropicalis, starting with the translation
initiation codon (Met) up to position -340
Figure 2.3. Sequence alignment of the 3' UTR of P. lessonae, P. ridibundus, R.
catesbeiana, R.chensinensis, R. kukunoris, X. laevis and S. tropicalis albumin
genes
Figure 2.4. Comparison of 5' and 3' splice positions among P. lessonae, S.
tropicalis and X. laevis
Figure 2.5. Comparison of serum albumin exon lengths among P. lessonae, S.
tropicalis, X. laevis, G. gallus, R. norvegicus, M. musculus and human
Figure 2.6. Comparison of serum albumin intron lengths among P. lessonae, S.
tropicalis, X. laevis, G. gallus, R. norvegicus, M. musculus and human
Figure 2.7. Structure of the PL Tc1-1ORY element and its consensus
sequence
Figure 2.8. Structure of PL Tc1-10Xt element and its consensus sequence53
Figure 2.9. Structure of PL Tc1-RT element and its consensus sequence
Figure 2.10. Structure of PL Tc1-PP element and its consensus sequence
Figure 2.11. Structure of PL Tc1-DR1 element and its consensus sequence61
Figure 2.12. Structure of PL Tc1-FR3 element and its consensus sequence64

Figure 2.13. Structure of the PL hAT-2-TS element and its consensus sequence.

Figure 2.14. Structure of the PL hAT-2-ET element and its consensus sequence.

Figure 2.15. Structure of the PL hAT-2-OG element and its consensus sequence.

Figure 3.4. Maximum clade credibility tree constructed using Bayesian phylogenetic analysis of nuclear SAI-1 + RanaCR1 alleles from western Palearctic and eastern taxa of Eastern Palearctic water frogs (genus *Pelophylax*)..

LIST OF ABBREVIATIONS

WGD	Whole genome duplication
SD	Segmental duplication
TE	Transposable elements
HGT	Horizontal gene transfer
MHG	Main haplogroups
UTR	Untranslated terminal repeats
mtDNA	Mitochondrial DNA
nuDNA	Nuclear DNA
aa	Amino acids
kd	Kilodalton
bp	Base pairs
SA	Serum albumin
PL	Pelophylax lessonae
ORF	Open reading frame
ITR	Inverted terminal repeat
HP	Hepatocyte-specific promoter element
ADE	Amphibian downstream element
PLE	Poly(A)-limiting element
USE	U-rich upstream element
PAS	Polyadenylation site
DSE	Downstream element
CPSF	Cleavage and polyadenylation specifity factor
CstF	Cleavage stimulatory factor
IRE	Iron responsive element
LTR	Long terminal repeat

TLE	Tc1-like elements
NLS	Nuclear localization signal
DR	Direct repeat
HLE	hAT-like element
MSC	Messinian salinity crisis
SAI	Serum albumin intron
Myr	Million years
Ma	Million years ago
MAG	Main allele group
MPR	Mid-Pleistocene revolution

CHAPTER 1

GENERAL INTRODUCTION

1.1. Analysis of Molecular Evolution

Molecular evolutionary research covers a broad range of topics starting from gene and genome evolution to molecular phylogenetics (Li, 1997; Page and Holmes, 1998; Graur and Li, 2000; Nei and Kumar, 2000; Yang, 2006). Based on genetic information stored in molecules (DNA, RNA, proteins), evolutionary patterns can be analyzed and phylogenetic relationships among species and supraspecific taxa can be inferred. As can be demonstrated by an ever-growing amount of molecular data, molecular approaches which supplement traditional non-molecular methods are a pivotal part of the toolkit of evolutionary biologists and systematists. For Amphibia, internet sequence databases (GenBank, EMBL) currently contain about 3.326.318 entries of DNA and RNA sequences. DNA sequences in particular, because they bear the code of life, provide the most detailed information possible for any organism - the instructions for how each working part should be assembled and operate (Page and Holmes 1998). Progress in technological development, in particular the simplification of molecular techniques and the reduction in expense, will almost certainly lead to a further increase of molecular studies. Thus, molecular evolution is a constantly changing and evolving field because of the steadily growing number of sequenced genomes, both prokaryotic and eukaryotic, available to the scientific community and the public at large.

1.2. Mechanisms of Genome Evolution

1.2.1. Gene and Genome Duplications

Genome evolution occurs at distinct levels; beside small scale changes such as single base substitutions, insertions and deletions, gene and whole genome duplications (WGD) are possible. All of these mutation events provide raw genetic material on which natural selection can act upon.

Genome duplications have been playing an important role in the vertebrate genome evolution since it was suggested (Ohno, 1970) that the early vertebrates have undergone two rounds of whole genome duplication (2 R WGD) about 600 Ma which was later documented by analysis of *Hox* genes and *Hox* gene clusters (Holland et al., 1994; 1997). 3R of WGDs were proposed to occur in the stem lineage of ray-finned fish (Meyer and Van de Peer, 2005). These global events could extremely increase the number of gene copies and produce several distinct paralogs which provide novel raw material for evolutionary processes leading to genomic innovations (Zhang 2003); the globin gene family is an impressive example (reviewed by Storz et al. 2013).

These gene copies could remain or be lost differently according to their functions throughout evolution of vertebrate lineages (Blomme et al., 2006). Therefore the fate of paralogs can be different: i) both gene copies can remain and obtain new functions. (neofunctionalization); ii) they can acquire degenerative complementary mutations thus both are necessary to perform the original function of the ancestral gene (subfunctionalization) (Force et al., 1999); iii) they can maintain the same function by gene conservation through concerted evolution (Li, 1997); iv) they can evolve to pseudogenes, mostly functionless, which are not deleterious and not eliminated by selection (Zhang, 2003), and provide the organism genetic diversity (Gu, 2003). In the last case, it increases tolerance to

changing environmental conditions decreasing the probability of extinction (Crow and Wagner, 2006).

Rather than WGD, gene duplications can also arise from segmental duplications (SDs), unequal crossing over and transpositions events. SDs are characterized by low copy number and can change from few base pairs to many megabases. They have played an important role in the evolution of the human and primate genomes, for example many rapidly evolving exons are found within SDs (Lorente-Gardos et al., 2013). SDs can be arranged as adjacent duplications, interspersed intrachromosomally or interchromosomally (Ramos and Ferrier, 2012). It was suggested that distinct DNA-based mechanisms are responsible for these arrangements. For instance, non-homologous end-joining (NHEJ) more likely results in adjacent duplications (Meisel, 2009) while non-allelic homologous recombination (NAHR) between repetitive elements on the genome accounts for interspersed duplications (Fiston-Lavier et al., 2007). In the second mechanism, unequal crossing over produces linked duplicated genes in the chromosome which can include many genes, a single gene or the part of a gene. Retrotransposition mechanism, in contrast, generates gene copies randomly inserted into the genome and not linked to the original copy. Because mRNA is reverse transcribed into cDNA which is inserted into the genome, many regulatory elements such as promoter are not transcribed during this process. Thus, new gene copies generally become pseudogenes which do not contain regulatory sequences necessary for transcription (Zhang, 2003).

1.2.2. Transposable Elements

Transposable elements (TEs) are important drivers of genome evolution and are intrinsic components of regulatory networks (Feschotte 2008). TE-derived sequences such as promoters (Van de Lagemaat et al., 2003; Marino-Ramirez et al., 2005; Cohen et al., 2009; Conley et al., 2008; Jern and Coffin, 2008), polyadenylation signals and termination sites (Roy-Engel et al., 2005; Lee et al.,

2008; Conley and Jordan, 2012), and smRNAs (Smalheiser and Torvik, 2005) are involved in regulation of gene expression at both the transcriptional and post-transcriptional level (Feschotte, 2008; Medstrand et al., 2005; Rebollo et al., 2012).

Transposable elements are classified into two groups according to their structure and mode of transposition: i) Retrotransposons (class I elements) which mobilize via reverse-transcription using an RNA intermediate, and ii) DNA transposons (class II elements) which replicate without an RNA intermediate, either by a cutand-paste mechanism (DNA transposons), by rolling circle DNA replication (helitrons), or by so far unknown mechanisms (politrons/mavericks) (e.g. Levin and Moran 2011; Rebollo et al., 2012). Among the Class I elements two major subclasses are recognized: (1) retroelements (REs) with long terminal repeats (LTRs) and (2) elements without LTRs (non-LTR REs) (Rebollo et al., 2012; Deininger and Batzer, 2002).

The copy number of TEs is significantly related with the genome size (Kidwell, 2002). For instance, TEs make up only small parts of the fruit fly (*Drosophila melanogaster*) and the worm (*Caenorhabditis elegans*) genomes (15-22% and 12%, respectively). In contrast, half or even more than half of the vertebrate genomes is dominated by distinct classes of TEs. For example, 47% of the genome of giant salamander, the largest genome in vertebrates (14 Gb to 74 Gb), is dominated by TEs, particulary by LTR retrotransposons (Sun et al., 2011). In human, 45% of the genome contains TEs (Chenais et al., 2012).

In addition to their impact on genome size variation, TEs can influence the dynamics of the genome in different ways. Activity of TEs may result in both destructive and constructive alterations (Kidwell and Lisch, 2000). Deleterious effects occur if TEs insert into coding sequences which can lead to premature termination of the peptide sequence and thus, disrupt gene function (Chenais et al., 2012). Many TE derived sequences, however, can act as cis-regulatory

elements at promoter and enhancer regions (Jordan et al., 2003; Cohen et al., 2009; Teng et al., 2011); they can also function as an antisense promoter for genes that are located at 5' of the complete element (Kazazian, 2004). Moreover, TEs may evolve to new protein coding genes in a process called exonization (Sela et al., 2010). Alzohairy et al. (2013) reviewed how TEs became parts of functional genes in eukaryotic genomes in the process of "molecular domestication". As a result of a fusion between TEs and host genes chimeric proteins can originate as demonstrated for the SETMAR gene (Roman et al., 2007). Recently, the availability of transcriptome data from the European water frog species *P. lessonae* and its comparison to transcriptome data obtained from *Silurana* revealed that TEs, particularly LTR retroelements, may play important roles during embryogenesis (Grau et al., 2014).

Moreover, TEs can carry an intragenic poly(A) signal such as human L1 and Alu elements (Roy-Engel et al., 2005) which results in premature cleavage of the transcripts. Besides their promoter and intragenic poly(A) functions, these elements can cause 5' and 3' transduction during insertion because they can carry sequences from a promoter upstream of the complete element and downstream of the poly(A) signal (Kazazian, 2004). Lastly, they can contain some characteristics which may result in microRNAs, as known from miniature inverted repeat transposable elements (MITEs) (Feschotte, 2008). It was also suggested that precursor of mammalian microRNAs are ancient MIR (SINE) and L2 (LINE) elements (Smalheiser and Torvik, 2005). At chromosomal level, TEs are potential sources for several types of chromosomal rearrangements. For example, the human genome is rich in L1 and Alu elements which make quite likely non-allelic homologous recombination; mispairing between these non-LTR elements can result in deletions, segmental duplications, inversions or translocations (Konkel and Batzer, 2010).

1.2.3. Exon Shuffling

Exon shuffling is an important molecular mechanism for the formation of new genes and also for genome and proteome evolution (Liu and Grigoriev, 2004) in complex eukaryotic genomes because increase in genome size is correlated with increase in intron length and the number of repetitive elements which make more likely potential of exon shuffling by intronic recombination (Keren et al., 2010). Exon shuffling gives rise to a new exon combination produced by recombination between intronic sequences of unrelated genes, which assorts exonic sequences randomly (Gilbert, 1978) or exon duplication occurs within the same gene (Kondrashov and Koonin, 2001; Long et al., 2003).

Non-coding intronic sequences are long and harbour many types of repetitive elements which make them recombination hotspot regions via mismatch or nonhomologous recombination (Patthy, 1999; Kolkman and Stemmer, 2001). Exon shuffling can be symmetric and asymmetric according to intron phase, which indicate the position of a given intron within a codon. Phase 0 introns are located between two codons, phase 1 introns lie between the first and second nucleotide of the codon, and phase 2 intron is found between the second and third nucleotide of the codon (França et al., 2012). Thus, symmetric exons have the same phase in their flanking introns including three groups 0-0, 1-1, 2-2 while asymmetric exons are 0-1, 0-2, 1-0, 1-2, 2-0 and 2-1. It was suggested that only symmetric exons can be subjected to exon shuffling, tandemly duplicated or deleted with preserving the reading frame (Kolkman and Stemmer, 2001; França et al., 2012).

Exon shuffling significantly contributed to metazoan evolution and to the development of multidomain proteins. These proteins function in tissue remodelling processes, cell-to-cell and cell-to-matrix interactions, cell-to-cell communication, that is, are mainly essential for multicellularity of metazoa (Patthy, 1999). It was documented that ancient domains are flanked by phase 0

introns and they have a tendency of distribution to take place in the central part of proteins. In contrast, modern domains are frequently flanked by phase 1 introns, and they are mainly found in their carboxyl and amino ends (Vibranovski et al., 2005; 2006; França et al., 2012). In addition, genes reflecting high degree exon shuffling events could provide novel sources for protein-protein interactions; both interactions among multiple types of other domains and interactions with other copies of themselves during metazoan evolution (Cancherini et al., 2010).

1.2.4. Horizontal Gene Transfer

Horizontal gene transfer (HGT), also referred as lateral gene transfer, is a nonsexual movement of genetic material between closely or distantly related genomes in contrast to vertical transmission of genetic information from parents to offspring (Keeling and Palmer, 2008). The transfer of genes can occur via distinct ways such as bacteria, viruses, transposons, other vectors and direct contacts or initial hybridization (Syvanen, 2012). Particularly, early developmental stages in multicellular eukaryotes provide a weak link for entry of foreign genes into new genomes (Huang, 2013). HGT is important in bacterial genome evolution (Frost et al., 2005) while its role in eukaryotes is greatly variable because several types of transfers are possible. In the simple transfer, genes from bacteria to eukaryotes move and duplicate or replace the homologous copies. In the serial transfer, transfer from bacteria to eukaryotes follows the second eukaryotic lineage which results in complex relationships. In the novel gene transfer, a new function is provided to the eukaryotic lineage (Keeling and Palmer, 2008; Rajarapu, 2014). It can show a patchy distribution among eukaryotic lineages because of differential loss or gain. Therefore, its role in genome evolution is variable. HGT of transposable elements among eukaryotic lineages is one of the best example how they impact on genome evolution from a neutral variation to genome-wide innovation which can result in speciation or adaptation (Schaack et al., 2010). Moreover, transferred genes in distinct eukaryote species take part in parasitism, disease resistance, new metabolic pathways, nutrient and energy metabolism, and many other new functions, supporting evidence in the adaptative evolution of eukaryotes (Haegeman et al., 2011; Fitzpatrick, 2012; Zhu et al., 2011).

There are three main approaches to predict HGT in eukaryotes. Initially, codonbased approach is based on comparison of G+C content and codon usage of recipient with donor organism after HGT because organisms have specific base frequency and codon usage characteristics (Becq et al., 2010; Azad and Lawrence, 2011). This approach is suitable for recent HGT rather than ancient one (Lawrence and Ohman, 1997). Secondly, the BLAST-based approach is to use database search to find out the most similar sequence to a query sequence of a gene (Keeling and Palmer, 2008; Whitaker et al., 2009). Lastly, phylogenetic approach is based on conflict between a gene tree and species phylogeny. Any incongruence between two phylogenetic trees is an indicator of HGT, which requires well sampling of homologous sequences, their multiple alignment, phylogenetic tree construction and topology comparison. Each approach has distinct drawbacks (Keeling and Palmer, 2008; Whitaker et al., 2009; Schönknecht et al., 2013, Rajarapu, 2014)

1.3. Molecular Phylogeography

Phylogeography studies how historical processes shaped the current distribution of species by utilizing phylogenetic analysis of genetic data and their geographic distribution (Avise et al., 1987, reviewed later by Hickerson et al., 2010). It answers questions about a single species such as species delimitation (Knowles and Carstens, 2007), past hybridization, introgression, hybrid zones (Hewitt, 2001) and multiple species named as comparative phylogeography which studies how past common geological and climatic events have affected the contemporary species diversity by looking at overlapping historical patterns of genetic divergence and gene flow in time and space (Hickerson et al., 2010). Many early phylogenetic studies were based on mainly mitochondrial DNA (mtDNA), while recently there is a considerable transition to multilocus markers. Although distinct genes have distinct evolutionary rates which could give rise to distinct phylogenetic signals, application of data partition methods on concatenated datasets can help to resolve the problem (Nylander et al., 2004). Brito and Edwards (2009) suggested a large number of multiple and independent loci can overcome the effect of coalescent variation and HGT events and resolve incongrunce between the gene tree and the species tree.

1.3.1. Phylogenetics and Phylogeography of the Eastern Mediterranean Water Frogs (genus *Pelophylax*)

The discovery of the hybrid nature of the common European edible frog, *Pelophylax esculentus*, and its taxonomic separation from one of its parental species, the pool frog, *P. lessonae* (Berger, 1967; 1968) has intensified the research on western Palearctic water frogs resulting in acquisition of large genetic data sets on the basis of which diverse questions on the evolutionary genetics, systematics, and ecology of western Palearctic water frogs (reviewed by Graf & Polls Pelaz 1989; Plötner 2005).

Water frogs in the Eastern Mediterranean area present a huge genetic, organismal, phylogenetic, and ecological diversity. Despite their high similarity in morphology, they show considerable genetic divergence, indicating the existence of several cryptic species. Distinct levels of genetic divergence are present, some lineages are considered to represent distinct evolutionary species (Beerli et al., 1996; Plötner et al., 2010, Akın et al., 2010b). Water frogs are found in all types of freshwater habitats, and because of their sensitive skin structure they are easily affected from environmental changes. All make them hence as a candidate group to study the impact of past historical geological and climatic events on contemporary phylogeography and molecular evolution (Beerli et al., 1996, Plötner et al., 2010, Akın et al., 2010b).

The first phylogenetic study of water frogs from eastern Mediterranean region (although samples are only from the Aegean region) based on 31 protein electrophoretic loci indicated four main groups which are (1) *P. cretensis* and *P. epeiroticus* group, close to (2) *P. perezi* and *P. saharicus* outgroup and (3) *P. lessonae* and *P. shqipericus* group close to (4) a diverse group comprising of *P. cerigensis*, *P. cf.bedriagae* and *P. ridibundus*. The relationships within the last group could not be resolved (Beerli et al., 1996).

A more comprehensive study based on mtDNA (12 rDNA and ND3), including water frogs from Anatolia, Syria, Jordan, and Central Asia, indicated that water frogs from Cyprus and Central Asia represent distinct species. Moreover, Anatolian frogs did not belong to either *P. ridibundus* from Europe or *P. bedriagae* from Syria; the Anatolian clade comprised several subclades (one of them represented the *caralitanus* lineage) and the so-called Ceyhan lineage from the Çukurova plain. All Anatolian lineages formed a single huge clade together with *P. cretensis* and *P. epeiroticus*. In addition to this clade, two additional clades are present: one clade consists of *P. lessonae, P. bergeri* and *P. shqipericus*; the other includes *P. saharicus* and *P. perezi* (Plötner and Ohst, 2001; Plötner et al., 2001). The following study (Lymberakis et al., 2007) based on distinct mtDNA genes (16S rRNA and cytochrome b) verified the previous results and revealed new finding about *P. cerigensis* and *P. kurtmuelleri*, however, their phylogenetic relationships with respect to *P. bedriagae* and P. *ridibundus* remained unresolved in this research.

Certainly the most detailed studies from eastern Mediterranean region using an intensive data set (612 samples) on the basis of mtDNA (ND3 and ND2) documented an extreme genetic diversity which was represented by six main haplogroups (MHGs) (Plötner et al., 2010; Akın et al., 2010b) (Fig 1). MHG1 includes *P. ridibundus* and *P. kurtmuelleri* from European and Balkan populations, respectively. MHG2 indicates the Levant populations (*P. bedriagae*). MHG3 is unique for Cyprus water frogs, described as a new species *P. cypriensis*

(Plötner et al., 2012). MHG4 and 5 are distributed around the Çukurova plain in Anatolia. MHG6 is the greatest group inhabiting an area from western Anatolia to the Caspian Sea. It is subdivided into four subgroups (MHG6a-d). MHG6a is found in the southwestern Anatolia, representing P. cf. *caralitanus*. MHG6b only occurs in western Antalya, and on the islands Rhodos and Karpathos. It was described as a distinct species, P. cf. *cerigensis* (Beerli et al., 1994), its systematic status, however, is not yet clear. MHG6c is distributed from western Anatolia to the Caspian Sea. MHG6d was found in the river systems Euphrates and Tigris (Plötner et al., 2010; Akın et al., 2010b). As for P. cf. *cerigensis*, the systematic status of these two MHGs still remains unclear (Fig 1.1).



Figure 1.1. Geographic distribution of water frogs' main haplogroups (MHGs) and subgroups (ad) of water frogs in Anatolia and neighbouring regions.

1.4. The Present Study

As indicated above water, the frog complex presents a biologically fascinating array of questions; my theses is focused on molecular evolution and phylogeography of eastern Mediterranean region water frog complex. The main aim was to evaluate phylogenetic and phylogeographic hypotheses formulated on the basis of mtDNA (e.g. Plötner et al., 2010; Akın et al., 2010a;b) and intron 1 of the serum albumin gene (Plötner et al. 2009; 2012).

The goal of the first manuscript was to analyze the structure and molecular evolution of the serum albumin gene of the pool frog *P. lessonae*. The complete sequence of the albumin gene including all exons and introns, upstream sequence of the 5' UTR and downstream sequence of the 3' UTR were presented. Potential regulatory elements and functionally important motifs in the 5' and 3' UTR, and in the exon-intron junctions were documented by aligning sequences from closely and distantly related amphibian species. A comparison of exon and intron lengths of the pool frogs with other vertebrate species provided insight about molecular evolution of the albumin gene. Moreover, the entire scaffold of *P. lessonae* was scanned for retroelements which were described and systematized. Finally, phylogenetic relationships between selected water frog species were tested using both nucleotide and amino acid sequences of the albumin exons.

The task of second manuscript was to study phylogeographic patterns of mitochondrial genes (ND2 and ND3) and the nuclear marker serum albumin intron 1 (SAI-1) in water frog complex of the eastern Mediterranean region. These results revealed discordances in the tree topology, the number of genetic stocks, the level of genetic differentiation, the times of divergence and the patterns of geographic distribution. These discordances between two markers are discussed in the light of distinct processes that are incomplete lineage sorting, retention of ancestral polymorphism, introgression after secondary contact in concert with sexbiased or adaptive processes.
1.5. References

Akın, Ç., Bilgin, M., Bilgin, C.C. (2010a) Discordance between ventral colour and mtDNA haplotype in the water frog *Rana (ridibunda) caralitana*, 1988 Arıkan. *Amphibia-Reptilia* 31: 9–20.

Akın, Ç., Bilgin, C.C., Beerli, P., Westaway, R., Ohst, T., Litvinchuk, S.N., et al. (2010b) Phylogeographic patterns of genetic diversity in eastern Mediterranean water frogs were determined by geological processes and climate change in the Late Cenozoic. *J Biogeogr* 37: 211–2124.

Alzohairy, A.M., Gyulai, G., Jansen, R.K., Bahieldin, A. (2013) Transposable elements domesticated and neofunctionalized by eukaryotic genomes. *Plasmid* 69: 1-15.

Avise, J.C., Arnold, J., Ball, R.M., Bermingham, E., Lamb, T., Neigel, J.E., Reeb, C.A., Saunders, N.C. (1987) Intraspecific phylogeography: the mitochondrial DNA bridge between population genetics and systematics. *Annu Rev Ecol Syst* 18: 489-522.

Azad, R.K. and Lawrence J.G. (2011) Towards more robust methods of alien gene detection. *Nucleic Acids Res* 39: e36.

Becq, J., Churlaud, C. Deschavanne, P. (2010) A benchmark of parametric methods for horizontal transfer detection. *PLoS One* 5: e9989.

Beerli, P., Hotz, H., Tunner, H., Heppich, S., Uzzell, T. (1994) Two new water frog species from the Aegean islands Crete and Karpathos (Amphibia, Salientia, Ranidae). *Notulae Naturae, Academy of Natural Sciences of Philadelphia* 470:1-9.

Beerli, P., Hotz, H. & Uzzell, T. (1996) Geological dated sea barriers calibrate protein clock for Aegean water frogs. *Evolution* 50: 1676-1687.

Berger, L. (1967) Embryonal and larval development of F1 generation of green frogs different combinations. *Acta Zool Cracov* 12(7): 123-160.

Berger, L. (1968) Morphology of the F1 generation of various crosses within *Rana esculenta*-complex. *Acta Zool Cracov* 13: 301-324.

Blomme, T., Vandepoele, K., Bodt, S., Simillion, C., Maere, S., Peer, Y.V. (2006) The gain and loss of genes during 600 million years of vertebrate evolution. *Genome Biol* 7(5): R43, doi:10.1186.

Brito, P.H. and Edwards, S.V. (2009) Multilocus phylogeography and phylogenetics using sequence-based markers. *Genetica* 135: 439-455.

Cancherini, D.V., França, G.S., de Souza, S.J. (2010) The role of exon shuffling in shaping protein-protein interaction networks. *BMC Genomics* 11(Suppl 5): 1-13.

Chenais, B., Caruso, A., Hiard, S., Casse, N. (2012) The impact of transposable elements on eukaryotic genomes: from genome size increase to genetic adaptation to stressful environments. *Gene* 509: 7-15.

Cohen, C.J., Lock, W.M., Mager, D.L. (2009) Endogenous retroviral LTRs as promoters for human genes: a critical assessment. *Gene* 448: 105-114.

Conley, A.B., Piriyapongsa, J., Jordan, I.K. (2008) Retroviral promoters in the human genome. *Bioinforma Oxf Engl* 24: 1563-1567.

Conley, A.B., Jordan, I.K. (2012) Cell type-specific termination of transcription by transposable element sequences. *Mob DNA* 3: 15.

Crow, K.D. and Wagner, G.P. (2006) What is the role of genome duplication in the evolution of complexity and diversity? *Mol Biol Evol* 23(5): 887-892.

Deininger, P.L., Batzer, M.A. (2002) Mammalian retroelements. *Genome Res* 12: 1455-1465.

Feschotte, C. (2008) Transposable elements and the evolution of regulatory networks. *Nat Rev Genet* 9: 397-405.

Fiston-Lavier, A.S., Anxolabehere, D., Quesneville, H. (2007) A model of segmental duplication formation in Drosophila melanogaster. *Genome Res* 17: 1458-1470.

Fitzpatrick, D.A. (2012) Horizontal gene transfer in fungi. *FEMS Microbiol Lett* 329: 1-8.

Force, A., Lynch, M., Picket, F.B., Amores, A., Yan, Y.L., Postlethwait, J. (1999) Preservation od duplicate genes by complementary, degenerative mutations. *Genetics* 151: 1531-1545.

França, G.S., Cancherini, D.V., de Souza, S.J. (2012) Evoloutionary history of exon shuffling. *Genetica* 140: 249-257.

Frost, L.S. et al., (2005) Mobile genetic elements: the agents of open source evolution. *Nat Rev Microbiol* 3: 722-732.

Gilbert, W. (1978) Why genes in pieces? *Nature* 271: 501.

Graf, J.D., Polls-Pelaz, M. (1989) Evolutionary genetics of the *Rana esculenta* complex. In: *Evolution and ecology of unisexual vertebrates* p. 289-302. Dawley, R.M., Bogart, J.P., Eds, Albany.

Grau, J.H., Poustka, A.J., Meixner, M., Plötner, J. (2014) LTR retrolements are intrinsic components of transcriptional networks in frogs. *BMC Genomics* 15 (626): 1-16.

Graur, D. and Li, W.-H. (2000) Fundamentals of molecular evolution. Sinauer Associates, Sunderland, Massachusetts.

Gu, X. (2003) Functional divergence in protein (family) sequence evolution. *Genetica* 118: 133-141.

Haegeman, A., Jones, J.T., Danchin, E.G.J. (2011) Horizontal gene transfer in nematodes: a catalyst for plant parasitism? *Mol Plant Microbe Interact* 24: 879-887.

Hewitt, G.M. (2001) Speciation, hybrid zones and phylogeography or seeing genes in space and time. *Mol Ecol* 10: 537-549.

Hickerson, M.J., Cartens, B.C. et al. (2010) Phylogeography's past, present, and future: 10 years after Avise, 2000. *Molecular Phylogenet Evol* 54: 291-301.

Holland, P.W., Garcia-Fernandez, J., Williams, N.A., Sidow, A. (1994) Gene duplications and the origins of vertebrate development. *Dev Suppl* 125-133.

Holland, P.W. (1997) Vertebrate evolution: something fishy about Hox genes. *Curr Biol* 7: 570-572.

Huang, J. (2013) Horizontal gene transfer in eukaryotes: the weak-link model. *Bioessays* 863-875.

Jern, P., Coffin, J.M. (2008) Effects of retroviruses on host genome function. *Annu Rev Genet* 42: 709-732.

Jordan, I.K., Rogozin, I.B., Glazko, G.V., Koonin, E.V. (2003) Origin of a substantial fraction of human regulatory sequences from transposable elements. *Trends Genet* 19: 68-72.

Kazazian, H.H. (2003) Mobile elements: drivers of genome evolution. *Science* 303: 1626-1632.

Keeling, P.J. and Palmer, J.D. (2008) Horizontal gene transfer in eukaryotic evolution. *Nat Rev Genet* 9: 605-618.

Keren, H., Lev-Maor, G., Ast, G. (2010) Alternative splicing and evolution: diversification, exon definition and function. *Nat Rev Genet* 1-11.

Kidwell, M.G. (2002) Transposable elements and the evolution of genome size in eukaryotes. *Genetica* 115: 49-63.

Kidwell, M.G., and Lisch, D.R. (2000) Transposable elements and host genome evolution. *TREE* 15(3): 95-99.

Knowles, L.L., Carstens, B.C. (2007) Delimiting species without monophyletic gene trees. *Sys Biol* 56: 887-895.

Kolkman, J.A. and Stemmer, W.P.C. (2001) Directed evolution of proteins by exon shuffling. *Nat Biotechnol* 19: 423-428.

Kondrashow, F.A. and Koonin, E.V. (2001) Origin of alternative splicing by tandem exon duplication. *Hum Mol Genet* 10: 2661-2669.

Konkel, M.K., Batzer, M.A. (2010) A mobile threat to genomic stability: the impact of non-LTR retrotransposons upon the human genome. *Semin Cancer Biol* 20: 211-221.

Lawrence, J.G. and Ohman, H. (1997) Amelioration of bacterial genomes: rates of change and exhange. *J Mol Evol* 44: 383-397.

Lee, J.Y., Ji, Z., Tian, B. (2008) Phylogenetic analysis of mRNA polyadenylation sites reveals a role of transposable elements in evolution of the 3'-end of genes. *Nucleic Acids Res* 36: 5581-5590.

Levin, H.L., Moran, J.V. (2011) Dynamic interactions between transposable elements and their hosts. *Nat Rev Genet* 12: 615-627.

Li, W.-H. (1997) Molecular Evolution. Sinauer Associates, Sunderland, Massachusetts.

Liu, M., Grigoriev, A. (2004) Protein domains correlate strongly with exons in multiple eukaryotic genomes - evidence of exon shuffling? *Trends Genet* 20: 399-403.

Long, M., Betran, E., Thornton, K., Wang, W. (2003) The origin of new genes: glimpses from the young and old. Nat Rev Genet, 4(11): 865-875.

Lorente-Galdos, B., Bleyhl, J., Santpere, G., Vives, L., Ramirez, O., Hernandez, J., Anglada, R., Cooper, G.M., Navarro, A., Eichler, E.E., Marques-Bonet, T. (2013) Accelerated exon evolution within primate segmental duplications. *Genome Biol* 14(R9): 1-12. Lymberakis, P., Poulakakis, N., Manthalou, G., Tsigenopoulos, C.S., Magoulas, A., Mylonas, A. (2007) Mitochondrial phylogeography of Rana (Pelophylax) populations in the eastern Mediterranean region. *Mol Phylogenet Evol* 44: 115-125.

Mariño-Ramírez, L., Lewis, K.C., Landsman, D., Jordan, I.K. (2005) Transposable elements donate lineage-specific regulatory sequences to host genomes. *Cytogenet Genome Res* 110: 333-341.

Meisel, R.P. (2009) Repeat mediated gene duplication in the *Drosophila pseudoobscura* genome. *Gene* 438: 1-7.

Medstrand, P., van de Lagemaat, L.N., Dunn, C.A., Landry, J.-R., Svenback, D., Mager, D.L. (2005) Impact of transposable elements on the evolution of mammalian gene regulation. *Cytogenet Genome Res* 110: 342-352.

Meyer, A., Van de Peer, Y. (2005) From 2R to 3R: evidence for a fish-specific genome duplication (FSGD). *Bioessays* 27: 937-945.

Nei, M., Kumar, S. (2000) Molecular Evolution and Phylogenetics. Oxford University Press, New York.

Nylander, J.A.A., Ronquist, F., Huelsenbeck, J.P., Nieves-Aldrey, J.L. (2004) Bayesian phylogenetic analysis of combined data. *Syst Biol* 53: 47-67.

Ohno, S. (1970) Evolution by gene duplication. New York: Springer-Verlag.

Page, R.D.M., and Holmes, E.C. (1998) Molecular evolution: a phylogenetic approach. Blackwell, Osney Mead, Oxford, U.K.

Patthy, L. (1999) Genome evolution and the evolution of exon-shuffling - a review. *Gene* 238: 103-114.

Plötner, J. (2005) *Die westpaläarktischen Wasserfrösche*. Von Märtyrern der Wissenschaft zur biologischen Sensation. Z f Feldherpetologie, Beiheft 9, Laurenti, Bielefeld.

Plötner, J., Ohst, T., Böhme, W., Schreiber, R. (2001) Divergence in mitochondrial DNA of Near Eastern water frogs with special reference to the systematic status of Cypriot and Anatolian populations (*Anura, Ranidae*). *Amphibia – Reptilia* 22: 397-412.

Plötner, J., Ohst. T. (2001) New hypothesis of the western Palearctic water frog complex (Anura: Ranidae). *Mitt Mus Nat kd Zool Reihe* 77: 5-21.

Plötner, J., Köhler, F., Uzzell, T., Beerli, P., Schreiber, R., Guex, G.-D. & Hotz, H. (2009) Evolution of serum albumin intron-1 is shaped by a 5' truncated non-long terminal repeat retrotransposon in western Palearctic water frogs (Neobatrachia). *Mol Phylogenet Evol* 53: 784–791.

Plötner, J., Uzzell, T., Beerli, P., Akın, Ç., Bilgin, C. C., Haefeli, C., Ohst, T., Köhler, F., Schreiber, R., Guex, G.-D., Litvinchuk, A. N., Westaway, R., Reyer, H.-U., Hotz, H., (2010) Genetic divergence and evolution of reproductive isolation in eastern Mediterranean water frogs. In: Glaubrecht, M., Schneider, H. (Eds.), Evolution in action. *Case studies in adaptive radiation and the origin of biodiversity. Special volume from the SPP 1127 "Radiations – Genesis of Biological diversity" of the DFG.* Springer, Heidelberg, Berlin.

Plötner, J., Baier, F., Akın, Ç., Mazepa, G., Schreiber, R., Beerli, P., Litvinchuk, S.N., Bilgin, C.C., Borkin, L., Uzzell, T. (2012) Genetic data reveal

that water frogs of Cyprus (genus Pelophylax) are an endemic species of Messinian origin. *Zoosyst Evol* 88: 261-283.

Rajarapu, G. (2014) Horizontal gene transfer. *Phylogenetics and Evolutionary Biology* 2(2):1-3.

Ramos, O.M., Ferrier, D.E.K. (2012) Mechanisms of gene duplication and translocation and progress towards understanding their relative contributions to animal genome evolution. *International Journal of Evolutionary Biology* 1-10, doi:10.1155/2012/846421.

Rebollo, R., Romanish, M.T., Mager, D.L. (2012) Transposable elements: an abundant and natural source of regulatory sequences for host genes. *Annu Rev Genet* 46: 21-42.

Roman, Y., et al., (2007) Biochemical characterization of a SET and transposase fusion protein. Metnase: its DNA binding and DNA cleavage activity. *Biochemistry* 46: 11369-11376.

Roy-Engel, A.M., et al. (2005) Human retroelements may introduce intragenic polyadenylation signals. *Cytogenet Genome Res* 110: 365-371.

Schaack, S., Gilbert, C., Feschotte, C. (2010) Promiscuous DNA: horizontal transfer of transposable elements and why it matters for eukaryotic evolution. *Trends Ecol Evol* 25: 537-546.

Schönknect, G., Weber, A.P.M., Lercher, M.J. (2013) Horizontal gene acquisitions by eukaryotes as drivers of adaptive evolution. *Bioessays* 36: 9-20.

Sela, N., Mersch, B., Hotz-Wagenblatt, A., Ast, G. (2010) Characteristics of transposable elements exonizationnwithin human and mouse. *PLoS One* 5: e10907.

Smalheiser, N.R., Torvik, V.I. (2005) Mammalian microRNAs derived from genomic repeats. *Trends Genet* 21: 322-326.

Storz, J.F., Opazo, J.O., Hoffman, F.G. (2013) Gene duplication, gene duplication, and the functional diversification of vertebrate globins. *Mol Phylogenet Evol* 66(2): 469-478.

Sun, C., et al., (2011) LTR retrotransposons contribute to genomic gigantism in plethodontid salamanders. *Genome Biol Evol* 4: 168-183.

Syvanen, M. (2012) Evolutionary implications of horizontal gene transfer. *Ann Rev Genet* 46: 341-358.

Tang, H. (2007) Genome assembly, rearrangement, and repeats. *Chem Rev* 107: 3391-3406.

Teng, L., Firpi, H.A., Tan, K. (2011) Enhancers in embriyonic stem cells are enriched for transposable elements and genetic variations associated with cancers. *Nucleic Acids Res* 39: 7371-7379.

Van de Lagemaat, L.N., Landry, J.-R., Mager, D.L., Medstrand, P. (2003) Transposable elements in mammals promote regulatory variation and diversification of genes with specialized functions. *Trends Genet TIG* 19:530-536.

Vibranovski, M.D., Sakabe, N.J., de Oliveira, R.S., de Souza, S.J. (2005) Signs of ancient and modern exon-shuffling are correlated to the distribution of ancient and modern domains along proteins. *J Mol Evol* 61: 341-350.

Vibranovski, M.D., Sakabe, N.J., de Souza, S.J. (2006) A possible role of exonshuffling in the evolution of signal peptides of human proteins. *FEBS Letters* 580: 1621-1624.

Whitaker, J.W., McConkey, G.A., Westhead, D.R. (2009) Prediction of horizontal gene transfers in eukaryotes: approaches and challenges. *Biochemical Soc T* 37: 792-795.

Yang, Z. (2006) Computational molecular evolution. Oxford University Press.

Zhang, J.Z. (2003) Evolution by gene duplication: an updatae. *Trends Ecol Evol* 18: 292-298.

Zhu, B., Lou, M-M., Xie, G-L., Zhang, G-Q. et al., (2011) Horizontal gene transfer in silkworm Bombyx mori. *BMC Genomics* 12: 248.

CHAPTER 2

MANUSCRIPT 1: MOLECULAR EVOLUTION AND CHARACTERIZATION OF THE SERUM ALBUMIN GENE OF THE POOL FROG, *PELOPHYLAX LESSONAE*

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2.1. Introduction

Serum albumin (SA) is a globular monomeric protein containing only one polypeptide chain of 605-610 amino acid residues. As the most abundant component of the plasma protein system SA is essential for maintaining the colloid osmotic (oncotic) pressure, allowing a proper distribution of body fluids between intravascular compartments and body tissues. It also serves as a transport protein for numerous endogenous and exogenous compounds, for example hormones, metals, fatty acids, and diverse metabolites (reviewed by Rothschild et al 1988; Fanali et al., 2012). In mammals and other vertebrates SA is synthesized in the liver (Peters and Anfinsen 1950, Miller et al. 1951, 1954); in frogs albumin is also expressed in skin (Zhang et al. 2005).

SA is encoded by a single gene except for the tetraploid frog species *Xenopus laevis* which possesses, as a result of genome duplication, two SA genes and the

dodecaploid *X. ruwenzoriensis* which even has three albumin encoding genes (Westley et al. 1981). All vertebrate SA genes so far analysed were composed of 15 exons and 14 intervening sequences (Sargent et al. 1981, Minghetti et al. 1986).

We here reported the complete SA gene sequence of the pool frog species *Pelophylax lessonae*. Based on comparisons with homologous sequences of other frog species, putative regulatory elements and functional motifs have been identified. Moreover, we compared exon and intron lengths of the water frog with other vertebrate species to get insight about molecular evolution of the albumin gene. We also described retroelements inserted in the entire scaffold of *P. lessonae*. As a highly polymorphic protein SA was often used as a genetic marker for systematic, phylogenetic, and population genetic studies in WPWFs (e.g. Tunner 1979, Uzzell and Hotz 1979, Uzzell 1979). Based on exon sequences we re-analyzed the phylogenetic relationships of selected species and compared our results with those obtained from studies on other mitochondrial and nuclear DNA markers (Plötner, 1998; Plötner & Ohst, 2001; Plötner, 2005; Plötner et al., 2007; 2009; 2010; 2012; Akın et al., 2010b).

2.2. Material and Methods

2.2.1. Sources of Albumin Sequences

The SA genes of the *Silurana tropicalis* (release v7.1) and *Xenopus laevis* genomes (release v7.2) were downloaded from Xenbase.org (Bowes et al., 2008). Albumin data of *Gallus gallus* (NC_006091, Hillier et al., 2004), *Rattus norvegicus* (NC_005113, Gibbs et al., 2004), *M. musculus* (NC_000071, Church et al., 2009) and *Homo sapiens* (NC_000004, Hillier et al., 2005) were obtained from GenBank. The *P. lessonae* SA gene was obtained from a genome assembly with an N50 of 21kb constructed from several next generation sequence libraries

of different lengths using SOAPdenovo genome assembler (Poustka, Grau and Plötner, unpublished). The SA gene was identified with BlAST searches using default parameters and the *Silurana* sequence as a query; it was included in a scaffold of ca. 64kb.

2.2.2. Alignment and Motif Searches

The alignment of the serum albumin exon sequences of *P. lessonae* with the coding sequence of *P. shqipericus*, the alignment of the 5' and 3' regions of the *P. lessonae* serum albumin gene with those of other amphibian species, and alignments of exonic and intronic sequences of the *P. lessonae* albumin gene with those of other vertebrate species were conducted in MEGA 6.06 (Tamura et al., 2013) using the ClustalW (Thompson et al., 1994) and Muscle algorithms (Edgar, 2004). Alignments were manually improved.

2.2.3. Search For Transposable Elements and Repetitive Sequences

To search for available interspersed repeats and low complexity DNA elements in the P. lessonae scaffold (64.077 bp), which included the complete albumin gene, the program Repeat Masker (Smit, Hubley and Green unpublished; RepeatMasker http://repeatmasker.org) program TRF at and the (http://tandem.bu.edu/trf/trf.html) (Benson, 1999) were used. Options used in the engine=cross Repeat Masker were search include search match; speed/sensitivity=slow; DNA source=vertebrate.

As a result of Repeatmasker searches, truncated transposable elements (TEs) were found in the *P. lessonae* scaffold which matched with complete TEs of vertebrate genomes stored in the Repbase repeat database (Jurka et al., 2005). These reference sequences were used as queries in BLAST searches against the *P. lessonae* genome to find copies of these elements specific to *P. lessonae*, termed as PL elements.

Because most of the PL elements are truncated, consensus sequences were reconstructed by aligning several hundred of these copies in the *P. lessonae* genome. Alignments of several copies of the complete elements were performed in MEGA 6.06 (Tamura et al., 2013) using ClustalW algorithm (Thompson et al., 1994) and Muscle algorithm (Edgar, 2004).

Each consensus element was screened for target site duplications, repeats, and transposase domains. Inverted terminal repeats of the derived consensus elements found the **EMBOSS** explorer were using einverted repeat (http://emboss.bioinformatics.nl/cgi-bin/emboss/einverted). ORF Finder (http://www.ncbi.nlm.nih.gov/gorf/gorf.html) was used to identify putative open reading frames from deduced nucleotide sequences and the types of proteins encoded by autonomous elements. Derived consensus elements and their translated proteins were blasted against nucleotide sequence and protein databases such as NCBI (http://www.ncbi.nlm.nih.gov/), UCSC genome browser (http://genome.ucsc.edu/), and EMBL (http://www.embl.de/) to identify and annotate element specific genes.

2.2.4. Phylogenetic Analyses Based on Protein Coding Exon Sequences

To reveal phylogenetic relationships among western Palearctic water frogs on the basis of albumin gene, nucleotide sequences of protein coding exons (up to the first stop codon) were used. Sequences of *Rana chensinensis, Rana kukunoris, X. laevis,* and *S. tropicalis* were included for outgroup comparisons. Sequence alignment based on the amino acid sequences was performed with MAFFT (http://mafft.cbrc.jp/alignment/server/) using the L-INS-i algorithm (Katoh et al., 2005).

In order to test whether positive selection is operating on the gene, a codon-based z-test was performed for all sequences (scope=overall average) on the basis of the modified Nei-Gojobori method (Nei and Kumar, 2000) as implicated in MEGA6.0 under the alternative hypothesis $d_N > d_S$ (positive selection) where d_N

is the number of non-synonymous substitutions per non-synonymous site and d_S is the number of synonymous substitutions per synonymous site (d_S), and the null hypothesis ($d_N = d_S$) where sequence evolution of protein coding exons is follow a neutral model of sequence evolution.

Genealogical relationships of SA sequences were reconstructed with maximum likelihood (ML) as implemented in MEGA 6.06 (Tamura et al., 2013) using the option 'partial deletion' for gaps/missing data treatment and 'Nearest-Neighbor-Interchange' as the heuristic search method. Nodal support was evaluated by bootstrapping (Felsenstein, 1985) with 1,000 replicates. The model that best describe the patterns of sequences of sequence evolution was selected on the basis of the Bayesian Information Criterion (BIC, Schwarz, 1978) and ML as implemented in MEGA. As a result the General Time Reversible (GTR) model (Tavare, 1986) with gamma-shaped rate variation (G=2.58) was chosen. (BIC=12505.7). Genetic distances were calculated on the basis of uncorrected p distance and the number of nucleotide and aa differences among distinct water frog species.

2.3. Results and Discussion

2.3.1. Structure of the Albumin Gene

The SA gene of the pool frog covers 32,460 bp starting from the putative cap site to the first poly(A) addition site. It consists of 15 exons, interrupted by 14 introns as in mammals and other frog species (Fig. 2.1, Appendices 2.1 and 2.2) (Sargent et al., 1981a,b; Minghetti et al., 1986; Schorpp et al., 1988). As in human, pool frog' introns are located within polypeptide chain between codon 3 and 1 and between 2 and 3 alternately. It is found in all three domains of SA protein, which supports the idea that albumin gene arose from a single primordial domain as a result of triplication (Sargent et al., 1981). In contrast to the SA gene lengths of

Xenopus laevis (11868 bp) and *Silurana. tropicalis* (12465 bp), the SA gene of pool frogs is significantly longer caused by much longer intronic sequences (Appendix 2.1) which are inserted by many retroelements (REs) (Table 2.1).



Figure 2.1. Schematic structure of pool frog'serum albumin gene from the Cap site of the first exon to the first polyadenylation site of the last exon. Translated exons are red marked; the partially translated exon 1 and exon 14 and the completely untranslated exon15 are yellow coloured. Introns are indicated by dark grey boxes.

2.3.1.1. The 5' Region of the Water Frog Albumin Gene

In their 5' region of the SA gene, from the cap site at position +1 to position -330, water frogs revealed considerable homology to Rana chensinensis, R. kukunoris, X. laevis and S. tropicalis (Fig. 2.2). Several conserved motifs are seen which may play functional roles for gene regulation in frogs and other vertebrates. The consensus sequence of the initiator element (Inr) (PyPyA₊₁NT/APyPy) of the water frog SA gene is identical to that of mammals (Bucher, 1990). The TATA box (TATAAATGT) located from -32 to -24 (Goldberg, 1979) is conserved in all six species. Only two base subtitutions are seen in X. laevis and S. tropicalis. A 13 bp hepatocyte-specific promoter element (HP1), with a consensus sequence GNTANTNNTNNNC (Schorpp, et al., 1988b), is located at positions -65 to -53. Except for three base changes in two positions, this element is extremely conserved, not only in the SA gene of the amphibians used in this study, but also in the promoters of the albumin and alpha-fetoprotein genes of chicken and mammals; this indicates an important function for hepatocyte specific expression for these genes and as a regulatory element during vertebrate evolution (Kugler et al., 1988; Schorpp, et al., 1988). Presence of a TATA box within a distance of 70

bp is required for the function of the HP element (Ryffel et al., 1989). A putative CAT box sequence, CCAAT was found at position -178; it is conserved among S. tropicalis, X. laevis, P. ridibundus and P. lessonae. The presence of a CAT box in the SA gene coincides with findings of Mantovani' (1998) who surveyed 96 unrelated SA promoters, including that of the X. laevis, but contradicts the statement of Schorpp et al. (1988a) who postulated the absence of a CAT box in the 68 kd and 74 kd albumin genes of X. laevis. Downstream from the CAT box, two conserved motifs were found. The first 10 bp motif with the consensus sequence AATYACCMAA, named amphibian downstream element I (ADE I), is located between positions -255 and -264. The second element (ADE II) is 37 bp long, located between positions -282 and -318, is highly conserved (81%) among X. laevis, S. tropicalis, P. ridibundus, and P. lessonae. It is partially overlapped by downstream element III (DE III) found in promoters of the albumin genes of rat, human, and mouse (Cereghini et al., 1987;1988), with which exhibits 59% of homology. Similar to amphibians, several regulatory elements were identified within a 150 bp region of human, rat and mouse SA promoters, including a TATA box, a so called proximal element (PE), a CAT box, DE I, II and III. These elements are thought to be involved in tissue-specific expression of the albumin gene (Cereghini et al., 1987). The specific functions of these elements, however, are not yet clear and have to be analysed experimentally.

	-340	-318	-281	-264	-255
			ADE II	4	ADE I
P.lessonae	TTTCAGACGCAG	ACTGGGATAC <mark>TGTCATAATCAAGG</mark>	ATCACTTTGCTTAACCTTTTGTTC	GTTTCACAGCCT <mark>AAT(</mark>	CACCAAACGCTTACATTAGTA
P.ridibundus	TTTCAGACGCAG	GACTGGGATAC <mark>TGTCATAATCAAGG</mark>	ATCACTTTGCTTAACCTTTTGTTC	GTTTCACAGCCT <mark>AAT(</mark>	CACCAAACGCTTACATTAGTA
R.chensinensis					
R.kukunoris					
X.laevis	ATTTGGGCACAT	ACCAATACACTAGTATACACAAAG	ATCAGTTTGCTTAACCTTTTGTTC	AGC <mark>AAT</mark>	<mark>FACCAAA</mark> GACGTTGACTAGCC
S.tropicalis	TTTGGGGCACAG	ACCAGTACA- <mark>TAGTATACACAAAG</mark>	ATCAGTTTGCTTAACCTTTTGTTCAC	6TAAT	TACCCAA GACTTTGCCTTGCC
	-241		-17	78	-143
			CAT	Г ВОХ	
P.lessonae	ΑΑΑΤΤΑΤΑΑΟΤΟ	GATCTACAAGGTCTTTCAA-	AAAATGTGTTG <mark>(</mark>		TTTACAACTGATTCAGTATTG
P.ridibundus	ΑΑΑΤΤΑΤΑΑΟΤΟ	GATCTACAAGGTCTTTCAA-	AAAATGTGTTG <mark>(</mark>		TTTACAACTGATTCAGTATTG
R.chensinensis					CAGTATTG
R.kukunoris				<mark>T</mark> GCAAGCCCACT	TTAACAATTGATTCAGTATTG
X.laevis	САТС	GCTAGGTTTTTTTTCACAAT	TTAAAAGGTTTTTCAAAATTCAGAAAA	CAATATA-GAGCAAG	CAG-CAATACGTTATT-

Figure 2.2. 5' regions of the serum albumin gene P. *lessonae, P. ridibundus, P. chensinensis, P. kukunoris, X. laevis, and S. tropicalis, starting with the translation initiation codon (Met) up to position -340, the start site transcription (+1). Dashes indicate gaps introduced to optimize in the alignment. Conserved regulatory motifs are red colour and underlined.*

HP ELEMENT

P.lessonae	TGTTTTGTCATTCTTTGT-G-TTTCTAGAAA-CGAGCCAAATTGTAATTAATTATAAAGCCAGTCTG <mark>GTTAATCATTT-C</mark> CAAATAGGA
P.ridibundus	TGTTTTGTCATTC-TTGT-G-TTTCTGGAAA-CGAGCCAAATTGTAATTAATTATAAAGCCAGTCTG <mark>GTTAATTATTT-C</mark> CAAATAGGA
R.chensinensis	TGTTTTGCCATTCTTTGT-G-TTTCTAGAGA-CGAGCCAAAATTTGTA-TTAATTATAAAGCCAGTCTG <mark>GTTAATTATTT-C</mark> TGAATAGGA
R.kukunoris	TGTTTTGGCATTCTTTGT-G-TTTTTAGAGA-CGAGCCAAAATTTGTA-TTAATTATAAAGCCAGTCTG <mark>GTTAATTATTT-C</mark> TGAATAGGA
X.laevis	TGACCTTAAAAGTTGATTGACATTAGGAAATTCCACAAAGCTAAAACAACTGCAAACAGAACAATTTGATAG <mark>GTTAATAATTTTC</mark> CAGATCTCT
S.tropicalis	TGACTTTAGGAAACTCCACAAAGCTTAAAAAACTGCAA-CAGAACAATTTGATAG <mark>GTTAATAATTTTC</mark> CAGATCTCT

	-32	+1	Met
	ΤΑΤΑ ΒΟΧ	CAP SITE	
P.lessonae	TGCAAAAATGG <mark>TATAAATGT</mark> CTGCCTTTCC	CTTAGCTGAT- <mark>TTAGGCT</mark> TCTAAGAGAACCTC	-CAAAAAACATTTGCCACC <u>ATG</u>
P.ridibundus	TGCAAAAATGG <mark>TATAAATGT</mark> CTGCCTTTCC	CTTAGCTGAT- <mark>TT<u>AGGCT</u>TCTAAGAGAACCTC</mark>	-CAAAAAACATTTGCCACC <u>ATG</u>
R.chensinensis	TGCAAAAATTG <mark>TATAAATGT</mark> CTGCCTTTCC	TTTAGCTGAT- <mark>TTAGGCT</mark> TCCAAGAGATCCTC	-CTAAAAACATTTGCAACC <u>ATG</u>
R.kukunoris	TGCAAAAATTG <mark>TATAAATGT</mark> CTGCCTTTCC	TTTAGCTGAT- <mark>TT<mark>AGGCT</mark>TCCAAGAGATCCTC</mark>	-CTAAAAACATTTGCAACC <u>ATG</u>
X.laevis	CTGAGCAATAG <mark>TATAAAACA</mark> AGAGGTATCA	ACTCATTTCAGA <mark>TCAGGCT</mark> TCTCAGAGGTCCCC	ACCCAATACATCTCCAGTCATG
S.tropicalis	CTTAGGAATAG <mark>TATAAAGCA</mark> AGAGGTATCA	ACTCATTTCAGA <mark>TCAGGCT</mark> TCTATGAGGTCCCC	ACCCAATACACCTCCAGTCATG

Figure 2.2. (continued).

2.3.1.2. The 3' Region of the Water Frog Albumin Gene

The 3' UTR region extends from the stop codon following exon 14 to the end of exon 15. It contains several conserved cis-regulatory elements (Fig. 2.3); 5' to 3', these are poly(A)-limiting elements a and b (PLEa, PLEb), U-rich upstream element 1 (USE1), poly(A) signal (PAS), U-rich upstream element 2 (USE2), polyadenylation site (PAS), and GU-rich downstream element (GU-rich DSE). PAS is the essential motif for 3' terminal cleavage and efficient polyadenylation, AAUAAA (Conway and Wickens, 1985). It is completely conserved among these species except that its position in R. chensinensis and R.kukunoris is shifted nine bases downstream; (the positions of these PAS were obtained from UTRscan (UTRdb and UTRsite, 2010). USE1 and USE2 are conserved U-rich upstream elements. They were found within 20 nt of both the 5' and 3' ends of PAS in all of these species. In P. lessonae conserved GU-rich downstream element (GU-rich DSE) was identified just of the poly(A) site (0). Although PAS is certainly needed for mRNA 3' end polyadenylation, the presence of these conserved USEs in all of these species (and of GU-rich DSE in *P. lessonae*) support their functional roles in efficient cleavage and polyadenylation. PAS and GU-rich DSE are known binding sites for cleavage and polyadenylation specificity factor (CPSF) and cleavage stimulatory factor (CstF), promoting enzymatic reactions of the 3'end processing (MacDonald et al., 1994; Murthy and Manley, 1995). Similarly, USE1, core upstream element (Hu et al., 2005) is a target site for specific protein complex that enhances element 3' end formation (Danckwardt et al., 2007). Graber et al. (1999) suggested that all yeast and plant 3'UTR have USE2 between the PAS and poly(A) site, indicating that it is evolutionary conserved element (Hu et al., 2005). Most of eukaryotes have this conserved pattern of USE1-PAS-DSE in their 3' UTR (Proudfoot, 2011). poly(A)-limiting elements (PLE_A and PLE_B) are two cisacting elements, firstly identified in X. laevis albumin pre-mRNA have also been found (PLE_A, between -176 and -159; and PLE_B, between -131 and -108). These elements in the last exon are independently involved in regulating of the length poly(A) tail on albumin pre-mRNA, and may cause its short (17-residue). Such nuclear control of poly(A)-tail length is common among vertebrates (Gupta et al., 1998; Gu et al., 1999). In the alignment, they showed nearly 70% homology with the elements found in *X. laevis*. Finally, in *S. tropicalis* iron responsive element (IRE) was found between -146 and -10 as a result of UTRscan (UTRdb and UTRsite, 2010). IRE are perhaps involved in mRNA stability: a low iron concentration may enhance the binding affinity of the IRE binding protein to the IREs, which may in turn inhibit attack by the nuclease on the cleavage site; when iron concentration increases, the IRE-binding protein is released and nucleolytic cleavage takes place (Binder et al., 1994). Functional roles of PLE_A and PLE_B in the poly(A) length control, in *P. lessonae, P. ridibundus* and in other species and IRE in albumin pre-mRNA in *S. tropicalis* should be also tested experimentally.

St	op Codon	PL	E _A				PLE _B	
		-176	-159			-131		-108
P.lessonae	UGACAU-GGAACAGGCCA-		AGAACAAGA	- CUGUGUUGAG	CCACUAGGUL	J <mark>AAACCAUCAI</mark>	JCAACUGGUC	<u>ACUGA</u>
P.ridibundus	UAACAU-GGAACAGGCCA-	<mark>AAAGCUGAU</mark>	AGAACAAGA	- CUGUGUUGAG	CCACUGGGUL	J <mark>AAACCAUCAI</mark>	JCAACUGGUC	<u>ACUGA</u>
R.catesbeiana	UAACAU-GGAACAGGCCA-	<mark>CAAGCUAAU</mark>	AUAACAAGA	-CAGCCUUGAG	CCACUAGGUL	J <u>AAAUCAUCAI</u>	JCAACUGGUC	<u>ACUGA</u>
R.chensinensis	UGACAU-GGAACAGGCCA-	<mark>CAAGCUGAU</mark>	AGAACAAGA	-CAGCCAUGAG	CCACUAGGUL	J <mark>AAAUCAUCAI</mark>	JUAACUGGUC	<u>AUUAA</u>
R.kukunoris	UGACAU-GGAACAGGCCA-		AGAACAAGA	-CAGCCAUGAG	CCACUAGGUL	J <mark>AAAUCAUCAI</mark>	JUAACUGGUC	AUUAA
X.laevis	UAA GAGUCCAUAAGAGCAAAGACCAG	CCUU <mark>CAAACUCAC</mark>	UGAGGAACACCUU	CCAUCUCUCAA	ACACAAGAAA	AAAAGUUCCU	JCAGCUGAAA	<u>AGA</u>
S.tropicalis	UAAGAGCAAAGGCCAG	CCUU <mark>CAAACUGAC</mark>	CAAGGAAGA	CCAUCUCU C AA	ACAC-AG-AA	AAA <mark>AAGUUUUCUI</mark>	JCAGCUGAAA	AU-GA
				USE1	PAS	5	USE2	
	-107			-52 -46	-38	-33 -23	3 -18	
P.lessonae	-UACAUGUCUCUG	AAGACUCAAGUAG	AAAAAAACAAUCA	<u>UUUUACU</u> G	CAAA AAUA	AAAUAAA	<u></u>	AU
P.ridibundus	-UACAUGUCUCUG	AAGACUCAAGUAG	AAAAAAACAAUCA	<u>UUUUACU</u> G	CAAA AAUA	AAAAAUAAA	<u>JUCUUU</u>	AU
R.catesbeiana	AUACAUGUUGCUGAUACAUGUCUCUG	AAGACUCAAGUAG	AAAAAAACUACCA	UCG <mark>UUUUACU</mark> G	CAAAAU <u>AAUA</u>		<u>JUCUUU</u>	AU
R.chensinensis	AUACAUGUUGCUGAUACAUGUCUCUG	AAGACUCAAGUAG	AAAAAAACUACCA	UCG <mark>UUUUACU</mark> G	CAAAAUAAUA	AGAAA AAUAAA	<u>JUCUUC</u>	A-
R.kukunoris	AUACAUGUCGCUGAUACAUGUCUCUG	AAGACUCAAGUAG	AAAAAAACUACCA	UCG <mark>UUUUACU</mark> G	CAAAAUAAUA	AGAAA	<u>JUCUUC</u>	A-
R.kukunoris X.laevis	AUACAUGUCGCUGAUACAUGUCUCUG GCAU-UUGCUCAG	AAGACUCAAGUAG ACCAUUCAACU	AAAAAAACUACCA	UCG <mark>UUUUACU</mark> G G <mark>UGUG - UU</mark> G	CAAAAUAAUA C AAUA	AGAAA <u>AAUAAA</u> AAAUAAA	JUCUUC GCAUUUAAAA	A- AAA-U

Figure 2.3. Sequence alignment of the 3' UTR of *P. lessonae, P. ridibundus, R. catesbeiana, R.chensinensis, R. kukunoris, X. laevis* and *S. tropicalis* albumin genes. Poly(A) site is accepted as position 0. Regulatory motifs both in the upstream and downstream of the poly(A) site are shown in underlined red colour.

Poly(A)				
	Site DSE (GU rich)			
	-1+1 +14			
P.lessonae	ga-c-ugg <u>auguucuuggugucug</u> uuuuuaug			
P.ridibundus	GA-C-UGG <mark>-</mark>			
R.catesbeiana	GA-C-CAG <mark>-</mark>			
R.chensinensis	GAAC-UGG <mark>-</mark>			
R.kukunoris	GAAC-UGG <mark>-</mark>			
X.laevis	ATATAGAG			
S.tropicalis	AUAUAGC			

Figure 2.3. (continued).

2.3.1.3. 5' and 3' Splice Junctions of the Water Frog's Albumin Gene

To identify consensus sequences at 5' and 3' splice junctions, all fourteen splice sites were aligned among *P. lessonae, S. tropicalis* and *X. laevis*. The consensus sequence of the first six nucleotide at 5' splice site is GTAAGT as indicated by Mount (1982). Upstream of the 3' splice site, the branch point sequence, the pyrimidine rich tract, and at the intron-exon junction C/T,AG are found and their consensus sequences are concordant with Mount (1982)'s findings. A putative branch point consensus sequences (T/CNT/CTA/GAT/C) including the reactive adenosine involve in the lariat formation during pre-mRNA splicing (Ruskin et al., 1984; Reed and Maniatis, 1985) is found in different positions in all species (Fig 2.4).

	Exon		Intro	n	Exon	
XL1	ACAG	<u>GTAAGC</u>CTTTAAATGCATTCATCGTTATTGAAATCCAAAA	XL1	TCATATATAT <u>TGATAAT</u> ACTTTATTATGCCTTTCATA <mark>CAG</mark>	ATGT	XL2
ST1	GCAG	GTAAGA TTTCAAATACATTAATCATTATTAAAATCCAAAG	ST1	TATATTTTAT <u>TTTTAAT</u> CCTTAATAATGCCTTTCTTG <mark>TAG</mark>	ATCA	ST2
PL1	CATG	GTAAGCAAGAATTTTTAGAACAGCAATCTTTATGTCCATA	PL1	TGTGTTTCCTATGCATTACATTTTTTATCTGTTTTTT	AAGA	PL2
XL2	GACT	GTAAGAATTGTATCTAAATTTACAATATATGCAAATAATT	XL2	TGGTGGATGAGTTCCTTATTCAGTATTATTGTGTTTGCAG	TACA	XL3
ST2	GACT	GTAAGAATTATTTCTACAAATAATTATAGTTTTTATGGT	ST2	TGGTGGATGAGT <u>TCCTAAT</u> TCAGTACTATTATTTTTG <mark>CAG</mark>	TACA	ST3
PL2	AATT	GTAAGTATTCAATTTATATAATGTGTTTTATTGTTCTGTA	PL2	TAGAATGAATATGTATTTACTATGCTGCTATGTCTTACAG	AGTG	PL3
XL3	CATA	GTGAGTAATTTTTTGTTTGTGTACTGAGCATATACCCAT	XL3	TTGTGCTATGTTTT <u>CACTAAT</u> TAAATTCACTATTTTA <mark>TAG</mark>	GGCA	XL4
ST3	AATT	GTAAGTAATTATAAGATTGTGTACTGTGCATATACATC	ST3	TTTGTGCTATTTTGCAAACTAATTTATTATTATTTCGCAG	GATA	ST4
PL3	AGCG	GTAAGC TCTCCTTGGCTCCTTCCCCGCAAGCTTTACATAA	PL3	CTGCCTT <u>CTTTAAT</u> TGTGTGTGTGTCTCTTGTGTCTG <u>CAG</u>	ATTG	PL4
XL4	CAGC	GTAATTATGAACAGTTTATTTATTTATTTATTTATTTTCCA	XL4	CTGGTTTGGATACTTACCTAAACTATTTATATGTCTA TAG	ATTC	XL5
ST4	CAGT	GTAAATATGAACACTTTATTTTATTTTTTCAAAACACATG	ST4	GGATACTTACCTAAACTCTCTTTTATTTTTTTGTCTA	GTTC	ST5
PL4	СТТА	GTGAGT TTATTTTAGTATTTTATATATACCATTGTACTTG	PL4	TTCAGGAT <u>TTGTAAT</u> GATTCTATTTTTTTTACATTT <mark>CAG</mark>	СТАС	PL5

Figure 2.4. Comparison of 5' and 3' splice positions among *P. lessonae, S. tropicalis* and *X. laevis*. Numbers of exons and introns are given next to species (XL, ST, and PL). Conserved elements at 5' and 3' splice sites are shown as underlined and red colour, and putative branch point positions are underlined.

	Exon		Intro	n	Exon	
XL5	AAAG	GTATAG TGTGCAGAAAAAGTAGTTTACCTATTCCATTTTA	XL5	GTCTTA <u>CTATGAT</u> GCCTTTTTTTTTAAATTTCCTTTA <mark>TAG</mark>	ATGA	XL6
ST5	AAAG	GTATAG TGTTCAGAAATCCAGTAATTTCACATTTCCCAGT	ST5	AT <u>TTCTAAT</u> CATGTTGTTAATTTTTTTTATTAACCTTA <mark>TAG</mark>	ATTA	ST6
PL5	ACGG	GTATGC CATTAAACTTCTTGAAATTAAACAATGTTATTGC	PL5	TTTTTTTTTTTTTTTGTTTGAAACAATACTTTCTTTTGCAG	ATGC	PL6
XL6	CACT	GTAAGT GGTTCTCTATACTGATGTGATTTGTAGCATGAAA	XL6	GTGTGTCTCTGCTCTTTTC <u>TTTCAAT</u> ATATCGTATTT <mark>CAG</mark>	AAAT	XL7
ST6	CAAT	GTAAGT TGCTATCTATACTGTCTAAAATGTGTACATTTGT	ST6	GTATGTATCTGTT <u>TCTTAAT</u> TTCTATATATTGTATTC <u>CAG</u>	AAAT	ST7
PL6	CACT	GTAAGTATCTATTCCCAATCCAGGAAATATATGTAGGGGA	PL6	TTTT <u>TTTCGAT</u> GCGGTAACTGCTTCTTTCTCATTTAT <mark>CAG</mark>	AACA	PL7
XL7	GAGG	GTAAGCAATTCTGGACAATAACAACCTGATAGAACACATC	XL7	TGATTTTTTTTTTTTGTGAAACTTTCTCTCACTCTGA <mark>TAG</mark>	CTGG	XL8
ST7	GAGG	GTAAGCAATTTTGATTAAGAGACCTGATGTATCACACCAG	ST7	GGTTGAACAACCC <u>CTTTAAT</u> AATCTTTTCCATTCTGA <mark>TAG</mark>	ATGG	ST8
PL7	AAGG	GTAAGCCTGTGCACCAAACATTTTAATATCTCCTTCATCA	PL7	<u>TACGAT</u> GTTATCATGCATATGTATTTTATATCTTTGA <mark>CAG</mark>	ATGG	PL8
XL8	GAAG	GTAAAA ATATGTTTTACATGTATTACACATTATCATATAT	XL8	TAATTTAATGCATGTAATTAATTATGCATGTGTTTTA <mark>TAG</mark>	ATAC	XL9
ST8	GCAG	GTAAAG ATAATACGTTATATATGTGTTACACATTGTAATA	ST8	AGTTACATTATTATTATGAATTGTGCCTGTTTTTTATAG	ATAT	ST9
PL8	CCCA	GTAAGTATACCTTTTAATGTTTATACAATTGCCATCAAGT	PL8	TTATGTCACTTTAAACAGTCATTTGCATTTGTTTCTCCCAG	СТТТ	PL9
XL9	TGGA	GTATGT TTATTTTACACTTTAAAGCCCTGTATGTACTTAA	XL9	GGAATTCTTGATTTACATC <u>TATTAAC</u> TCCTGTCATAT <u>TAG</u>	CCTG	XL10
ST9	TGCA	GTAGGT TTTTTTTTTAACTCTTTAAAACCAAGTGTGTACT	ST9	GGCCTCAT <u>TACTAAT</u> TAGAATTCTTATCCTTTTTAT <u>TAG</u>	GCTA	ST10
PL9	TGCC	GTATGTATATGAATGTTACTGTTACAGTCCAAATAGTAAA	PL9	TTTTTTTCATTTAAAGTATTTATTTTATTTTCATTT	ССТА	PL10
Figu	re 2.4. (c	continued).				

Exon		Intron	Exon	
XL10 ATGA	GTAAGT TTTACAGATACTGATAAACAGTTGGACAGTTGGA	XL10 TACATTTACTACATAAACATTAAGTATTTTTGGTTT <u>CAG</u>	CGCT	XL11
ST10 ATGA	GTAAGT TTCACAGATATAGATATCCTATTTCTTTGTTTAT	ST10 TAGTAAATCTGCCCCTAAGTTTTTTCTTCTTTTGTTT	GCTG	ST11
PL10 TTCA	GTAAGT TAAGCGTGTAGCATTTACAATTTTGAACATTTTA	PL10 <u>CATTAAC</u> ACTTTGCTAAGAACCGGTTTTCCTTTTTCCCAG	GCTG	PL11
XL11 AGAC	GTATGTATATATTGTTTATGTTTTGCTTAGATTACTGCAT	XL11 GTGTTTATATTGAAGTTATATTTTTTTCTTCCTTTTAAAAG	CTCA	XL12
ST11 AGAC	GTATGTATATATTGTTTGTTATGTTTTGCTTGGGTTTATG	ST11 T <u>TGTTGAT</u> ATTGATGTCCTGTTTTTCACTTCCCTTTAAAAG	CTTA	ST12
PL11 AAAG	GTAAGC TCTTCTTGTTTTCTATCCTTTCAATAGATTTTTA	PL11 CACTAAACC <u>TCTCAAT</u> GTTTTCTAATATTTTACATCA <mark>CAG</mark>	CTGG	PL12
XL12 AGAA	GTAAATATCTATAATTTCCCATTTTTTGTCATTATTTTTT	XL12 ATAATTATTTTCATATATATTTTTTATTATTTGTTTA	ATTC	XL13
ST12 AGAC	GTAAATATTTTTGCCATTATTGTTTGATGTAATTCTTTGA	ST12 TGAAATAACTTAATATATATGTATTTTCTTTGGTTATAG	ATTC	ST13
PL12 TGGT	GTAAGT TTTGGAAAAACAGCCATGAGAGGCTTTAAAGATT	PL12 TT <u>TCTTAAC</u> ATTGTTTAAAATTATATAATTTTCTTTA <mark>CAG</mark>	GTTG	PL13
XL13 AGAG	GTATGGCAACACTCTTGTGTTTTTGTATTATGAAAACAA	XL13 TTTATGTCTAATCTCATGTTTTATCTTATTTATCCCACAG	AAAC	XL14
ST13 AGAG	GTATGGCAACACTCTTGTCCTTTTTGTATCATGAAAACAA	ST13 CCTACTTACTGTCTTACTTCTTATCTTATTTACCTTGCAG	AAAA	ST14
PL13 GAAG	GTATGTAATGTGACTGCTGTGCAATGTGTATAATGCCAAC	PL13 <u>TTTAAT</u> AATAAATGAATTCTGTTATTTGTGTATATTT <u>TAG</u>	AAAG	PL14
XL14 CCAG	GTAAGA TTAGATGCCTAAAATCTACAACTTTATGCAAGTG	XL14 ACATTTGTCTTAGTTTAACACATTCTCTTTTGTTTTG	ССТТ	XL15
ST14 CCAG	GTAAGGTCAAATGCCTTAAACAGATGTGTACAACATTGTA	ST14 TGCAACAACTTCTAAAT <u>TCACAAT</u> TTCTTTTGTTGTG <mark>CAG</mark>	ССТТ	ST15
PL14 ACAG	GTAAGTCTTGCAGTGTGCATGAATGGTATATGTATTTGTA	PL14 TTTGCAGTGTATAAACCAGAACTTTCCTTTCTTCAAACCAG	GCCA	PL15
Figure 2.4. (d	continued).			

2.3.1.4. Exons and Introns of the Pool Frog's Albumin Gene

Exons of pool frog's albumin gene are very short and their lengths are nearly equal to those in *S. tropicalis, X. leavis, G. gallus, R. norvegicus, M. musculus* and human albumin genes except partially translated exon1, exon14 and untranslated exon15. Their average length in water frog's gene is approximately 133 bp, and ranging from 46 bp to 224 bp. (Fig. 2.5; Appendix 2.1).



Figure 2.5. Comparison of serum albumin exon lengths among *P. lessonae, S. tropicalis, X. laevis, G. gallus, R. norvegicus, M. musculus* and human.

In contrast to this length conservation in protein coding albumin exons among distinct vertebrate species, intron lenghts are highly variable. Remarkably, in comparison with *S. tropicalis*, *X. leavis*, and other vertebrate species, introns of *P. lessonae* are extremely long. Their mean length is 2175 bp and ranging from 95 bp to 4748 bp. In contrast, the mean lengths of introns for *S. tropicalis* and *X. laevis* are 734 bp and 703 bp respectively (Fig. 2.6; Appendix 2.1). Particularly, lengths of intron 3, 4, 5, 8, 10, 11 and 13 are more than three-fold comparing with *S. tropicalis*, *X. laevis* and other vertebrate species. In this study, we documented that these introns have many truncated copies of several distinct types of DNA

transposons and retrotransposons and simple repeats. Thus, these long intronic sequences of pool frog serum albumin gene indicate that the gene is potential target of several retroelement families.



Figure 2.6. Comparison of serum albumin intron lengths among *P. lessonae, S. tropicalis, X. laevis, G. gallus, R. norvegicus, M. musculus* and human.

2.3.2. Drivers of Serum Albumin Gene Evolution

Eukaryotic genomes, which are usually larger than prokaryotic genomes, contain distinct types of repeat families. These are considered an important sources of genomic rearrangements. Repeats include interspersed repeats (transposable elements, segmental duplications, and processed pseudogenes) and tandem repeats (microsatellites and minisatellites) (Tang, 2007). Transposable elements particulary have a considerable impact on genome size, genomic variation in both non-coding and coding parts, adaptation, and shaping the eukaryotic genome. These mobile repetitive elements form two classes: i) LTR (long terminal repeat) or retrovirus and Non-LTR retroelements, both mobilized by RNA intermediates

ii) DNA transposons, mobilized by DNA intermediates. Both groups have autonomous and nonautonomous elements (Kazazian, 2004; Chenais et al., 2012; Rebollo et al., 2012).

The current study showed that repetitive elements are also important drivers of the evolution of the serum albumin gene because truncated elements of several distinct families of DNA transposons, non-LTR and LTR retroelements, and tandem repeats and simple repeats were found upstream of the 5' UTR, downstream of the 3' UTR and within individual albumin introns (Table 2.1). Truncated DNA transposons come from the Tc1Mar-Tc1, hAT-Charlie, RC/Helitron, hAT-Tip100 families, and truncated non-LTR elements are grouped within the LINE/CR1, LINE/L1-Tx1 and LINE/Penelope families. A single truncated LTR element, belonging to the Gypsy family was found. The prominent characteristics of these truncated elements and their corresponding families are explained in detail below and in Table 2.1.

Class	Family	Element	Program	Start	Stop	Length (bp)	Insertion Site
		Tc1-10_Xt-1	RM	7444	7233	212	upstream of the 5'UTR
		Tc1-10_Xt-2	RM	6878	6477	402	upstream of the 5'UTR
		TC1_RT-3	RM	2851	2667	185	upstream of the 5'UTR
		TC1_PP-4	RM	2683	2549	135	upstream of the 5'UTR
		Tc1-1Ory-5	RM	2423	817	1607	upstream of the 5'UTR
	TcMar-Tc1	TC1_FR3-6	RM	5555	5922	368	intron 4
		TC1_FR3-7	RM	5907	6293	387	intron 4
s		TC1DR1-8	RM	15364	16365	1002	intron 8
son		TC1-13_Xt-9	RM	21772	21958	187	intron 11
ods		TC1-13_Xt-10	RM	24984	25579	596	intron 11
ans		TC1DR1-11	RM	47511	48454	944	downstream of the 3'UTR
A tr		hAT-2_TS-1	RM	2417	2497	81	intron 3
N		hAT-2_ET-2	RM	17880	17965	86	intron 10
	hAT Charlia	hAT-2_ET-3	RM	18007	18167	161	intron 10
		hAT-2_TS-4	RM	18724	18821	98	intron 10
		hAT-2_OG-5	RM	18818	18892	75	intron 10
		hAT-2_TS-6	RM	18882	18988	107	intron 10
		Helitron-1_DR-1	RM	15120	15223	104	intron 8
	DC/Holitron	Helitron-1_DR-2	RM	16581	16668	88	intron 8
	KC/Helltroll	Helitron-1_DR-3	RM	47172	47262	91	downstream of the 3'UTR
		Helitron-1_DR-4	RM	48617	48706	90	downstream of the 3'UTR

Table 2.1. Classification of truncated repetetive elements obtained from Repeat Masker (RM) and TRF programs' searches and length and insertion sites of these elements within the albumin gene starting from the upstream 5' UTR to downstream of the 3' UTR according to start point (+1)

Table 2.1 (co	ntinued).
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Class	Family	Element	Program	Start	Stop	Length (bp)	Insertion Site
		hAT-32_CPB	RM	2710	2778	69	intron 3
DNA transposons	hAT-Tip100	hAT-N33_CPB	RM	54341	54387	47	downstream of the 3'UTR
		hAT-N33_CPB	RM	55040	55078	39	downstream of the 3'UTR
LTR retroelements	Gypsy	Gypsy-24-1_XT	RM	36356	36646	291	downstream of the 3'UTR
		CR1_1a_Xt-1	RM	281	346	66	intron 1
		CR1-2_XT-2	RM	4879	5006	128	intron 4
Non I TD	LINE/CR1	CR1_1a_Xt-3	RM	43276	43336	61	downstream of the 3'UTR
NUII-LIK retrolements		CR1-2_XT-2-4	RM	43376	45276	1901	downstream of the 3'UTR
retrotements		CR1-2_XT-7	RM	45464	46171	708	downstream of the 3'UTR
	LINE/L1-Tx1	KibiDR1	RM	14122	14219	98	intron 8
	LINE/Penelope	Penelope-5_XT	RM	50802	50951	150	downstream of the 3'UTR
		(GTTCTGTCACACT)n	TRF	7465	7441	25	upstream of the 5'UTR
		(AATA)n	RM	5877	5849	29	upstream of the 5'UTR
		(TATTT)n	RM	5347	5321	27	upstream of the 5'UTR
S		(ATCAGTGCCCATCAGGTCAGCCTC)n	TRF	2438	2499	62	intron 3
the	Simple repeat	(ATCAT)n	RM	2499	2525	27	intron 3
Ó		(CTCCCCCCAGTACAGACCTCT)n	TRF	2889	2971	83	intron 3
		(T)n	RM	9214	9235	22	intron 5
		(GGGCAAGTGGGCCCCCC)n	TRF	11144	11184	41	intron 6
		(A)n	RM	12455	12475	21	intron 7

Table 2.1. (continued)
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Class	Family	Element	Program	Start	Stop	Length (bp)	Insertion Site
Others	Simple repeat	(TATT)n	TRF	13262	13289	28	intron 8
		(AGATAA)n	RM	13446	13487	42	intron 8
		(AGTTATTTTGAGG)n	TRF	16491	16518	28	intron 8
		(TGATGGGCACTGATAGGCTGCAC)n	TRF	18722	18985	264	intron 10
		(CAAGTATAACCC)n	TRF	21657	21681	25	intron 11
		(AT)n	RM	22452	22497	46	intron 11
		(AAATAT)n	RM	23904	23937	34	intron 11
		(T)n	RM	26656	26681	26	intron 12
		(CAGTG)n	RM	27415	27440	26	intron 12
		(CATTG)n	RM	36237	36264	28	downstream of the 3'UTR
		(A)n (TRF)	TRF	39061	39089	29	downstream of the 3'UTR
		(AT)n (TRF)	TRF	40804	40907	104	downstream of the 3'UTR
		(ATATACA)n	RM	46412	46458	47	downstream of the 3'UTR
		(TATATTA)n	RM	46790	46871	82	downstream of the 3'UTR
		(CTGGTCACTCTGCATTGAAGGGGGGCACG)n	TRF	49137	49226	90	downstream of the 3'UTR
	Low complexity	GA-rich	RM	17014	17076	63	intron 8
		A-rich	RM	18644	18690	47	intron 10
		A-rich	RM	32729	32770	42	downstream of the 3'UTR
		A-rich	RM	46503	46540	38	downstream of the 3'UTR
		A-rich	RM	52475	52506	32	downstream of the 3'UTR

2.3.2.1. DNA Transposons

2.3.2.1.1. Tc1Mar-Tc1 Family

This family is widespread in both plants and animals including, invertebrates and vertebrates (Plasterk et al., 1999). The length of complete Tc1-like elements (TLE) ranges between 1 and 5 kb. They have a single transposase-encoding gene (282-345 aa), are flanked by two inverted terminal repeats (ITR), variable in their lengths (Plasterk et al., 1999). Transposase proteins usually have two main domains: a DNA-binding domain in the N-terminal region and a catalytic domain in the carboxy-terminal region region; these are separated by a nuclear localization signal (NLS). The DNA binding domain contains one or two helixturn-helix motif, which are responsible for recognition and binding of ITRs; the catalytic domain includes DDD or DDE motifs, which carry out DNA cleavege and joining reactions during transposition (Benjamin et al., 2007; Muńoz-Lopez and Garcia-Pérez, 2010). ITRs contains three functional motifs: the cleavage site, a DNA transposon binding site, and an enhancer (Benjamin et al., 2007). In the P. lessonae scaffold containing the albumin gene, 11 truncated Tc1-like elements (TLE) were found. As a result of search using the program Repeatmasker in the Repbase repeat database (Jurka et al., 2005), these were found to represent seven distinct intact TLE.

2.3.2.1.1.1. PL Tc1-10ry Element

The single copy of Tc1-1Ory is the longest (1607 bp) Tc1-like element (TLE) found in the albumin gene. To obtain a consensus sequence for this elements, the reference sequence of Tc1-1Ory (Pocwierz-Kotus et al., 2007) from the Repbase database was blasted against the *P. lessonae* genome; 1366 hits were returned. 230 of these hits longer than 600 bp were aligned to derive a consensus sequence named as PL Tc1-1Ory. The overall structure of the element resembles Tc1-like elements in fish and amphibians (Leaver, 2001): it has an open reading frame
(ORF) encoding an intact transposase with 339 aa, which is flanked by nearly identical inverted terminal repeats of 219 bp. Each ITR contains 22 bp direct repeats. PL Tc1-1Ory thus belongs to the inverted repeat/direct repeat (IR/DR) group of Tc1-like elements described by Izswak et al. (1995). In the region between the left ITR and the transposase initiation codon, two eukaryotic promoter motifs, a CCAAT box (binds to the transcription factor NFY) (Mantovani, 1998) and a TATAA box (Bucher, 1990) are recognized. In the right ITR region, a putative poly(A) signal (PAS) and iron responsive element (IRE) were found as a UTR scan (Fig 2.7). Both the nucleotide sequence and the aa sequence of the transposase aa are nearly identical (99% identity) to the element in Atlantic salmon (Salmo salar; positions 120142 to 121747 in GenBank GQ505859.1; Lukacs et al., 2010) with 14 nucleotides and 4 aa changes in ITR regions and ORF. Moreover, presence of several other active elements in P. lessonae genome indicates very recent horizontal transmission of this element from salmon to pool frog P. lessonae. The copy of the element found in the upstream of the 5' UTR of the albumin gene is also nearly identical to the consensus sequence of the element only with very small insertions or deletions in ITRs and only two aa changes, which is predicted that it is also naturally functional element. This element is also very similar to passport, a native Tc1 transposon in flattfish (Clark et al., 2009), a preference for insertion into genes, can be useful tool for the manipulation of vertebrate genome.



Figure 2.7. Structure of the PL Tc1-1ORY element and its consensus sequence. A) Structure of PL Tc1-1ORY and of its copy in the serum albumin gene. The white coloured rectangular boxes represent inverted terminal repeats; the gray coloured rectangle represents the transposase encoding region. B) The consensus sequence of the 1634 bp PL Tc1-1ORY element. The translated transposase is indicated below the ORF. The putative CAT box, TATA box and poly(A) signal are underlined and shown by bold letters, and the region indicating a putative IRE is marked by blue arrows. At the beginning and end of the sequence, ta repeat is duplicated genomic target site, which is used by Tc1-like transposons. Regions of the sequence representing ITRs are underlined with a black colour and DRs within them are shown by red colour letters.

B

taCAGTGCCTTGCGAAAGTATTCGGCCCCCTTGAACTTTCGACCTTTTGCCACATTTCAGGCTTCAAACAT AAAGATATAAAACTGTAATTTTTTGTGAAGAATCAACAACAAGTGGGACACAATCATGAAGTGGAACGAAAT TTATTGGATATTTCAAACTTTTTAACAAATAAAAAACTGAAAAATTGGGCGTGCAAAATTATTCAGCCCCT CAT box

TTACTTTCAGTGCAGCAAACTCTCTCCAGAAGTTCAGTGAGGATCTCTGAATGAT TATA box

GACTAATGATGATAAATAGAATCCACCTGTGTGTAATCAAGTCTCCG<u>TATAAAT</u>GCACCTGCACTGTGATAG TCTCAGAGGTCCGTTTAAAGCGCAGAGAGCATCatgaagaacaaggaacacaccaggcaggtccgagatact M K N K E H T R Q V R D T

gttgtggagaagtttaaagccggatttggatacaaaaagatttcccaagctttaaacatcccaaggagcact V V E K F K A G F G Y K K I S Q A L N I P R S T gtgcaagcgataatattgaaatggaaggagtatcagaccactgcaaatctacgaagacctggccgtccctct V Q A I I L K W K E Y Q T T A N L R R P G R P S aaactttcagctcatacaaggagaagactgatcagagatgcagccaagaggcccatgatcactctggatgaa K L S A H T R R R L I R D A A K R P M I T L D E ctgcagagatctacagctgaggggggagactctgtccataggacaacaatcagtcgtatactgcacaatct L Q R S T A E V G D S V H R T T I S R I L H K S ggcctttatggaagaggtggcaagaagaccatttctaaagatatccataaaagtgttgtttaaagttt

G L Y G R V A R R K P F L K D I H K K C C L K F gccacaagccacctgggagacacaccaaacatgtggaagaaggtgctctggtcagatgaaaccaaaatcgaa A T S H L G D T P N M W K K V L W S D E T K I E ctttttggcaacaatgcaaaacgttatgtttggcgtaaaagcaacacagctcatcaccctgaacaaccatc L F G N N A K R Y V W R K S N T A H H P E H T I cccactgtcaaacatggtggtggcagcatcatggtttgggcctgcttttcttcagcagggacagggaagatg PTVKHGGGSIMVWACFSSAGTGKM gttaaaattgatgggaagatggatggagccaaatacaggaccattctggaagaaaacctgatggagtctgca V K I D G K M D G A K Y R T I L E E N L M E S A aaagacctgagactgggacggagatttgtcttccaacaagacaatgatccaaaacataaagcaaaatctaca K D L R L G R R F V F Q Q D N D P K H K A K S T atggaatggttcacaaataaacatatccaggtgttagaatggccaagtcaaagtccagacctgaatccaatc MEWFTNKHIQVLEWPSQSPDLNPI ENLWKELKTAVHKRSPSNLTELEL ${\tt ttttgcaaggaggaatgggcaaaaatttcagtctctcgatgtgcaaaactgatagagacataccccaagcga}$ F C K E E W A K I S V S R C A K L I E T Y P K R cttacagctgtaatcgcagcaaaaggtggcgctacaaagtattaaCT<u>TAAGGGGGCTGAATAA</u>TTTTGCACG L T A V I A A K G G A T K Y * Poly(A) Signal

CCCAATTTTTCAGTTTTTATTTGTTAAAAAAGTTTGAAATATCCAATAAATTTCGTTCCACTTCATGATTG ↓ IRE TGTCCCACTTGTTGTTGATTCTTCACAAAAAATTACAGTTTTATATCTTTATGTTTGAAGCCTGAAATGTGG ↓ CAAAAGGTCGAAAAGTTCAAGGGGGCCGAATACTTTCGCAAGGCACTGta 1634 bp

Figure 2.7. (continued).

2.3.2.1.1.2. PL Tc1-10Xt Element

This element has two truncated copies (212 and 402 bp) upstream of the 5' UTR of the albumin gene. To obtain a consensus sequence of this element, the reference sequence of the Tc1-10Xt (Eagle) was blasted against the *P. lessonae*

genome; the search returned 628 hits. Because the consensus element is very short, particulary at the 5' region, all copies of the element from 50 bp to 1346 bp were used to derive a consensus sequence of the element, named as PL Tc1-10Xt. Although it has an intact transposase open reading frame (339 aa), both the left ITR and the transcriptional regulator region (between the left ITR and the transposase initiation codon) contain deletions more than 50 bp. No copy of a PL Tc1-10Xt element intact for these regions was found in the *P. lessonae* genome. Thus, the length of the left ITR is 125 bp. In contrast, the ORF flanking the right ITR is 216 bp in long, with 12 bp imperfect DRs. In the transcriptional regulatory region, only the TATA box is identified; and in the 3' UTR, only the IRE (Fig 2.8). Both complete consensus nucleotide sequence of the element and its transposase coding ORF and regions show the highest similarity to Tc1-10Xt (Eagle) element (Silurana tropicalis - consensus sequence from Repbase, Pocwierz-Kotus et al. 2007). Similarity between the aa sequences of the transposases is 91% (309/339). This low similarity even in transposase coding ORFs and many in frame shifts and stop codons in ORFs in several copies of the element in *P. lessonae* genome, and a great deletion in 5' UTR and many small insertions and deletions in both the 5' and 3' UTR indicate ancient transmission of these elements from S. tropicalis to P. lessonae or they have inherited from the common ancestor. Accumulations of mutations through time could make it as an inactive element in P. lessonae genome. Two truncated copies of the element were found upstream of the 5' UTR of the albumin gene. The longer copy extends from the end of the ORF to the 3' end of the right ITR whereas short copy covers a very small region within the ORF.



Figure 2.8. Structure of PL Tc1-10Xt element and its consensus sequence. A) Structure of the PL Tc1-10Xt element and its two copies in the serum albumin gene. White coloured rectangles indicate inverted terminal repeats; and gray coloured rectangles represent transposase encoding region. B) The consensus sequence of the 1612 bp PL Tc1-10Xt element. The transposase translation is indicated below the ORF. The putative TATA box is underlined and shown by bold letters, and the region of the putative IRE is marked by blue arrows. At the beginning and end of the sequence, ta repeat is duplicated genomic target site, which is used by Tc1-like transposons. Regions of the sequence representing ITRs are underlined with a black colour and DRs within them are shown by red coloured letters.

B

$\underline{\texttt{taCAGTGCCTTGAAAAAGTATTCATACCCCTTGAAATTTTCCACATTTTGTCATGTTACAACCAAAAACGTA}$
AATGTATTTTATTGGGATTTTATGTGATAGACCAACACAAAGTGG
ACAAATAAATAAATAAATAAATAAAAAAAAAAAA
TTTTAAATAGAGTCCACCTG

TATA box

TGTGTAATTTAATCTCAG<u>TATAAAT</u>ACAGCTGTTCTGTGAAGCCCTCAGAGGTTTGTTAGAGAACCTTAGTG AACAAACAGCATCatgaaggccaaggaacacaccaggacaggtcagggataaagttgtggagaagtttaaagca M K A K E H T R Q V R D K V V E K F K A gggttaggttataaaaaaatatcccaagctttgaacatctcacggagcactgttcaatccatcatcaaaaa G L G Y K K I S Q A L N I S R S T V Q S I I Q K

tggaaagagtatggcacaactgcaaacctaccaagacatggccgtccacctaaactgacaggccgggcaagg W K E Y G T T A N L P R H G R P P K L T G R A R agagcattaatcagagaagcagccaagaggcccatggtaactctggaggaggctgcagagatccacagctcag R A L I R E A A K R P M V T L E E L Q R S T A Q gtgggagaatctgtccacaggacaactattagtcgtgcactccacaatctggcctttatggaagagtggca V G E S V H R T T I S R A L H K S G L Y G R V A agaagaaagccattgttgaaagaaagccatagaagtcccgtttgcagttgcagtggagagccatgtgggggac R R K P L L K E S H K K S R L Q F A R S H V G D acagcaaacatgtggaagaaggtgctctggtcagatgagaccaaaattgaactttttggcctaaaagcaaaa TANMWKKVLWSDETKIELFGLKAK cgctatgtgtggcggaaaactaacactgcacatcaccctgaacaccatccccaccgtgaaacatggtggt R Y V W R K T N T A H H P E H T I P T V K H G G ggcagcatcatgttgtggggatgcttttcttcagcagggacagggaagctggtcagagttgatgggaagatg G S I M L W G C F S S A G T G K L V R V D G K M gatggagccaaatacagggcaatcttagaagaaaacctgttagagtctgcaaaagacttgagactggggcgg D G A K Y R A I L E E N L L E S A K D L R L G R aggttcaccttccagcaggacaacgaccctaaacatacagccagagctacaatggaatggtttagatcaaag R F T F Q Q D N D P K H T A R A T M E W F R S K catattcatgtgttagaatggcccagtcaaagtccagacctaaatccaattgagaatctgtggcaagacttg H I H V L E W P S Q S P D L N P I E N L W Q D L aaaattgctgttcacagacgctctccatccaatctgacagagcttgagctattttgcaaagaagaatgggca KIAVHRRSPSNLTELELFCKEEWA aaaatgtcactctctagatgtgcaaagctggtagagacatccccaaaaagacttgcagctgtaattgcagcg K M S L S R C A K L V E T S P K R L A A V I A A K G G S T K Y * TGTAAAAAATGTTGAAAACCATTTATCATTTTCCTTCCACTTCACAATTATGTGCCACTTTGTGTTGGTCTA J IRE TCACATAAAATCCCAATAAAATACATTTACGTTTTTGGTTGTAACATGACAAAATGTGGAAAATTTCAAGGG J GTATGAATACTTTTTCAAGGCACTGta 1612 bp

Figure 2.8. (continued).

2.3.2.1.1.3. PL Tc1-RT Element

This element has one very short truncated copy (185 bp) upstream of the 5' UTR of the albumin gene, which overlaps a very short region in the ORF (Fig 2.9). To obtain consensus sequence of this element, the Repbase reference sequence of Tc1-RT was blasted against the *P. lessonae* genome. The search returned 695 hits.

Because the element is considerably truncated particularly at the 5' region, all copies of the element from 100 bp to 1473 bp were used to derive a consensus sequence, named as PL Tc1-RT. Although it contains an intact transposase open reading frame (339 aa), all copies of it in the P. lessonae genome have a deletion about of 50 bp in their left ITRs. An internal DR 16 bp long follows this deletion. The predicted length of left ITR is thus about 210-220 bp. In contrast, the right ITR carries an imperfect 19 bp DRs; and the predicted length is about 210-220 bp. This uncertainity in lengths of both DRs and ITRs can be explained by accumulation of mutations through time after the insertion of original element. This element, like PL Tc1-1Ory element, belong to the inverted repeat/ direct repeat (IR/DR) group of Tc1-like elements described by Izswak et al. (1995). In the transcriptional regulatory region, only the CAT box is identified, probably because of accumulation of mutations through time (Fig 2.9). Both the complete consensus nucleotide sequence of the element and its transposase coding ORF show the greatest similarity to Tc1-RT element (Rana temporaria - consensus sequence from Repbase; Leaver, 2001). Similarity between the aa sequences of the transposases is 98% (339/333), even though the consensus sequence of PL Tc1-RT has an intact transposase ORF while the ORF of Tc1-RT has a stop codon and is thus not complete. None of copies of PL Tc1-RT in the P. lessonae genome has an intact ORF. Moreover, the long deletion in the left ITR, and several types of mutations (insertions and deletions) in both ITRs and the transcriptional regulatory region again could indicate an ancient transmission of this element from Rana temporaria to the P. lessonae genome. Or the direction of transmission not clear, probably both R. temporaria and P. lessonae have inherited this from the same common ancestor.



Figure 2.9. Structure of PL Tc1-RT element and its consensus sequence. A) Structure of type PL Tc1-RT element and its copy in the serum albumin gene. The white coloured rectangles indicate inverted terminal repeats; and gray coloured rectangles represents the transposase encoding region. B) The consensus sequence of the 1646 bp PL Tc1-RT element and positions where nucleotide sequences are translated indicated below the ORF. The putative CAT box is underlined and shown by bold letters. At the beginning and end of the sequence, ta repeat is duplicated genomic target site, which is used by Tc1-like transposons. Regions of the sequence representing ITRs are underlined with a black colour and DRs within them are shown by red colour letters.

B

 taCAGTGCCTTGCAAAAGTATTCACCCCCTTGACTTTTGCTACCTATTTTGTTACATTACAGCCTTAAGTTCAA

 TGTTTTGTTAATCTGAATTTTATGTGATGGATGGATCAGAACACAATAGTCTAAGTTGGTGAAGT

 ------CTGAAAATTGGCATGTGCGTATGTATTCACCC

CAT box

<u>CCTT</u>TGTTATGAAGCCCATAAAAAGCTCTGGTGCAA<u>**CCAAT</u>TACCTTCAGAAGTCACATAATTAGTGAAATG</u> ATGTCCACCTGTGTGCAATCTAAGTGTCACATGATCTGTCATTACATATACACACCCTTTTTTGAAAGGCCCC AGAGGCTGCAACACCTAAGCAAGAGGCACCACTAACCAAACACTGCCatgaagaccaaggaactctccaaa M K T K E L S K</u>**

caagtaagggacaatgttgttgagaagtacaagtcagggttaggttataaaaaaatatccaaatctttgatg Q V R D N V V E K Y K S G L G Y K K I S K S L M atccccaggagcaccatcaaatctatcataaccaaatggaaagaacatggcacaacagcaaacctgccaaga I P R S T I K S I I T K W K E H G T T A N L P R gacggccgcccaccaaaactcacggaccgggcaaggagggcattaatcagagaggcagcacaagaacctaag D G R P P K L T D R A R R A L I R E A A Q R P K gtaaccctggaggagctgcagagttccacagcagagactggagtatctgtacataggacgacaataagccgt V T L E E L Q S S T A E T G V S V H R T T I S R acgctccatagagttgggctttatggcagagtggccagaagaaagccattacttcagcaaaaaccaaaatg T L H R V G L Y G R V A R R K P L L S A K N K M gcacgttttgagtttgcgaaaaggcatgtgggagactcccaaaatgtatggaggaaggtgctctggtctgat A R F E F A K R H V G D S Q N V W R K V L W S D gagactaaaattgaacttttcggccatcaaagaaaacgctatgtctggcgcaaacccaacacatcacatcac E T K I E L F G H Q R K R Y V W R K P N T S H H ccaaagaacaccatccccacagtgaaacatggtggtggcagcatcatgctgtggggatgtttttcagcagcc P K N T I P T V K H G G G S I M L W G C F S A A G T G K L V R V E G K M D G A K Y R D I L E Q N ctgtaccactctgtgcgtgatttgaggctaggacggaggttcaccttccagcaggacaatgaccccaaacac L Y H S V R D L R L G R R F T F Q Q D N D P K H actgctaaagcaacacttgagtggtttaaggggaaacatgtaaatgtgttggaatggcctagtcaaagccca TAKATLEWFKGKHVNVLEWPSOSP gacctcaatccaatagaaaatctgtggtcagacttaaagattgctgttcacaagcgcaaaccatccaacttg D L N P I E N L W S D L K I A V H K R K P S N L aaggagctggagcagttttgcaaggaggaatgggcaaaaatcccagtggtaagatgtggcaagctcatagag K E L E Q F C K E E W A K I P V V R C G K L I E acttatccaaagcgacttggagctgtgattgccgcaaaaggtggctctacaaagtattgaCTTTAGGGGGGGT TYPKRLGAVIAAKGGSTKY* AGGTTGTGAGGCAACAAAACACGAAAAATGCCAA<mark>GGGGGTGAATACTTTTGCAAGGCA</mark>CTGta 1646bp

Figure 2.9. (continued).

2.3.2.1.1.4. PL Tc1-PP Element

This element has one very short truncated copy (135 bp) upstream of the 5' UTR of the albumin gene; it overlaps a very short region in the ORF and the right ITR (Fig 2.10). To obtain a consensus sequence of this element, the reference sequence of Tc1-PP was blasted against the *P. lessonae* genome; the search returned 547 hits. Because the element is truncated at both the 5' and 3' regions, all copies of the element from 100 bp to 1283 bp were used to derive a complete

consensus sequence, named as PL Tc1-PP element. It has an intact transposase open reading frame (339 aa), but both its the left and particularly right ITRs (27 bp are present) are extremely truncated. Therefore, none of DRs is found. In the transcriptional regulatory region, two eukaryotic promoter motifs, a CCAAT box and a TATAA box are recognized (Fig 2.10). Both the consensus nucleotide sequence of the element and its transposase coding ORF and regions show the greatest similarity to the Tc1-PP element (Pleuronectes platessa - consensus sequence from Repbase database, Leaver 2007). Similarity between the aa sequences of the transposases is 87% (296/339). Extreme truncations in both ITRs and several mutations (indels) in the transposase encoding ORF indicate ancient transmission of this element from a marine flatfish species (*Pleuronectes platessa*) to P. lessonae. Because all copies of this element in the P. lessonae genome contain extremely truncated ITRs, it seems possible that the original element inherited by *P. lessonae* already had very short ITRs or they could considerably accumulate mutations through time, and remain as fossil elements that it cannot be possible to recognize them.





Figure 2.10. Structure of PL Tc1-PP element and its consensus sequence. A) Structure of the PL Tc1-PP element and its copy in the serum albumin gene. White coloured rectangles indicate inverted terminal repeats; and gray coloured rectangle represents transposase encoding region. B) The consensus sequence of the 1298 bp PL Tc1-PP element. Positions where nucleotide sequences are translated indicated below. The putative CAT box and TATA box are underlined and shown by bold letters. Regions of the sequence representing truncated ITRs are underlined with a black colour.

ATGGTTTTCAAAATTTTTTACAAATAAATATCTGAAAAGTGTGGCGTGCATTTGTATTCAGCCCCCTTTACT

Cat Box

CTGATACCCCTAACTAAAATCTAGTGGAA<u>CCAAT</u>TGCCTTCAG------AGTCACCTAATTAGT

Tata Box

AAATAGAGTCCACCTGTGTGTAATTTAATCTCAG<u>TATAAAT</u>ACAGCTGTTCTGTGAAGCCCTCAGAGGTTTG TTAGAGAACCTTAGTGAACAAACAGCATCatgaaggccaaggaacacaccagacaggtcagggataaagtt M K A K E H T R Q V R D K V gtggagaagtttaaagcagggttaggttataaaaaaatatcccaagctttgaacatctcacggagcactgtt V E K F K A G L G Y K K I S Q A L N I S R S T V caatccatcatccgaaaatggaaagagtatggcacaactgcaaacctaccaagacatggccgtccacctaaa O S I I R K W K E Y G T T A N L P R H G R P P K ctgacaggccgggcaaggaggagcattaatcagagaagcagccaagagggcccatggtaactctggaggagctg L T G R A R R A L I R E A A K R P M V T L E E L cagagatccacagctcaggtggggagaatctgtccacaggacaactattagtcgtgcactccacaaatctggc Q R S T A Q V G E S V H R T T I S R A L H K S G L Y G R V A R R K P L L K E S H K K S R L Q F A agaagccatgtgggggacacagcaaacatgtggaagaaggtgctctggtcagatgagaccaaaattgaactt R S H V G D T A N M W K K V L W S D E T K I E L tttggcctaaaagcaaaacgctatgtgggggaaaactaacactgcacatcaccctgaacaccatcccc F G L K A K R Y V W R K T N T A H H P E H T I P accgtgaaacatggtggtggcagcatcatgttgtggggatgcttttcttcagcagggacagggaagctggtc TVKHGGGSIMLWGCFSSAGTGKLV agagttgatgggaagatggatggagccaaatacagggcaatcttagaagaaaacctgttagagtctgcaaaa R V D G K M D G A K Y R A I L E E N L L E S A K gacttgagactggggcggaggttcaccttccagcaggacaacgaccctaaacatacagccagagctacaatg D L R L G R R F T F Q Q D N D P K H T A R A T M gaatggtttagatcaaagcatattcatgtgttagaatggcccagtcaaagtccagacctaaatccaattgag E W F R S K H I H V L E W P S Q S P D L N P I E

Figure 2.10. (continued).

Figure 2.10. (continued).

2.3.2.1.1.5. PL Tc1-DR1 Element

This element has two truncated copies, one in intron 8 (1002 bp) and one downstream of the 3' UTR (944 bp). Because the reference sequence of Tc1-DR1 obtained from Repbase contains only the ORF, this incomplete reference sequence was blasted against GenBank; the search yielded a significant match to a longer element (1642 bp) found in Atlantic salmon (Salmo salar - EF467298, GenBank, from 34140 to 35412, complementary strand; Yazawa et al., 2008). This longer reference sequence was blasted against the *P. lessonae* genome. The search returned 2984 hits; 352 hit sequences longer than 1000 bp were used to derive a consensus sequence. The consensus sequence of the element (PL Tc1-DR1) has an intact transposase ORF (340 aa) flanked by identical ITRs about 221 bp each. The left ITR contains 20 bp and the right ITR contains 19 bp DRs respectively. The element thus belongs to the inverted repeat/ direct repeat (IR/DR) group of Tc1-like elements described by Izswak et al. (1995). In the region between the left ITR and the transposase initiation codon, only one eukaryotic promoter motif, a CCAAT box, is recognized (Fig 2.11). Both the complete consensus nucleotide sequence of the element and its transposaseencoding ORF show the greatest similarity to the element (EF467298.1) found in Salmo salar (Yazawa et al., 2008). Similarity between aa sequences of transposases is 97% (328/340). This high level of similarity together with the presence of several copies of the element in the P. lessonae genome indicates very

recent horizontal transmission of this element from salmon to water frog, which is still mobilizing within *P. lessonae* genome. Both copies of the element in the serum albumin gene correspond to the transposase coding ORF containing insertions and deletions that result in frame shifts and stop codons (Fig 2.11A)



Figure 2.11. Structure of PL Tc1-DR1 element and its consensus sequence. A) Structure of PL Tc1-DR1 element and its copies in the serum albumin gene. The white coloured rectangles indicate inverted terminal repeats; gray coloured rectangle represents transposase encoding region. B) The consensus sequence of the 1650 bp PL Tc1-DR1 element. The transposase translation is indicated below the ORF. The putative CAT box is underlined and shown by bold letters. Regions of the sequence representing ITRs are underlined and DRs within them are shown by red colour letters.

B

CAT Box

<u>CCTAA</u>GTGAAAATGTCCAAATTGGGC<u>CCAAT</u>TAGCCATTTTCCCTCCCGGTGTCATGTGACTCGTTAGTGT</u> TACAAGGTCTCAGGTGTGAATGGGGAGCAGGTGTGTTAAATTTGGTGTTATCGCTCTCACTCTCATACTG GTCACTGGAAGTTCAACatggcacctcatggcaaagaactctctgaggatgtgaaaaaaagaattgttgct</u>

						М	Α	Ρ	Н	G	Κ	Е	I	L	S	Е	D	۷	/	К	Κ	R	Ι		V	А	
ct	aca	taa	aga	tgg	cct	agg	gcta	ata	aga	aga	ttg	cca	aa	gad	cc	tga	aad	tg	gag	ctg	gca	gca	cg	gt	ggo	caa	g
L	Н	Κ	D	G	L	G	Y	Κ	K	I	А	. I	<	т	L	K	L	-	S	С	S	Т	,	V	А	К	
ac	cat	aca	gcg	gtt	taa	cag	gga	cag	gtt	cca	ctc	aga	aa	cag	ggc	ctc	gco	at	gg	tc	gac	caa	ag	aa	gtt	gag	t
Т	I	Q	R	F	Ν	R	Т	G	S	Т	Q) [N	R	Ρ	R	ŀ	ł	G	R	Ρ	K		К	L	S	

gcacgtgctcagcgtcatatccagaggttgtctttgggaaatagacgtatgagtgctgccagcattgctgca A R A Q R H I Q R L S L G N R R M S A A S I A A E V E G V G G Q P V S A Q T I R R T L H Q I G L catggctgtcgtcccagaaggaagcctcttctaaagatgatgcacaagaaagcccgcaaacagtttgctgaa H G C R P R R K P L L K M M H K K A R K Q F A E gacaagcagactaaggacatggattactggaaccatgtcctgtggtctgatgagaccaagataaacttattt D K Q T K D M D Y W N H V L W S D E T K I N L F ggttcagatggtgtcaagcgtgtgtggcggcaaccaggtgaggagtacaaagacaagtgtgtcttgcctaca G S D G V K R V W R Q P G E E Y K D K C V L P T ${\tt gtcaagcatggtggtgggggtgtcatggtctggggctgcatgagtgctgccggcactgggggagctacagttc}$ V K H G G G S V M V W G C M S A A G T G E L Q F attgagggaaccatgaatgccaacatgtactgtgacatactgaagcagagcatgatccccttccggaga I E G T M N A N M Y C D I L K Q S M I P S L R R ctgggccgcagggcagtattccaacatgataacgaccccaaacaccacctccaagacgaccactgccttgcta L G R R A V F Q H D N D P K H T S K T T A L L aagaagctgagggtaaaggtgatggactggccaagcatgtctccagacctaaaccctattgagcatctgtgg K K L R V K V M D W P S M S P D L N P I E H L W ggcatcctcaaacggaaggtggaggagcgcaaggtctctaacatccaccagctccgtgatgtcgtcatggag G I L K R K V E E R K V S N I H Q L R D V V M E gagtggaagaggactccagtggcaacctgtgaagctctggtgaactccatgcccaagagggttaaggcagtg E W K R T P V A T C E A L V N S M P K R V K A V ctggaaaataatggtggccacacaaaatattgaCACTTTGGGCCCAATTTTGGACATTTTCACTTAGGGGTGT LENNGGHTKY* ACACTGTTATACAAGCTGTACACTCACTACTTTACATTGTAGCAAAGTGTCATTTCTTCAGTGTTGTCACAT GAAAAGATATAAAAAATATTTACAAAAATGTGAGGGGTGTACTCACTTTTGTGAGATACTGTATGT

1650 bp

Figure 2.11. (continued).

62

2.3.2.1.1.6. PL Tc1-FR3 Element

In the intron 4 of the serum albumin gene contains two truncated copies (368 and 387 bp) of this element. To obtain a consensus sequence of this element, the reference sequence of the Tc1-FR3 from Repbase database (Smith, 2002) were blasted against the P. lessonae genome; the search returned 1569 hits; 384 copies of the element larger than 400 were alinged to derive a consensus sequence. The consensus sequence of the element (PL Tc1-FR3) has an intact transposase ORF (343 aa), which is flanked by short imperfect ITRs (29 bp) that contain very short DRs (5 bp). In the transcriptional regulatory region, the CAT box and TATA box are identified (Fig 2.12). Both the complete consensus nucleotide sequence of the element and its transposase-encoding ORF show the greatest similarity to the element found in Takifugu rupripes (Fugu - assembly HE591539, from 5007707 to 5008942, complementary strand). Similarity between aa sequences of the transposases is 84 % (287/343). This low similarity in the entire element, many in frame shifts and stop codons in ORFs and long truncations both the 5' and 3' UTR in several copies of the element in the P. lessonae genome, indicate ancient transmission of this element from fish to water frog. Two truncated copies of the element found in the intron 4 of the albumin gene overlap with the last region of the ORF (Fig 2.12).



Figure 2.12. Structure of PL Tc1-FR3 element and its consensus sequence. A) Structure of PL Tc1-FR3 type element and its copies in the serum albumin gene. The white coloured rectangules indicate inverted terminal repeats and gray coloured rectangle represents the transposase encoding region. B) The consensus sequence of the 1248 bp PL Tc1-FR3. The transposase translation is indicated below the ORF. The putative CAT box and TATA box are underlined and shown by bold letters. Regions of the sequence representing ITRs are underlined and DRs within them are shown by red colour letters.

B

																		C	AT I	Box			
CACTATATTGCCAAAAGTATTGGGACACCCCCCCAAATCATTGGATTCAGGTGTTCCACGGCCCCCCCC																							
	Tata Box																						
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CA	LALAGGIG <u>IAIAAAA</u> ILAAGLALLIAGGLAIGLAGALIGLIILIALAAALAIIIGIGAAAGAATgggTCgC																						
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> +	- ++c	c++	act	act			tcc		atc	2201	Fa+1	- 2 01	Faat		- at a			-+		7 - 2	act		2262
aı	atttccttgctactaaatattccacggtcaactgttagtggtattataacaaagtggaagcaactgggaaca																						
Ι	S	L	L	L	Ν	Ι	Ρ	R	S	Т	V	S	G	Ι	Ι	Т	К	W	Κ	Q	L	G	Т
ac	agca	aac	tca	gcca	acga	aag	tgg	tag	gcca	acg	taaa	aat	gaca	agag	gcgg	gggt	ca	gcgo	cat	gct	gaa	gcg	caca
т	А	т	Q	Р	R	S	G	R	Р	R	К	М	т	Е	R	G	Q	R	М	L	К	R	т
gt	gcg	cag	aag	tcg	ccaa	act	gtc	tgca	aga	gtca	aata	agci	taca	agad	ct	cca	aact	ttc	gtg	tgg	cct	tca	gatt
v	R	R	s	R	0	L	s	A	E	s	I	A	т	D	L	0	т	s	С	G	L	0	I
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ag	Caci	aac	aac	agu	grg	Lago	aga	gct	LCa	Lgg	aat	388		LCa	LSR	-cg	ageo	age	LgC	acc	Caa	gcc	ιιαι
S	Т	Т	Т	V	R	R	Е	L	Н	G	М	G	F	Н	G	R	А	А	А	S	К	Ρ	Y
at	cac	caa	gtg	caa	tgca	aaa	gcg	tcg	gat	gca	gtgg	gtg	taaa	agca	acgo	cg	cca	ctg	gac [.]	tct	aga	gca	gtgg
Ι	т	к	С	Ν	А	К	R	R	М	Q	W	С	К	А	R	R	Н	W	т	L	Е	Q	W
ag	agacgtgttctctggagtgacgaatcacgcttctctgtctg																						
D	D	v		ы	c	п	E	c	D	E	c	v	ы	0	c	п	c	D	v	ы	v	ы	D
N	N	v	L	W	J	U	L	ر	N	1	J	v	VV	ų	J	U	u	N	v	W	v	W	N
tt	ttgccaggagaacggtacttgcctgactgcattgtgccaagtgtaaagtttggtggaggggggattatggtg																						

L P G E R Y L P D C I V P S V K F G G G G I M V tggggttgttttttcaggggttgggcttggccccttagttccagtgaagggaactcttaaggcgtcagcatac W G C F S G V G L G P L V P V K G T L K A S A Y caagacattttggacaatttcatgctcccaactttgtgggaacagtttggggatggccccttcctgttccaa Q D I L D N F M L P T L W E Q F G D G P F L F Q catgactgcgcaccagtgcacaaagcaaggtccataaagacatggatgagcgagtttggggtggaggaactt H D C A P V H K A R S I K T W M S E F G V E E L gactggcctgcacagagtcctgacctcaacccgatagaacacctttgggatgaattagagcggagactgcga D W P A Q S P D L N P I E H L W D E L E R R L R gccaggccttctcgtccaacatcagtgcctgacctcacaaatgcgcttctggaagaatggtcaaacattccc R P S R P T S V P D L T N A L L E E W S N I P atagacacactcctaaaccttgtggacagccttcccagaagagttgaagctgttatagctgcaaagggtggg I D T L L N L V D S L P R R V E A V I A A K G G РТОУ* TCCCAATACTTTTGGCAATATAGTG 1248bp

Figure 2.12. (continued).

2.3.2.1.1.7 Structural Characteristics of ITR and Transposase Genes of P. lessonae consensus TLEs

The characteristic of *P. lessonae* ITRs and transposase genes are summarized in Table 2.2. All ITRs are well conserved (85-100% identity), varying from 29 bp (PL Tc1-FR3) to 221 bp (PL Tc1-DR1). Like other vertebrate TLEs such as PPTN, RTTN, SSTN (Leaver, 2001) and Frog Prince transposons (Miskey et al., 2003), they have DRs in the extremities of their ITRs, that act as binding site for the transposase. As comparing to PL Tc1-DR1 and PL Tc1-1Ory, PL Tc1-FR3 ITRs has very short DRs (5 bp). In addition, PL Tc1-1Ory, PL Tc1-Xt and PL Tc1-RT contain a highly conserved motif "CAGTG" at the 5' end, and "GTCAC" at the 3' end, of ITRs (Brezinsky et al., 1990). The size of 5' UTRs ranges from 72 bp (PL Tc1-FR3) to 187 bp (PL Tc1-RT). Except PL Tc1-DR1 and PL Tc1-FR3,

they all have a TA duplication at the beginning and end of the element, a sign of a Tc1-mariner insertion event. In all PL elements, the transposase ORF, which range from 339aa to 343 aa and have a highly conserved [D,D(34)E] motif in the catalytic domain, is uninterrupted and intact.

Name			PL Tc1-1Ory	PL Tc1-10Xt	PL Tc1-RT	PL Tc1-PP	PL Tc1-DR1	PL Tc1-FR3	
Size (bp)			1634	1612	1646	1298	1650	1248	
	ITR size	Left	219	125*	189*	67*	221	29	
	(bp)	Right	219	216	192*	27*	221	29	
ITR	% identity ITR		99	99	85	*	100	100	
III	DR size (bp)		22	12	16-19	*	19-20	5	
	CAGTG motif		Yes	Yes	Yes	*	No	No	
sase gene	ORF quality		intact 339 aa	intact 339 aa	intact 339 aa	intact 339 aa	intact 340 aa	intact 343 aa	
Transpo	5' UTR size (bp)		174	144*	187	164	156	72	
TA duplica.			Yes	Yes	Yes	*	No	No	

Table 2.2. Main features of *P. lessonae* consensus TLEs.

2.3.2.1.2. hAT/Charlie Family

DNA transposons of the hAT superfamily are widespread in fungi, plants and animals (Calvi et al., 1991; Rubin et al., 2001), including such vertebrates such as frogs, lizards, snakes and mammalians (Pace et al., 2008; Novick et al., 2010). Recently, the group has been subdivided into two distinct subfamilies (Ac and Buster; Arensburger et al., 2011). Elements are generally 2.5 to 5 kb long. An 8 bp target site duplication (TSD) is formed during transposition. The ORF encoding transposase is flanked by two short ITRs (5-27 bp), which share a low sequence similarity in their consensus sequence (Kempken and Windhofer, 2001; Rubin et al., 2001). According to Hickman et al. (2005) and Arensburger et al. (2011) hAT transposon families contain four domain in their transposase: an N-terminal domain carrying a BED zinc finger; a DNA-binding domain involved in oligomerization; a catalytic domain having the first two carboxylates of the catalytic triad (DD); and a long insertion with several α -helices that ends with a glutamate residue (E) in close proximity to the catalytic domain. Lastly, the catalytic triad (DDE) is completed (Hickman et al., 2005). In the *P. lessonae* scaffold, six truncated hAT-like elements (HLE), representing three distinct HLEs were identified.

2.3.2.1.2.1. PL hAT-2-TS Element

This element has three very short truncated copies (81, 98 and 107 bp) in introns 3 and 10 of the albumin gene; each overlaps a very short region in the transcriptional regulatory region (Fig 2.13). To obtain consensus sequence of this element, the reference sequence of the hAT-2-TS from the Repbase database (Novick et al., 2010) was blasted against the *P. lessonae* genome, which returned 943 hits. Because the consensus element is considerably truncated in both the UTR and ORF parts, particularly at the 5' end, all copies of the element from 56 bp to 1580 bp were used to derive a complete consensus sequence, named as PL hAT-2-TS element. Although an intact transposase coding ORF (601 aa) is present, there is very long deletion (469 bp) in the 5' UTR region; it is not found any copies of the element in the complete *P. lessonae* genome. It contains very short ITRs about 16 bp long. In the transcriptional regulatory region, only the TATA box is identified as a result of online search for promoter and functional

motifs using NsiteM (http://linux1.softberry.com/berry.phtml) (Solovyev et al., 2010). Both complete consensus nucleotide sequence of the element and its transposase coding ORF and regions show the highest similarity to element in the tarsier (*Tarsius syrichta* - consensus sequence from Repbase; Novick et al., 2010). Similarity between aa sequences of transposases is 95% (570/601). None of the copies of PL hAT-2-TS element in the *P. lessonae* genome has an intact ORF. Moreover, the long deletion in the 5' UTR, and several other insertions and deletions in both UTRs, the transcriptional regulatory region, and ORF indicate very ancient transmission of this element to *P. lessonae* genome. It is not clear how this element can invade from a mammalian species to amphibian, however, recently it was indicated that parallel transfer in tetrapod genomes are very common which could be explained through the way of prey to predator transmission (Novick et al., 2010). Or much more likely some form of this element was present in the very ancient common ancestor of tarsiers and frogs, and has been inherited by both (with changes, mostly deleterious).



Figure 2.13. Structure of the PL hAT-2-TS element and its consensus sequence. A) Structure of PL hAT-2-TS element and its copies in the serum albumin gene with black coloured triangle indicating inverted terminal repeats and gray coloured rectangles representing the transposase encoding region. B) The consensus sequence of the 3538 bp PL hAT-2-TS element and with translations of the nucleotide sequences indicated below. The putative TATA box is underlined and shown by bold letters. Regions of the sequence representing ITRs are underlined with black.

Tata Box

M M S K K R K I D S E C R V F K E Q W T Y gattactttttcacgcagtacaaagaaagagctgtttgtctgatatgccagaatatagtgtctgtgttcaag D Y F F T Q Y K E R A V C L I C Q N I V S V F K gaatacaatttgcgtcgacactacaaaactcaacataaagataaatatgattgtttggtcggacaagtgaga E Y N L R R H Y K T Q H K D K Y D C L V G Q V R

Figure 2.13. (continued).

B

aaagaaaaaatattaaaactgaaaaatacattgacaactcagcaaaatacttttgtgaagcaaaagcagcta K E K I L K L K N T L T T Q Q N T F V K Q K Q L aatatatcatcattgcgagcaagttttcaagttgccaagctcatagcgtgcactggcagaccattcgtagag N I S S L R A S F O V A K L I A C T G R P F V E G E F V K E C L L S V A K E M C P E K A D L F S acagtgagtctttcaggacctacaattacacgaagaattgaagaaatgggagaaaatttgcatctgcatttg T V S L S G P T I T R R I E E M G E N L H L H L ONSSKKLCYFSLALDESNDVRDSA caacttctaattttcattcgtgggacaaatgaatatttcgaagtcacagaggagcttgctgcactgaaaagc Q L L I F I R G T N E Y F E V T E E L A A L K S I K G T T T G E D I Y E K V C Q T M N D L E L D tgggctaaactagccagtgtgacaactgatggtgctcctagcatggtggggtctatgaaaggagtggttgcaW A K L A S V T T D G A P S M V G S M K G V V A cgcattaaaaaagagatggacaaacacaaccattcacatccaatagccatacactgcctcatccaacaa R I K K E M D K H N H S H P I A I H C L I H Q Q gcactgtgttgtaaatcattgaagttggactctgtcatgaaaattgtggtatcttgtgttaacttcattagaA L C C K S L K L D S V M K I V V S C V N F I R A H A L N H R Q F Q E F L S E L N V A Y E D I L Y H T E V R W L S R G R V L K R F Y D L L P Q V tctgcttttatgctttcgaaaaacaaagaagtaccagagctcaaagatgcagaatggaaatggcaccttgcc S A F M L S K N K E V P E L K D A E W K W H L A tttctgacagatgtaacagagctactcaacagtttcaatgttcaacttcaaggaaagggaagctcatctgt F L T D V T E L L N S F N V Q L Q G K G K L I C gatatgcattcacatgtgaaagcatttcaagtaaaattagacctcctcattaaacaagtgaaggaaaaac D M H S H V K A F O V K L D L L I K O V K E E N F C H L P T T Q N L L A E K P A V A F P N K T C

Figure 2.13. (continued).

gtggatttactagaaattttgcaaaaggagtttcaatttagatttaaagagcttcatctccatgaacaggac V D L L E I L O K E F O F R F K E L H L H E O D atacagcttttccggaacccattttctgttgacattgaaattgttgatccgatttaccaaatggaattggct I Q L F R N P F S V D I E I V D P I Y Q M E L A gaactacagaattgtgactctctgaaagacgcattcaaatcaagcagccttactaatttctatgcatctctc E L Q N C D S L K D A F K S S S L T N F Y A S L ccctctgagacatatcctaatctcaggaaccatgcactcaaaattgcaaccatctttggcagcacctatgtc P S E T Y P N L R N H A L K I A T I F G S T Y V tgtgaacagacttttttcccgaatgaaacatctgaaatctccaaccagatccagactaactgatgaacacttg C E Q T F S R M K H L K S P T R S R L T D E H L catcacttgctacgactagcagtgacaaatatggaaccggatattgaccatctcattagccaaaagcaggcc H H L L R L A V T N M E P D I D H L I S Q K Q A catacttcccattgaAATACTGGTAAGTTTCTTTATTTAACTTTACTTGTTCTTCATTTTAAATATTGTATT нт S н * TGTTCCCGTTTTGTTTTTTCACTTCAAAATAAGATATGTGCAGTGTGCATAGGAATTTGTTCACAGTTTTT TTTTTTTTAAACTATAGTCCGGCCCTCCAATGGTCTGAGGGACAGTGAACTGGCCCCCTGTTTAAAAAG<u>GTT</u> TGAGGACCCCTG 3538 bp

Figure 2.13. (continued).

2.3.2.1.2.2. PL hAT-2-ET Element

This element has two very short truncated copies (86 and 161 bp long) in intron 10 of the serum albumin gene, which overlap a very short region in the transcriptional regulatory region (Fig 2.14). To obtain a consensus sequence of this element, the reference sequence of the hAT-2-ET from Repbase database (Novick et al., 2010) were blasted against the *P. lessonae* genome; the search returned 695 hits. Because the element is considerably truncated in both the UTRs and ORF parts, particularly at the 5' end, all copies of the element from 44 bp to 1797 bp were used to derive a complete consensus sequence (PL hAT-2-ET). Although an intact transposase coding ORF (601 aa) is present, there is very long deletion (about 188 bp)in the 5' UTR region, this is not found in any copies of the

element in the complete *P. lessonae* genome. Perhaps this deletion was present at the time of first insertion into *P. lessonae* genome. Only the right ITR, about 16 bp long, is identified on the basis of reference sequences while the left ITR is not known because of the long deletion in this region. In the transcriptional regulatory region, only the TATA box is found. Both complete consensus nucleotide sequence of the element and its transposase coding ORF and regions show the highest similarity to element in the lesser hedgehog (*Echinops telfairi* - consensus sequence from Repbase; Novick et al., 2010). Similarity between aa sequences of transposases is 93% (556/601). None of copies of PL hAT-2-ET element on *P. lessonae* genome has an intact ORF. Moreover, the transcriptional regulatory region, and the ORF indicate very ancient transmission of this element to *P. lessonae* genome. Or the element was present in the very ancient common ancestor of both species.

A



Figure 2.14. Structure of the PL hAT-2-ET element and its consensus sequence. A) Structure of PL hAT-2-ET element and its copies in the serum albumin gene with black coloured triangles indicating inverted terminal repeats and gray coloured rectangles representing transposase encoding region. B) The consensus sequence of the 2914 bp PL hAT-2-ET element and with translations of the nucleotide sequences indicated below. The putative TATA box is underlined and shown by bold letters. Region of the sequence representing ITR is underlined with black.

TAATGCGAATGGTCAGCGGTCAGTGTTAATGCTAATGGTCAGTGCTCAGTATTAATGCTAATGGTCAGCGGT CAGTATTAATGCAAATGGTCAGCGCTCAGTGTTAATGCAAATGGTCAGTGCTCAGTATTAATGCAAATGGTC AGCGCTCAGTATTAATGCAAATGGTCAGTGCTCAGTGTTAATGCAAATGGTCAGCGGTCAGTGTTAATGCAA ATGGTCAGTGCTCAGTGTTATCGCATGGGGGGCCCCAAACTGGTAATCTGCCTAGGGCCCCATGGGAACTTA ------CAGACAGCCGAGGAGTAGAAAACCCAATTTAATTGACAGTAAGTGCATTTATATTCTGATTG CTATTCAGTTGTATGATGTTGTATGTTGTGTGCTGTGTAAGCCCCGGTTCACACTGTTGCGATCTCTGAGCA GTGCGAGTTCAGCCATATGCTTGTATGGCTGAACTTGCATTAGATTCGGAAGAAAAAGGCACACGTGCCTT CTTTTTCCTGCAGTGGAATCGGATCGCATGGGTCTTCTCACCCATGCGATCCTGTGCGAGTTCAC

TatA Box

ATGTCTCTGTGTGGGCAAAGTTATTGCTGGTATATTGTTTTTGTAGCGCTG**TATATA**TATTGGTATTTTA CTAATAGCAATTTGGAATCCCTAGGAAACAatgatatcaaaaaaagaaaaattgactctgagtgcagggta M I S K K R K I D S E C R V F K E Q W T Y D Y F F T Q Y K E R A V C L I C Q aatatagtgtctgtgttcaaggaatacaatttgcgtcgacactacgaaactcaacataaagataaatatgat N I V S V F K E Y N L R R H Y E T Q H K D K Y D tgtttggtcggacaagtgagaaaagataaaatattaaaactgaaaaatggattgacaactcagcaaaatact C L V G Q V R K D K I L K L K N G L T T Q Q N T tttgtgaagcaaaagcagctaaatatatcatcattgcgagcaagttttcaagttgccaagctcatagcgtgc F V K Q K Q L N I S S L R A S F Q V A K L I A C T G R P F V E G E F V K E C L L S V A K E M C P gagaaggctgatttatttagtacagtgagtctttcaggacctacaattacacgaagaattgaagaaatggga E K A D L F S T V S L S G P T I T R R I E E M G ENLHL UNSSKKLCYFSLALDES aatgatgttcgtgattctgcacaacttctaattttcatttgtgggacaaatgaatatttcgaagtcacagag N D V R D S A Q L L I F I C G T N E Y F E V T E

Figure 2.14. (continued).

B

gagcttgctgcactgaaaagcatcaaaggaacaacaacaggagaggatatctatgaaaaagtttgccaaact E L A A L K S I K G T T T G E D I Y E K V C O T atgaatgatttggagctggactgggctaaactagccagtgtgacaactgatggtgctcctagcatggtgggg M N D L E L D W A K L A S V T T D G A P S M V G ${\tt tctatgaaaggagtggttgcacgcattaaaaaagagatggacaaacacaaccattcacatccaatagccata}$ S M K G V V A R I K K E M D K H N H S H P I A I cactgcctcatccaccaacaagcactgtgttgtaaatcattgaagttggactctgtcatgaaaattgtggta H C L I H Q Q A L C C K S L K L D S V M K I V V S C V N F I R A H A L N H R Q F Q E F L S E L N gttgcctatgaagatattctgtaccacacagaagtccgttggctgagtcgaggagagagttttgaaacgttttV A Y E D I L Y H T E V R W L S R G R V L K R F tatgacttacttccacaggtttctgcttttatgctttcgaaaaacaaagaagtaccagagctcaaagatgca Y D L L P Q V S A F M L S K N K E V P E L K D A gaatggaaatggcaccttgcctttctgacagatgtaacagagctactcaacagtttcaatgttcaacttcaa E W K W H L A F L T D V T E L L N S F N V Q L Q ggaaaggggaagctcatctgtgatatgcattcacatgtgaaagcatttcaagtaaaattagacctcctcatt G K G K L I C D M H S H V K A F Q V K L D L L I aaacaagtgaaggaggaaaacttctgccatctccccacgactcaaaaccttttggcggaaaaaccagcagtt K Q V K E E N F C H L P T T Q N L L A E K P A V gcattcccaaacaaaacatgtgtggatttactagaaattttgcaaaaggagtttcaatttagatttaaagag A F P N K T C V D L L E I L Q K E F Q F R F K E cttcatctccatgaacaggacatacagcttttccggaacccattttctgttgacattgaaattgttgatccg L H L H E Q D I Q L F R N P F S V D I E I V D P atttaccaaatggaattggctgaactacagaattgtgactctctgaaagacgcattcaaatcaagcagcctt I Y Q M E L A E L Q N C D S L K D A F K S S S L actaatttctatgcatctctccccctctgagacatatcctaatctcaggaaccatgcactcaaaattgcaaccatgcaccatgcactcaaaattgcaaccatgcactcaaaattgcaaccatgcaccatgcactcaaaattgcaaccatgcactaaaattgcaaccatgcactaaaattgcaaccatgcactaaaattgcaaccatgcactaaattgcaaccatgcactaaaattgcaaccatgcactaaaattgcaaccatgcactaaaattgcaaccatgcactaaattgcaaccatgcactaaattgcaaccatgcactaaattgcaaccatgcactaaattgcaaccatgcactaaattgcaaccatgcactaattgcaaccatgcactaaattgcaaccatgcactaaattgcaaccatgcactaaattgcaaccatgcactgcactaaattgcaaccatgcactgcaT N F Y A S L P S E T Y P N L R N H A L K I A T atctttggcagcacctatgtctgtgaacagacttttttcccgaatgaaacatctgaaatctccaaccagatcc I F G S T Y V C E O T F S R M K H L K S P T R S agactaactgatgaacacttgcatcacttgctacgactagcagtgacaaatatggaaccggacattgaccat R L T D E H L H H L L R L A V T N M E P D I D H

Figure 2.14. (continued).

Figure 2.14. (continued).

2.3.2.1.2.3. PL hAT-2-OG Element

This element has only one very short truncated copy (75 bp) in intron 10, which overlaps a very short region in the transcriptional regulatory region (Fig 2.15). To obtain a consensus sequence of this element, the reference sequence of the hAT-2-OG from Repbase database (Novick et al., 2010) was blasted against the P. lessonae genome; the search returned 514 hits. Because the element is considerably truncated in both the UTRs and ORF parts, all copies of the element from 60 bp to 1797 bp were used to derive a complete consensus sequence (PL hAT-2-OG). Although it contains an intact transposase coding ORF (601 aa), it has a deletion of about 41 bp before the translation initiation and two short deletions in the 5' UTR region. These parts are not found in any copies of the element in the P. lessonae genome. Only the right ITR (15 bp) is identified on the basis of the reference sequence. Both the complete consensus nucleotide sequence of the element and its transposase coding ORF and regions show the highest similarity to element in the greater galago (Otolemur monteiri - consensus sequence from the Repbase; Novick et al., 2010). Similarity between aa sequences of transposases is 92% (601/553). None of copies of PL hAT-2-OG element in the P. lessonae genome has an intact ORF. Additionally, several other insertions and deletions both in UTRs, the transcriptional regulatory region and ORF indicate very ancient transmission of this element to P. lessonae genome.



Figure 2.15. Structure of the PL hAT-2-OG element and its consensus sequence. A) Structure of thePL hAT-2-OG element and its copies in the serum albumin gene with Black coloured triangle indicate inverted terminal repeats; and gray coloured rectangles represent transposase encoding region. B) The consensus sequence of the 2790 bp PL hAT-2-OG element and positions where nucleotide sequences are translated indicated below the ORF. Region of the sequence representing ITR is underlined with a black colour.

B

M I S K K R K

L T T Q Q N T F V K Q K Q L N I S S L R A S F Q V A K L I A R T G R P F V E G E F V K E C L L S V A K E M C P E K A D L F S T V S L S G P T I T R R I E E M G E N L H L H L Q N S S K K L C Y F tcattggcactcgacgaaagcaatgatgttcgtgattctgcacaacttctaattttcattcgtgggacaaat S L A L D E S N D V R D S A Q L L I F I R G T N gaatatttcgaagtcacagaggagcttgctgcactgaaaagcatcaaaggaacaacaacaggagggatatc E Y F E V T E E L A A L K S I K G T T T G E D I ${\tt tatgaaaaagtttgccaaactatgaatgatttggagctggactgggctaaactagccagtgtgacaactgat}$ Y E K V C O T M N D L E L D W A K L A S V T T D ggtgctcctagcatggtggggtctatgaaaggagtggttgcacgcattaaaaaagagatggacaaacacaac G A P S M V G S M K G V V A R I K K E M D K H N cattcacatccaatagccatacactgcctcatccaccaacaagcactgtgttgtaaatcattgaagttggac H S H P I A I H C L I H Q Q A L C C K S L K L D S V M K I V I S C V N F I R A H A L N H R Q F Q gaatttctgtctgagctaaatgttgcctatgaagatattctgtaccacagaagtccgttggctgagtcga E F L S E L N V A Y E D I L Y H T E V R W L S R gggagagttttgaaacgtttttatgacttacttccacaggtttctgcttttatgctttcgaaaaaacaaagaa G R V L K R F Y D L L P Q V S A F M L S K N K E gtaccagagctcaaagatgcagaatggaaatggcaccttgcctttctgacagatgtaacagagctactcaac V P E L K D A E W K W H L A F L T D V T E L L N agtttcaatgttcaacttcaaggaaaggggaagctcatctgtgatatgcattcacatgtgaaagcatttcaa S F N V Q L Q G K G K L I C D M H S H V K A F Q gtaaaattagacctcctcattaaacaagtgaaggagaaaacttctgccatctccccacgactcaaaacctt V K L D L L I K Q V K E E N F C H L P T T Q N L L A E K P A V A F P N K T C V D L L E I L Q K E tttcaatttagatttaaagagcttcatctccatgaacaggacatacagcttttccggaacccattttctgtt F Q F R F K E L H L H E Q D I Q L F R N P F S V

Figure 2.15. (continued).

Figure 2.15. (continued).

2.3.3. Sequence Evolution and Phylogenetic Implications

The following calculations are based only on protein coding exon sequences of WPWFs. Nucleotide frequencies of exons 1-14 and the corresponding transition/transversion rate (R) were estimated under the discrete GTR+G model (5 categories, G=0.05) with MEGA6.06 (Tamura et al., 2013). All positions with less than 95 % site coverage were eliminated resulting in a final data set of 1587 positions. The estimated nucleotide frequencies were A=23.7 %, T=25.2 %, C=17.8 %, and G=22.3 %; the R-value amounted to 0.73, which was used as an input parameter in a codon-based z-test to test whether positive selection is operating on the gene. The z-test was performed for all sequences (scope=overall average) on the basis of the modified Nei-Gojobori method (Nei and Kumar, 2000) as implicated in MEGA6.06 under the alternative hypothesis $d_N > d_S$ (positive selection) where d_N is the number of non-synonymous substitutions per non-synonymous site and d_S is the number of synonymous substitutions per

synonymous site. As a result, the null hypothesis $(d_N = d_S)$ could not be rejected (p=0.161), i.e. sequence evolution of protein coding exons is expected to follow a neutral model of sequence evolution.

Based on the full sequence length of 1815 bp (from the start to the first stop codon), 58 variable sites (3.2%) were found; 34 (1.9%) were parsimony informative. The corresponding values for the 605 aa are 43 (7.1%) and 27 (4.5%), respectively. Pairwise sequence comparisons revealed 8 to 39 nucleotide and 7 to 33 aa differences (Appendices 2.3 and 2.4).

ML analysis of albumin exons clearly supports the sister group relationship of *P. lessonae* and *P. bergeri* and indicates a close relationships of *P. ridibundus* and *P. cf. bedriagae* from Anatolia as already suggested by previous studies (e.g. Plötner et al., 2010; Akin et al., 2010b; Plötner et al., 2012). Moreover, *P. epeiroticus* is closely related to *P. ridibundus/P.* cf. *bedriagae* which coincides with the results of phylogenetic reconstructions based on the first intron of the SA gene (Plötner et al., 2009; 2012) and our mtDNA phylogenies (e.g. Plötner, 1998, Plötner and Ohst 2001; Plötner et al., 2010; Akin et al., 2010b). In contrast to former hypotheses, however, the SA exon data do not support a sister group relationship between *P. shqipericus* and *P. lessonae+P. bergeri*. Instead, *P. shqipericus* represents a distinct lineage; this corresponds to the results of protein electrophoretic investigations (Beerli et al., 1996).



Figure 2.16. Phylogenetic relationships of main western Palearctic water frog species based on nucleotide sequence of protein-coding albumin exons. Numbers on the branches show bootstrap values.

2.4. References

Akın, Ç., Bilgin C.C., Beerli, P., Westaway, R., Ohst, T., Litvinchuk, S.N., Uzzell, T., Bilgin, M., Hotz, H., Guex, G.-D., Plötner, J. (2010b) Phylogeographic patterns of genetic diversity in eastern Mediterranean water frogs were determined by geological processes and climate change in the Late Cenozoic. *J Biogeogr* 37: 211–2124.

Arensburger, P., Hice, R.H., Zhou, L., Smith, R.C., Tom, A.C., Wright, J.A., Knapp, J., O'Brochta, D.A., Craig, N.L., Atkinson, P.W. (2011) Phylogenetic and functional characterization of the hAT transposon superfamily. *Genetics* 188: 45-57.

Beerli, P., Hotz, H. & Uzzell, T. (1996) Geological dated sea barriers calibrate protein clock for Aegean water frogs. *Evolution* 50: 1676-1687.

Benjamin, B., Yves, B., Corinne, A.-G. (2007) Assembly of the Tc1 and mariner transposition initiation complexes depends on the origins of their transposase DNA binding domains. *Genetica* 130: 105-120.

Benson, G. (1999) Tandem repeats finder: a program to analyze DNA sequences. *Nucleic Acids Res* 27 (2): 573-580.

Binder, R., Horowitz, J.A., Basilion, J.P., Koeller, D.M., Klausner, R.D., Harford, J.B. (1994) Evidence that the pathway of transferrin receptor mRNA degradation involves an endonucleolytic cleavage within the 3'UTR and does not involve poly(A) shortening. *EMBO J* 13: 1969-1980.

Bowes, J.B., Snyder, K.A., Segerdell, E., Gibb, R., Jarabek, C., Noumen, E., Pollet, N., Vize, P.D. (2008) Xenbase: a Xenopus biology and genomic resource. *Nucleic Acids Res* 36(Database issue): D761-767.

Brezinsky, L., Wang, G.V., Humphreys, T., Hunt, J., (1990) The transposable element Uhu from Hawaiian Drosophila-member of the widely dispersed class of Tc1-like transposons. *Nucleic Acids Res* 18: 2053–2059.

Bucher, P. (1990) Weight matrix descriptions of four eukaryotic RNA polymerase II promoter elements derived from 502 unrelated promoter sequences. *J Mol Biol* 212: 563–578.

Calvi, B.R., Hong, T.J., Findley, S.D., Gelbart, W.M (1991) Evidence for a common evolutionary origin of inverted repeat transposons in Drosophila and plants: hobo, Activator and Tam3. *Cell* 66: 465-471.

Cereghini, S., Raymondjean, M., Carranca, A.G., Herbomel, P., Yaniv, M. (1987) Factors involved in control of tissue-specific expression of albumin gene. *Cell* 50: 627-638.

Cereghini, S., Blumenfeld, M., Yaniv, M. (1988) A liver-specific factor essential for albumin transcription differs between differentiated and dedifferentiated rat hepatoma cells. *Gene Dev* 2: 957-974.

Church, D.M., Goodstadt, L., Hillier, L.W., Zody, M.C., Goldstein, S. et al. (2009) Lineage-specific biology revealed by a finished genome assembly of the mouse. *PLoS Biol* 7(5): 1-16.

Chénais, B., Caruso, A., Hiard, S., Casse, N. (2012) The impact of transposable elements on eukaryotic genomes: from genome size increase to genetic adaptation to stressful environments. *Gene* 509: 7-15.

Clark, K.J., Carlson, D.F., Leaver, M.J., Foster, L.K., Fahrenkrug, S.C. (2009) Passport, a native Tc1 transposon from flatfish, is functionally active in vertebrate cells. *Nucleic Acids Res* 37(4): 1239-1247.

Conway, L., Wickens, M. (1985) A sequence downstream of AAUAAA is required for formation of simian virus 40 late mRNA 3' termini in frog oocytes. *Proc Natl Acad Sci* 82: 3949-3953.

Danckwardt, S., Kaufmann, I., Gentzel, M., Foerstner, K.U., Ganzert, A.S., Gehring, N.H., Neu-Yilik, G., Bork, P., Keller, W., Wilm, M., Hentze, M.W., Kulozik, A.E. (2007) Splicing factors stimulate polyadenylation via USEs at non-canonical 3'end formation signals. *EMBO* 26: 2658-2669.

Edgar, R.C. (2004) MUSCLE: multiple sequence alignment with high accuracy and high throughput. *Nucleic Acids Res* 32(5): 1792-1797.

Fanali, G., Masi, A., Trezza, V., Marino, M., Fasano, M., Ascenzi, P. (2012)Human serum albumin: from bench to bedside. *Mol Aspects Med* 33: 209-290.

Felsenstein, J. (1985) Confidence limits on phylogenies: An approach using the bootstrap. *Evolution* 39:783-791.

Gibbs, R.A., Weinstock, G.M., Metzker, M.L., Muzny, D.M. et al., (2004) Genome sequence of the brown norway rat yields insights into mammalian evolution. *Nature* 428 (6982): 493-521.

Graber, J.H., Cantor, C.R., Mohr, S.C., Smith, T.F. (1999) In silico detection of control signals: mRNA 3'end-processing sequences in diverse species. *Proc Natl Acad Sci* 96: 14055-14060.

Goldberg, M.L. (1979) Sequence analysis of *Drosophila* histone genes. Ph.D. dissertation, Stanford University, California.

Gu, H., Gupta, J.D., Schoenberg, D.R. (1999) The poly(A)-limiting element is a conserved cis-acting sequence that regulates poly(A) tail length on nuclear premRNAs. *Pro Natl Acad Sci* 96: 8943-8948.

Gupta, J.D., Gu, H., Chernokalskaya, E., Gao, X., Schoenberg, D.R. (1998) Identification of two cis-acting elements that independently regulate the length of poly(A) on Xenopus albumin pre-mRNA. *RNA* 4: 766-776.

Hickman, A.B., Perez, Z.N., Zhou, L., Musingarimi, P., Ghirlando et al., (2005) Molecular architecture of a eukaryotic transposase. *Nat Struct Mol Biol* 12: 715-721.

Hillier, L.W., Miller, W., Birney, E., Warren, W. et al. (2004) Sequence and comparative analysis of the chicken genome provide unique perspectives on vertebrate evolution. *Nature* 432 (7018): 695-716.

Hillier, L.W., Graves, T.A., Fulton, R.S., Fulton, L.A., Pepin, K.H. et al. (2005) Generation and annotation of the DNA sequences of human chromosomes 2 and 4. *Nature* 434 (7034): 724-731.

Hu, J., Lutz, C.S., Wilusz, J., Tian, B. (2005) Bioinformatic identification of candidate cis-regulatory elements involved in human mRNA polyadenylation. *RNA* 11: 1485–1493.

Izsvak, Z., Ivics, Z. & Hackett, P. B. (1995) Characterization of a Tc1-like transposable element in zebrafish (Danio rerio). *Mol Gen Genet* 247: 312–322.

Jurka, J., Kapitonov, V.V., Pavlicek, A., Klonowski, P., Kohany, O., Walichiewicz, J. (2005) Repbase Update, a database of eukaryotic repetitive elements. *Cytogenet Genome Res* 110:462-467.
Katoh, K., Kuma, K.-I., Toh, H., Miyata, T. (2005) MAFFT version 5: improvement in accuracy of multiple sequence alignment. *Nucleic Acids Res* 33(2): 511-518.

Kazazian, H.H. (2004) Mobile elements: drivers of genome evolution. *Science* 303: 1626-1632.

Kempken, F., Windhofer, F. (2001) The hAT family: a versatile transposon group to plants, fungi, animals, and man. *Chromosoma* 110: 1-9.

Kugler, W., Wagner, U., Ryffel, G.U. (1988) Tissue specifity of liver gene expression: a common liver-specific promoter element. *Nucleic Acids Res* 16 (8): 3165-3174.

Leaver, M. (2001) A family of Tc1-like transposons from the genomes of fishes and frogs: evidence for horizontal transmission. *Gene* 271: 203-214.

Lukacs, M.F., Harstad, H., Bakke, H.G., Beetz-Sargent, M., McKinnel, L., Lubieniecki, K.P., Koop, B.F., Grimholt, U. (2010) Comprehensive analysis of MHC class I genes from the U-, S-, and Z- linaeges in Atlantic salmon. *BMC Genomics* 11: 154.

MacDonald, C.C., Wilusz, J. Shenk, T. (1994) The 64-kilodalton subunit of the CstF polyadenylation factor binds to pre-mRNAs downstream of the cleavage site and influences cleavage site location. *Mol Cell Biol* 14: 6647-6654.

Mantovani, R. (1998) A survey of 178 NF-Y binding CCAAT boxes. *Nucleic Acids Res* 26 (5): 1135-1143.

Miller, L.L., Bly, C.G., Watson, M.L., Bale, W.F. (1951) J Exp Med 95: 531.

Miller, L.L., Bale, W.F. (1954) J Exp Med 99: 125.

Miskey, C., Izsvak, Z., Plasterk, R.H.A., Ivics, Z., (2003) The Frog Prince: a reconstructed transposon from Rana pipiens with high transpositional activity in vertebrate cells. *Nucleic Acids Res* 31: 6873-6881.

Minghetti, P.P., Ruffner, D.E., Kuang, W.-J., Dennison, O.E., Hawkins, J.W., Beattie, W.-G., Dugaiczyk, A. (1986) Molecular structure of the human albumin gene is revealed by sequence within q11-22 of chromosome 4. *J Biol Chem* 261: 6747-6757.

Mount, S.M. (1982) A catalogue of splice junction sequences. *Nucleic Acids Res* 10(2): 459-472.

Munoz-Lopez, M. Garcia-Perez, J.L. (2010) DNA transposons: nature and applications in genomics. *Curr Genomics* 11: 115-128.

Murthy, K.G. Manley, J.L. (1995) The 160-kD subunit of human cleavagepolyadenylation specifity factor coordinates pre-mRNA 3'end formation. *Gene Dev* 9: 2672-2683.

Nei, M. & Kumar, S. (2000) *Molecular Evolution and Phylogenetics*. Oxford University Press, New York.

Novick, P., Smith, J., Ray, D., Boissinot, S. (2010) Independent and parallel lateral transfer of DNA transposons in tetrapod genomes. *Gene* 449: 85-94.

Pace, J.K., 2nd, Gilbert, C., Clark, M.S., Feschotte, C. (2008) Repeated horizontal transfer of a DNA transposon in mammals and other tetrapods. *Proc Natl Acad Sci USA* 105: 17023-17028.

Peters, T.Jr., Anfinsen, C.B. (1950) Production of radioactive serum albumin by liver slices. *J Biol Chem* 182: 171-179.

Plasterk, R., Izsvak, Z., Ivics, Z. (1999) Resident aliens: the Tc1/mariner superfamily of transposable elements. *Trends Genet* 15(8): 326-332.

Plötner, J. (1998) Genetic diversity in mitochondrial 12S rDNA of western Palearctic water frogs (*Anura, Ranidae*) and implications for their systematics. *J Zool Syst Evol Research* 36: 191-201.

Plötner, J. (2005) Die westpaläarktischen Wasserfrösche. Von Märtyrern der Wissenschaft zur biologischen Sensation. Laurenti Verlag, Bielefeld.

Plötner, J., Ohst, T. (2001) New hypotheses on the systemtics of the western Palearctic water frog complex (Anura, Ranidea). *Mitt Mus Natkd Berl Zool Reihe* 77 (1): 5-21.

Plötner, J., Köhler, F., Uzzell, T., Beerli, P. (2007) Molecular systematics of amphibians. In: Heatwole, H., Tyler, M. (Eds.), Amphibian Biology, vol. 7. Phylogeny and Systematics. Surrey Beatty & Sons, Chipping Norton, pp. 2672–2756.

Plötner, J., Köhler, F., Uzzell, T., Beerli, P., Schreiber, R., Guex, G.-D. & Hotz, H. (2009) Evolution of serum albumin intron-1 is shaped by a 5' truncated non-long terminal repeat retrotransposon in western Palearctic water frogs (Neobatrachia). *Mol Phylogenet Evol* 53: 784–791.

Plötner, J., Uzzell, T., Beerli, P., Akın, Ç., Bilgin, C.C., Haefeli, C., Ohst, T.,
Köhler, F., Schreiber, R., Guex, G.-D., Litvinchuk, A. N., Westaway, R.,
Reyer, H.-U. & Hotz, H. (2010) Genetic divergence and evolution of
reproductive isolation in eastern Mediterranean water frogs. Evolution in action.
Case studies in adaptive radiation and the origin of biodiversity. Special volume

from the SPP 1127 'Radiations – Genesis of Biological diversity' of the DFG (ed. by M. Glaubrecht), pp. 373–403. Springer, Heidelberg, Berlin.

Plötner, J., Baier, F., Akın, Ç., Mazepa, G., Schreiber, R., Beerli, P., Litvinchuk, S.N., Bilgin, C.C., Borkin, L., Uzzell, T. (2012) Genetic data reveal that water frogs of Cyprus (genus Pelophylax) are an endemic species of Messinian origin. *Zoosyst Evol* 88: 261-283.

Pocwierz-Kotus, A., Burzynski, A., Wenne, R. (2007) Family of Tc1-like elements from fish genomes and horizontal transfer. *Gene* 390: 243-251.

Proudfoot, N.J. (2011) Ending the message: poly(A) signals then and now. *Gene Dev* 25: 1770-1782.

Rebollo, R., Romanish, M.T., Mager, D.L. (2012) Transposable elements: an abundant and natural source of regulatory sequences for host genes. *Annu Rev Genet* 46: 21-42.

Reed, R. and Maniatis, T. (1985) Intron sequences involved in lariat formation during pre-mRNA splicing. *Cell* 41: 95-105.

Rothschild, M.A., Oratz, M., Schreiber, S.S. (1988) Serum albumin. *Hepatology* 8: 385–401.

Rubin, E., Lithwick, G., Levy, A.A. (2001) Structure and evolution of the hAT transposon superfamily. *Genetics* 158: 949-957.

Ruskin, B., A. Krainer, T. Maniatis, Green, M.R. (1984) Ex- cision of an intact intron as a novel lariat structure during pre-mRNA splicing in vitro. *Cell* 38: 317-331.

Ryffel, G.U., Kugler, W., Wagner, U., Kaling, M. (1989) Liver specific gene transcription in vitro: the promoter elements HP1 and TATA box are necessary and sufficient to generate a liver-specific promoter. *Nucleic Acids Res* 17(3): 939-953.

Sargent, T.D., Jagodzinski, L.L., Yang, M., Bonner, J. (1981a) Fine structure and evolution of the rat serum albumin gene. *Mol Cell Biol* 1: 871-883.

Sargent, T.D., Yang, M., Bonner, J. (1981b) Nucleotide sequence of cloned rat serum albumin messenger RNA. *Proc Natl Acad Sci USA* 78: 243-246.

Schorpp, M., Döbbeling, U., Wagner, U., Ryffel, G.U. (1988a) 5'-Flanking and 5'-Proximal Exon Regions of the Two Xenopus Albumin Genes: Deletion Analysis of Constitutive Promoter Function. *J Mol Biol* 199: 83-93.

Schorpp, M., Kugler, W., Wagner, U., Ryffel, U. (1988b). Hepatocyte-specific promoter element HP1 of the Xenopus albumin gene interacts with transcriptional factors of mammalian hepatocytes. *J Mol Biol* 202: 307-320.

Schwarz, G.E. (1978) Estimating the dimension of a model. *Annals of Statistics* 6(2): 461–464

Solovyev, V.V., Shahmuradov, I.A., Salamov, A.A. (2010) Identification of promoter regions and regulatory sites. *Methods Mol Biol.* 674: 57-83.

Tamura, K., Stecher, G., Peterson, D., Filipski, A., and Kumar, S. (2013) MEGA6: Molecular Evolutionary Genetics Analysis Version 6.0. *Mol Biol Evol* 30: 2725-2729.

Tang, H. (2007) Genome assembly, rearrangement, and repeats. *Chem Rev* 107: 3391-3406.

Tavaré, S. (1986) "Some Probabilistic and Statistical Problems in the Analysis of DNA Sequences". *Lectures on Mathematics in the Life Sciences (American Mathematical Society)* 17: 57–86.

Thompson, J.D., Higgins, D.G., Gibson, T.J. (1994) CLUSTAL W: improving the sensitivity of progressive multiple sequence alignment through sequence weighting, position-specific gap penalties and weight matrix choice. *Nucleic Acids Res* 22 (22): 4673-4680.

Tunner, H.G. (1979) The inheritance of morphology and electrophoretic markers from homotypic crosses of the hybridogenetic *Rana esculenta*. *Mitteilungen aus dem zoologischen Museum in Berlin,* 55 (1): 89-109.

UTRdb and UTRsite (RELEASE 2010) a collection of sequences and regulatory motifs of the untranslated regions of eukaryotic mRNAs. *Nucleic Acids Res* 38 (Database Issue): D75-D80.

Uzzell, T. (1979b) Immunological distances between the serum albumins of Rana ridibunda and Rana lessonae. *Proceedings of the Academy of Natural Sciences of Philadelphia,* 130: 1-10.

Uzzell, T. (1982) Immunological relationship of western Palearctic water frogs (Salientia: Ranidae). *Amphibia-Reptilia* 3: 135-143.

Uzzell, T., Hotz, H. (1979) Electrophoretic and morphological evidence for two forms of green frogs (Rana esculenta complex) in peninsular Italy (Amphibia, Salientia). *Mitteilungen aus dem zoologischen Museum in Berlin,* 55 (1): 13-27.

Westley, B., Wyler, T., Ryffel, G., Weber, R. (1981) *Xenopus laevis* serum albumins are encoded in two closely related genes. *Nucleic Acids Res* 9(15):3557-3574.

Yazawa, R., Cooper, G.A., Hunt, P., Beetz-Sargent, M., Robb, A.,Conrad,
M., McKinnel, L., So, S., Jantzen, S., Phillips, R.B., Davidson, W.S.,
Koop,B.F. (2008) Striking antigen recognition diversity in the Atlantic salmon Tcell receptor alpha/delta locus. *Dev Comp Immunol* 32 (3): 204-212.

Zhang, Y.-X., Lai, R., Lee, W.-H., Zhang, Y. (2005) Frog albumin is expressed in skin and characterized as a novel potent trypsin inhibitor. *Protein Sci Sep* 14(9): 2469-2477.

2.5. Appendices

Table 2.3. Comparison of SA exon and intron lengths among distinct vertebrate species. Start and stop site positions of individual exon and intron within the serum albumin gene of *P. lessonae*, the lengths of each exon and intron of the serum albumin gene for *P. lessonae*, *S. tropicalis*, *X. leavis*, *G. gallus*, *R. norvegicus*, *M. musculus* and *H. sapiens*, and the average lengths of exons and introns of the albumin gene for each species are given in this table.

P.lessonae		LENGTH (bp)							
E/ INT	START	STOP	P. lessonae	S. tropicalis	X. laevis	G. gallus	R. norvegicus	M. musculus	H. sapiens
E1	1	116	116	117	117	93	94	125	118
Int1	117	634	518	494	487	1487	704	707	709
E2	635	698	64	67	67	61	58	58	58
Int2	699	1802	1104	636	866	754	912	921	1454
E3	1803	1935	133	133	133	133	133	133	133
Int3	1936	4450	2515	585	717	919	1616	1151	1832
E4	4451	4656	206	206	203	215	212	212	212
Int4	4657	7004	2348	541	604	705	952	657	549
E5	7005	7137	133	133	133	133	133	133	133
Int5	7138	10080	2943	1112	878	1066	903	888	824
E6	10081	10178	98	98	98	98	98	98	98
Int6	10179	12023	1845	561	385	1277	1519	1372	1587
E7	12024	12153	130	130	130	130	130	130	130
Int7	12154	12730	577	618	620	879	986	1007	1293
E8	12731	12939	209	215	215	215	214	215	215
Int8	12940	17408	4469	660	718	549	747	806	1399
E9	17409	17541	133	133	133	133	133	133	133
Int9	17542	17636	95	211	624	828	1070	1106	1088
E10	17637	17734	98	98	98	98	98	98	98
Int10	17735	20907	3173	1767	792	137	1145	1166	1177
E11	20908	21046	139	139	139	139	139	139	139
Int11	21047	25794	4748	946	1310	518	327	317	418
E12	25795	26018	224	224	224	224	224	224	224
Int12	26019	27796	1778	1090	766	530	1000	1882	1192
E13	27797	27929	133	133	133	133	133	133	133
Int13	27930	30742	2813	141	142	1019	574	569	614
E14	30743	30788	46	55	65	57	62	68	68
Int14	30789	32317	1529	921	935	276	1062	1123	770
E15	32318	32460	143	134	135	30	139	144	163
Mean	Exo	ns	133,67 ±49,71	134,33 ±48,92	134,87 ±47,50	126,13 ±57,75	133,33 ±50,57	136,2 ±49,01	137,00 ±49,62
±SD	Intro	ons	2175,36 ±1399,94	734,5 ±412,99	703,14 ±274,17	781,71 ±369,56	901,21 ±407,40	976,57 ±382,06	1064,71 ±426,52

Table 2.4. Alignment of *P. lessonae* scaffold 10.7 with the serum albumin cDNA of *P. shqipericus*. Yellow coloured regions indicate exonic positions, grey coloured regions show intronic sequences, green coloured areas represent DNA transposons belonging distinct families; pink coloured regions indicate distinct types of LTR and non-LTR retroelements; blue coloured regions show simple repeats and low comlexity elements; and conserved regulatory motifs in both 5' and 3' regions of the albumin genes are also shown.

U40452_Pshqipericus		[100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AACACTTGGTTCTAAACTGTATACACATGCAATAAACATAATGCATACAGGAAGGA$	[100]
U40452_Pshqipericus		[200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AGCCGGCATACACAGTCTGATCCTAATGGATCGTCTTGGGGATGTCCGGGGAGTACTCACGGATCATTTGTCAATGTCTGTTTGCCTGGATAATTCGTTG}$	[200]
U40452_Pshqipericus		[300]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GCGTTGCCGTTTTGCTTCCCAGGGCAGTAGTGGATGATAAAGTCAAAAGGCTGCAGCGCCAAACTCCAGTATCTTCTGGCCCATAGTCTGTCGGTAGGAG}$	[300]
U40452_Pshqipericus		[400]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTTGCGCACAGTCCCCTCCAAAACGCACAGTGACAGTGGGTTCTGTCACACTGTTCTGTC	[400]
U40452_Pshqipericus		[500]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAAGACTTGAGACTGGGGGGGGGGGGGTTCACCTTCCAGCAGGGGAACGACCCTAAACATACAGCCAGAGCTACAATGAAATGGTTTAAATCAAAGCATATT	[500]
U40452_Pshqipericus		[600]
<pre>scaffold39565_10.7_Plessonae</pre>	CATGTGTTAGAATGGCCCAGTCAAAGTCCAGACCTAGATCCAATTGAGAATCGGTGGCAAGACTTGAAAATTGGAGTAAACTGGAGTGGACTGGAGTAAA	[600]
U40452_Pshqipericus		[700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TGCGAGGAATGTTTGAGAGTTTAGGAGCCCAAAGGAATTGGTTTTTATACAACTGCTCCTCCTTCCT$	[700]
U40452_Pshqipericus		[800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TGGTAAGCAGCCTCTTTAGAGCACTTTGTTTGGTGTTGTGTGTG$	[800]
U40452_Pshqipericus		[900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATCAGAATATAATTTACTCAGCACTGCTTGTTTGGATTTTACTATTAATCTTTCTGCTATCAACAGATTACTGCTAGCTGCTTTAGAGTGACATTTTAT$	[900]
U40452_Pshqipericus		[:	1000]
<pre>scaffold39565_10.7_Plessonae</pre>	GGTTCTTTCAGATGGAGCGCAGATGTT <mark>ATTTGTGTTAAGACTTGAAAATTGCTGTTCGTAGACACTCTCCATCCA</mark>	[:	1000]
U40452_Pshqipericus		[:	1100]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAAGAAGAATGGGCAAAAATGTCCCTCTCTAGATGTGCAAAGCTGGTAGAGACATCCCCAAAAAGACTTGCATCTGTAATTACAGCCAAATTCTAGAAA	[1100]

1140452 P shqipericus	Tc1-10_XT	12001
confold20565 10 7 D locconco		1200]
		1200]
U40452_Pshqipericus	[1300]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTTCACTATTATGTGCCACTTTGTGTCGGTCTATCACATAAAATCCCAATAAAATACATTTACGTTTTTGGTTGTAACATAACAAAATGTGGAAGATTT [1300]
U40452_Pshqipericus	[1400]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAGGGGTGTGAATACTTTCAAGGCATTGTATACGGAAATGAGGTGGCTCTACTGCATCAGATCATGTACTAGATACATGATCTGGCACTTCCAGGAAGG	1400]
U40452_Pshqipericus	[1500]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGTGTGTGCGCGCACTGCTGGGGGACCCAGCCCTTTTTGTGATTAAACACAGCAGATGTCGATCAGCAGGTGCCAATGGATTACCGCCGGCATCTGCCAA [1500]
U40452_Pshqipericus	[1600]
<pre>scaffold39565_10.7_Plessonae</pre>	TCACCGGGGAATTCACACAGGACAGAGCCATGCCTATGTAAACAAGGCAGAGCTCTGTCATGTCAGTTAAGGAAGTAATGGATTTTCTTTC	1600]
U40452_Pshqipericus	[1700]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGGGAATAAAAATCCATTACCTAATAGAATAAAATATCTTAATCACCGCAACTAAATATACAAAAAAATATATAAAAAAATTGTAATTTTAAAGTTGTGA [1700]
U40452_Pshqipericus	[1800]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTGATTGCTGCCAATACTCCACACGTATATTCACCTCACTTTCAATAATATCAGTGTCAAATTAGTACAGTGCAATCATCTATCT	1800]
U40452_Pshqipericus	[1900]
<pre>scaffold39565_10.7_Plessonae</pre>	ATAAAAAATCACTAAATATGTGTACACATATAACCTCCATATATTTCACAATCTGTGACAAAATATAACTTGCTCAAGTGAAAATTCCAAATTCATATGC [1900]
U40452_Pshqipericus	(AATA)n[2000]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTCATATGCACAAGATATATACCAGTG <mark>AATAAATAATATATATATATAAAAAAAAAA</mark>	2000]
U40452_Pshqipericus	[2100]
<pre>scaffold39565_10.7_Plessonae</pre>	TCCCAAAAGCGAGTGTTCTTGCCACCTCTGTGTATAACAATCAACTAAAAAGCAGGTGCTCATGTAATCCACCTCCAGTATTCACTAGCAGCTCACCTCC [2100]
U40452_Pshqipericus	[2200]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCTTTGACTCCTATTTCAGCAGAGAGTCAAAGTATAAAAAAGTATAAAATAGGCTCACTCA	2200]
U40452_Pshqipericus	[2300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGACCATTACTATTAGATGGTGGTAAGTCGGTGAATATTGCTTTATGCTGTATATAGCTGGCACGACTTGCAGTGCTGTTCGAAGTGAGTG	2300]

U40452_Pshqipericus	[24	100]
<pre>scaffold39565_10.7_Plessonae</pre>	TATACTTTTTTATAATAAATCTGCCTTTACGGAGAGCACTAGATGGTATGTCTTTTGTTTTATATGGCCGAGTAATATACAAGCACCGCATTAAGGAGA [24	ŀ00]
U40452_Pshqipericus	(TATTT)n[25	;00]
<pre>scaffold39565_10.7_Plessonae</pre>	GAGTGGGTTTCTGAAATACCCAGGAACATCCATTCATTCA	;00]
U40452_Pshqipericus	[26	500]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTGCATATGAATGCATGTGAATTTGAAGTTTTAACTTGAGCAAGTATTGCTCAAGTCATTTTGTCACAGATTGTGAAATGTATGGAGGTTATATGTGTA [26	500]
U40452_Pshqipericus	[27	/00]
<pre>scaffold39565_10.7_Plessonae</pre>	CACATATTTAGTGATTTTTTTATATATATGACACAATATTGAGATAGTTGATTGCGCTGTAGTAATTTGACACTGATAAAATCCATTACTTGCCTTAGT [27	/00]
U40452_Pshqipericus	[28	300]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAAGCACCTCACACAGTACACCACAACACTGTTAACTATATAAATGTATTAGTGTCACTGGTTCTCAAAAAATGTCAAAAGTGTCAGTTAGTGTTTGA [28	300]
U40452_Pshqipericus	[29	900]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGTCCACTGCAATATCACAGTCCTGCTATAAGTCACTGATCACCGCCATTACTAGTAAAAAAAA	900]
U40452_Pshqipericus	[30	900]
<pre>scaffold39565_10.7_Plessonae</pre>	ACCATAGTTTGCAGGTGCTTTAACTTTTGCGCAAACCAATCATTATATGCTAAATGGTATTTTTTTT	900]
U40452_Pshqipericus	[31	[00]
<pre>scaffold39565_10.7_Plessonae</pre>	TAATTTTGTCAAGAAATTAGATTTTTTACATTTTTTTTTT	.00]
U40452_Pshqipericus		200]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAATGCAGTGGTAATCAATTGTAGCAATATTTCCAGGCATTGGGCCTGATGTGGGTCTGCCATGGTTACCACCAACAGCACATTTAAGCTTGT [32	200]
U40452_Pshqipericus	[33	300]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCCCAAGTGTGAGATCTGAGACAATGAACAGTCCGGCATCTTTGGTTTTTAGTTTTATTGTAGGAACTTGGATGGGAGAAGGGAATGAGTGGGATACTC [33	300]
U40452_Pshqipericus	[34	100]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCTGGCTCAACTAGGTACAAATCTGGGACACTTGAGGAAAATGCCGCCTAGCGCTGTCTTGGCACTGGCTGAATGCCGACCAACGCTTACTGGCGCTTG [34	100]
U40452_Pshqipericus	[35	500]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAACCTGATTTGCGGTCTGTGAACTGAAACTTGACTTCTATTGCTTCCTCACAGCTCAAAAGTCCTGTTCACTGCTCCCTGTAGTCCTCTGCAACACTT [35	500]

U40452_Pshqipericus		[3600]
<pre>scaffold39565_10.7_Plessonae</pre>	AACACTTTACACTTCAGGGCCCAACCACTGGCACCCCTTTTACTCTTCAGGGTAATCTCTGGCACCCCCTTTCTATATTCCTGTACCGGCTAGCTA	[3600]
U40452_Pshqipericus		[3700]
<pre>scaffold39565_10.7_Plessonae</pre>	CTGGATCCCTCTGACCTTCTCCAGAGTGGTCCCCTCTCCTAGGCAGACCCCAGGGGCAGTTTGCAGTACTCATGACTCAGAATACTTTCCTGAAACTCGA	[3700]
U40452_Pshqipericus		[3800]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCCAACGCTCCACTGGTCCTCCGCACACCTGGATCCTTCTCTGCAGCTCAGCCCAGCCCTGTGTCCACTCAGGCCAGCCGCCTGGGACACCCGATACTT	[3800]
U40452_Pshqipericus		[3900]
<pre>scaffold39565_10.7_Plessonae</pre>	CCTGACAGGATAGACAGCCATCATCTCCCTCCTGGGACTCTGCTTCCCACCAAAGACAATCTGTCATGTGCTCCCCACAGACTTATATCCCCTCCCAGAG	[3900]
U40452_Pshqipericus		[4000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCTGTGCAGCAGAATCTGCCCTTCCAGGATGGATATAAGCCCTGGTGTGCTATTGGTCAGGTTTCCTCCACTGCCCAATGACAGCCCTCATTGGGATAC	[4000]
U40452_Pshqipericus		[4100]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTGGGGAGACTCCCCAACACAACAGGCACACCTGGGAACACATGCCAGCCTAGCTGAAAACATGGCCCTGCTGATAAGCAGACAAGGCCCACTATACTT	[4100]
U40452_Pshqipericus		[4200]
<pre>scaffold39565_10.7_Plessonae</pre>	TACCTAGCACCTACATTGGAGGGTGCTACACAATTATCACCAAAATAAAT	[4200]
U40452_Pshqipericus		[4300]
<pre>scaffold39565_10.7_Plessonae</pre>	ACATAACTGCACAATTGTATAGTTGGCTATACTTATTCTATGGATCACAGGAGTGCAGTTCATTTTGCACTCCTGTGACCCATTTTGGCTACTTAGCTGG	[4300]
U40452_Pshqipericus		[4400]
<pre>scaffold39565_10.7_Plessonae</pre>	TCCAGGTTCAGGAAAGAACGGCAGCTAGCTAGGCCTTTTAGAATGCCACTGAGAGCCTGAGCTAGCCACTCCTGCCCTCCCCCCACAGCCCAGCACTC	[4400]
U40452_Pshqipericus		[4500]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTGGGTGCTGGAGGAGCAGAGCAGAGAGCCAGTGAGTG	[4500]
U40452_Pshqipericus		[4600]
<pre>scaffold39565_10.7_Plessonae</pre>	CGCTTTGTTCTCGGGGTAGAGCCAGCGGTGGACAGTTGCAGCTGGGGATCGGTGCTGCAGCCATCTAGGGGGAGTATGATTTTTTTGCTATTTCCATAAT	[4600]
U40452_Pshqipericus		[4700]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTCTTTTAAGAAACCTGAGATTGGGCAAAAGTACACTTTCACCATCAGGGAAGCAACCTTACATCATACATCTCAACTGTCCCTCATTTGAATGAA	[4700]

U40452_Pshqipericus		[4800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TCCAATTTTTGAAGCCAGATCCCCCTGTCCCTCTTTTCCTTCTCAGTTGTCCCTCTTTTGGAAAGATGTATAGGCCTATCTAAGAAGTTAAATTACTTA}$	[4800]
U40452_Pshqipericus		[4900]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:constraint} ATGGTTTAGCCAATTATTTTTTTTTTTTTTTTTTTTTTT$	[4900]
U40452_Pshqipericus		[5000]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTATAAGGTGTTCCTTTTTCATCTCAGTTGAGAGGTATGCTTAAATGCGAAT <mark>CTAAAACCAAACTGTTGTGGTTTAAAAAACAAAATCCTAAAGGGCTT</mark>	[5000]
U40452_Pshqipericus	TCI-RT	[5100]
<pre>scaffold39565_10.7_Plessonae</pre>	GGAATGACCCATTCAACACCCACATATCAATTAAACTGAGATTCTGTGGCAAGACTTTAAAATTCCTGCTCATATATGGTCCCTTTTCAACCTAACAGAG	[5100]
U40452_Pshqipericus		[5200]
<pre>scaffold39565_10.7_Plessonae</pre>	CGATTTTGTCAAGAATAATTGGAAAAAACTGCAGTAATT <mark>GCAGCCAAAGGTGATTCTAGGTTTTGATTCAGGGGGATAAATCATGATGGGAACATAAAAT</mark>	[5200]
U40452_Pshqipericus	<i>ТС1-РР</i>	[5300]
<pre>scaffold39565_10.7_Plessonae</pre>	TACTGTTTTACTTTTTTCCTATTACCTTTTATGTTACAATAAAATATATGCATCT ATAACAGTCTGAGAGCGGTTACGTAGCACCCTGGGTTCTCATC	[5300]
U40452_Pshqipericus		[5400]
<pre>scaffold39565_10.7_Plessonae</pre>	CATACAGGTTGAGGGTTTCCAGGTTGTGGGTTTGGTACAGAGAAAACTGGTTGTTCAGTGTTCCCTGGAATTAGTCATGA	[5400]
U40452_Pshqipericus		[5500]
<pre>scaffold39565_10.7_Plessonae</pre>	GTATTCGGCCCCTTGAACTTTGCGACCCTTTGCCACATTTCAGGCTTCAAACATAAAGATATAAAACTGTAATTTTTTATGAAGAATCAACAACAAGTG	[5500]
U40452_Pshqipericus		[5600]
<pre>scaffold39565_10.7_Plessonae</pre>	GGACACAATCATGAAGTGGAACGAAATTTATTGGATATTTCAAACTTTAACAAATAAAAAACTGAAAAATTGGGCGTGCAAAATTATTCAGCCCCTTTAC	[5600]
U40452_Pshqipericus		[5700]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTCAGTGCAGCAAACTCTCTCCAGAAGTTCAGTGAGGATCTCTGAATGATCCAATGTTGACCTAAATGACTAATGATGATAAATAGAATCCACCTGTGT	[5700]
U40452_Pshqipericus		[5800]
<pre>scaffold39565_10.7_Plessonae</pre>	GTAATCAAGTCTCCGTATAAATGCACCTGCACTGTGATAGTCTCAGAGGTCCGTTTAAAGCGCAGAGAGCATCATGAAGAACAAGGAACACACCAGGCAG	[5800]
U40452_Pshqipericus		[5900]
scaffold39565 10.7 P. lessonae	GTCCGAGATACTGTTGTGGAGAAGTTTAAAGCCGGATTTGGATACAAAAAGATTTCCCAAGCTTTAAACATCCCAAGGAGCACTGTGCAAGCGATAATAT	[5900]

U40452_Pshqipericus		[6000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGAAATGGAAGGAGTATCAGACCACTGCAAATCTACGAAGACCTGGCCGTCCCTCTAAACTTTCAGCTCATACAAGGAGAAGACTGATCAGAGATGCAGC]	6000]
U40452_Pshqipericus		[6100]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAGAGGCCCATGATCACTCTGGATGAACTGCAGAGATCTACAGCTGAGGTGGGAGACTCTGTCCATAGGACAACAATCAGTCGTATACTGCACAAATCT] [6100]
U40452_Pshqipericus		[6200]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCCTTTATGGAAGAGTGGCAAGAAGAAAGCCATTTCTTAAAGATATCCATAAAAAGTGTCGTTTAAAGTTTGCCACAAGCCACCTGGGAGACACACCAA] [6200]
U40452_Pshqipericus		[6300]
<pre>scaffold39565_10.7_Plessonae</pre>	ACATGTGGAAGAAGGTGCTCTGGTCAGATGAAACCAAAATCAAACTTTTTGGCAACAATGCAAAACGTTATGTTTGGCGTAAAAGCAACACAGCTCATCA]	6300]
U40452_Pshqipericus		[6400]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCTGAACACCATCCCCACTGTCAAACATGGTGGTGGCAGCATCATGGTTTGGGCCTGCTTTTCTTCAGCAGGGACAGGGAAGATGGTTAAAATTGAT	Ι	6400]
U40452_Pshqipericus		[6500]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGAAGATGGATGGAGCCAAATACAGGACCATTCTGGAAGAAAACCTGATGGAGTCTGCAAAAGACCTGAGACTGGGACGGAGATTTGTCTTCCAACAAG]	6500]
U40452_Pshqipericus		[6600]
<pre>scaffold39565_10.7_Plessonae</pre>	ACAATGATCCAAAACATAAAGCAAAATCTACAATGGAATGGTTCACAAATAAACATATCCAGGTGTTAGAATGGCCAAGTCAAAGTCCAGACCTAAATCC] [6600]
U40452_Pshqipericus		[6700]
<pre>scaffold39565_10.7_Plessonae</pre>	AATCGAGAATCTGTGGAAAGAACTGAAAAACTGCTGTTCAAAAACGCTCTCCATCCA]	6700]
U40452_Pshqipericus		[6800]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGTCAGTCTCTCGATGTGCAAAACTGATAGAGACATACCCCAAGCGACTTACAGCTGTAATTACAGCAAAAGGTGGCGCTACAAAGTATTAACTTAAGG]	6800]
U40452_Pshqipericus		[6900]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGCTGAATAATATTGCACGCTCAATTTTTCAGGTTTTTATTTGTTAAAAAAGTTTGAAATATCCAATAAATTGCGCTCCACTTCATGATTGTGTCCCAC]	6900]
U40452_Pshqipericus		[7000]
<pre>scaffold39565_10.7_Plessonae</pre>	TTGTTGTTGATTCTTCACAAAAAAATTACAGTTTTATATCTTTATGTTTGAAGCCTGAAATGTGGAAAAAGGTCGAAAAGTTCAAGGGG	[7000]
U40452_Pshqipericus		[7100]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTCCAATTTTGGGTCGGGTTTTGCCAGTTACCAGCACAGCACTGACTG	[7100]

U40452_Pshqipericus	[7200]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGGACAGAGGGGGATCAGAGCCCAGAGGTCAAAGTGGGAGCCAATGCAGCAGTAAACTCTGAGGTATACAGAACATTTTGAGTTCTGTCTG	7200]
U40452_Pshqipericus	[7300]
<pre>scaffold39565_10.7_Plessonae</pre>	GAGACAAGGTCTAAGCCTGAGGGCCAAAGCAGCATGGCTACAACATGAAAGCCAGGGGTCTGAAGTATTTGCTTATGGCAGAGAGATGCTAAACAACCTC	7300]
U40452_Pshqipericus	[7400]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCATTGGTAACCCTTTTGTGTTTTTGTGAGCTTTCTTTTAAATAAA	7400]
U40452_Pshqipericus	[7500]
<pre>scaffold39565_10.7_Plessonae</pre>	ATAATGTAAAGCACTGTATGTGGGTTTCTCTTCTGTGAAAAGACGAACTATATTTTAATGTCTCAATACCTGACCTTTACAAGCATTTGGAAACCATGCA	7500]
U40452_Pshqipericus	[7600]
<pre>scaffold39565_10.7_Plessonae</pre>	GGTTTCAGACGCAGACTGGGATACTGTCATAATCAAGGATCACTTTGCTTAACCTTTTGTTCGTAGTTTCACAGCCTAATCACCAAACGCTTACATTAGT	7600]
U40452_Pshqipericus	CAAT Box [7700]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAATTATAACTGATCTACAAGGTCTTTCAAAAAATGTGTTG <mark>CCAAT</mark> CCAAGCCCACTTTACAACTGATTCAGTATTGTGTTTTGTCATTCTTTGTGTTT	7700]
U40452_Pshqipericus	TATA box:TATA(A/T)AA(G/A) [7800]
U40452_Pshqipericus scaffold39565_10.7_Plessonae	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [7800] 7800]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus Initition	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCCATAACAACGGAGTCCAGAC [7800] 7800] 7900]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTACCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGGAGAACCTCCAAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCCCCATAACAACGGAGTCAAGACC [7800] 7800] 7900] 7900]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGAGAGACCTCCAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCCATAACAACGGAGTCAAGAAC [ACTTACAAAAGCGACATCATg [[7800] 7800] 7900] 7900] 8000]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTACCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGAGAAACCTCCAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCCATAACAACGGAGTCCAAGAAC [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATGGTAACAAGAAATTTTAGAACAGCAATCTTTATGTCCATAACAAAGTGTGCTAAGTAGGTTTTGAGTGTTTTTCTGTCTG	7800] 7800] 7900] 7900] 8000]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTATATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGAGAAACCTCCAAAAAACATTTGCCACCATGAAGTGGGGCCACTCTGATTTGTCTGTTTATTCTCTCCCATAACAACGGAGTCAAGAC [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATGGTAAGAAGAAATTTTAGAACAGCAATCTTTATGTCCATAACAAAGGTATGAGTTTGAGTGTTTTTCTGTCTG	7800] 7800] 7900] 7900] 8000] 8000] 8100]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	TATA-box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGAGAAACCTCCAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCCATAACAACGGAGGTCAAGAAC [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATGGTAAGAAAACATTTTAGAACAGCAAATCTTTATGTCCATACAAAGGTAGGT	7800] 7800] 7900] 8000] 8000] 8100] 8100]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	TATA box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGAGAACCTCCAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCCATAACAACGGAGTCAAGAC [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATGGTAAGGAAGAATTTTAGAACAGCAAATCTTTATGTCCATAACAAAGGTTTGAGTGTTTTCTGTCTG	7800] 7800] 7900] 8000] 8000] 8100] 8100] 8200]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	TATA-box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTATAAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCATAACAACGGAGTCCAAGAC [GATTTAGGCTTCTAAGAGAAACCTCCAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCATAACAACGGAGGTCAAGGAC [GATTTAGGCTTCTAAGAGAAACCATCGAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCCATAACAACGGAGGTCAAGAC [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATGGTAAGAAGAATTTTAGAACAGCAAATCTTTATGTCCATACAAAATGTGCTAAGTAGGTTTGAGTGTTTTTCTGTCTG	7800] 7800] 7900] 8000] 8000] 8100] 8100] 8200] 8200]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus <i>Initition</i> scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	TATA-box:TATA(A/T)AA(G/A) [CTAGAAACGAGCCAAATTGTAATTAATTAATAAGCCAGTCTGGTTAATCATTTCCAAATAGGATGCAAAAATGGTATAAATGTCTGCCTTTCCCTTAGCT [site(+1:PyPyAN(T/A)PyPy ATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCATAACAACGGAGTCCAGAC [GATTTAGGCTTCTAAGAGAACCTCCAAAAAACATTTGCCACCATGAAGTGGGCCACTCTGATTTGTCTGTTTATTCTCTCCATAACAACGGAGTCAAGAC [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATg [ACTTACAAAAGCGACATCATGTAAGAAGAATTTTTAGAACAGCAAACTTTATGTCCATACAAATGTGCTAAGTAGGTTTTGAGTGTTTTTCTGTCTG	7800] 7800] 7900] 8000] 8100] 8100] 8200] 8200] 8200]

U40452_Pshqipericus		[8	3400]
<pre>scaffold39565_10.7_Plessonae</pre>	GAATATCCCCAACAAGGACACAACAACAGCAATAAAAATCACTACTTTATTGAAAATAAACATGCTCTGAAGTTACATTTTTTGTGACTGCTATGCTTAT	[8	3400]
U40452_Pshqipericus	ААБААСАТССАССАСАТСАССАССАССАСАТААТАААССАТАТТСТАААССАДСАССАССАСТСТСАААА	[8	3500]
<pre>scaffold39565_10.7_Plessonae</pre>	GTGTTTCCTATGCATTACATTTTTTATCTGTTTTTTTACAAGAACATCCACGGATAATAAACGATATTGTAAAGACAGTGGGAAAACCAGCTGTTGAAAA	[8	3500]
U40452_Pshqipericus	ATT	[8	3600]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTELAAGTATTCAATTTATATAATGTGTTTTATTGTTCTGTAAACACTTTGAAAATTGCATAAAATGCAAACTTCTAGATAAAACTGTTTTCAAAATTT	[8	3600]
U40452_Pshqipericus		[8	3700]
<pre>scaffold39565_10.7_Plessonae</pre>	TATAGTAGAACTATAGGCATTTTTTTTTTTTTTTTGGATAGAGGGAGG	[8	3700]
U40452_Pshqipericus		[8	3800]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTTCCTGTTTTATAACCAAAACAAGAAGTGAGAAGAAAACCCTCCAAAATAAGGTAATCCTTTGAGGACCCCCAGGCCATCAGAACTAGTGTCCCCATC	[8	3800]
U40452_Pshqipericus		[8	3900]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAAGATTTCCCTTCTATTGCTATTACAAAATTTGGGATTTCCTTTCACTTTTACTAACGGTAAACATGACAAATAGAGGGTGAATCTCGCTAATGGGGG	[8	3900]
U40452_Pshqipericus		[9	9000]
<pre>scaffold39565_10.7_Plessonae</pre>	CACAGACAGCAATAAAAACTCCCAGGTGTTCTACTGCCTCTCCATTCTGTTCAAAACTAAAGTAAAAGTTTTACCTTTAGTTATACTTTAAGGCTTGGTT	[9	9000]
U40452_Pshqipericus		[9	9100]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTATTAAAGTGATGATTTCTAATAGATTTCTTTTTTCCACTTCCTCATTATTCTTTGTTGCCTGGCTTGGTATGTCTAAATAAGGTTGAGCTATAA	[9	9100]
U40452_Pshqipericus		[9	9200]
<pre>scaffold39565_10.7_Plessonae</pre>	AAATTGTAGGCAAGGAAGAAGTTGGTGTTCTAGTAGAAACAAATAGATATTGGATTATTATTTTTTCAAATAGATATCGGATTATTATTATTATTACCATT	[9	9200]
U40452_Pshqipericus		[9	9300]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTATAGTTTAATCATCTCCCATAAAACCTACAAAATGTAACAGTTAATACACATGCATAACACGTTTCCATTATTAGTATGTAAAATCTGTACGTGAAAAAC	[9	9300]
U40452_Pshqipericus		[9	9400]
<pre>scaffold39565_10.7_Plessonae</pre>	ATACATAATTAAGCTATAGCAGGATACAGTATTCAGAACAAACA	[9	9400]
U40452_Pshqipericus		[9	9500]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGCATTAGTGGTTTTCATTATCCTTCTGATACAGTGAGGGTAACTGGTGTCTGGATGATGGTGGATATGGCAGGGCTTTGTGTACAAGGCCTCTTCATT	[9	9500]

U40452_Pshqipericus		[9600]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAGTGGCTCCATAACATATTCCTGTTTACAGCTACATTTTAAAGACAGATGTTGACACACAATGATTAGAATGAAT	[9600]
U40452_Pshqipericus	AGTGTTGGTGATGGTGGCACAGGATTTTGAAAAATGCTCACTGGATGAACATCTGAAAGTCCAGGCAAAAATTATAGAGGCTGTTGATAACTG	[9700]
<pre>scaffold39565_10.7_Plessonae</pre>	CTTACAGAGTGTTGGTGATGGTGGCACAGGATTTTGAAAAATGCTCACTGGATGAACATCTGAAAGTCCAGGCAAAAATTATGGAGGCTGTTGATAACTG	[9700]
U40452_Pshqipericus	CGAAAAACATCCAGAGGAAGCTGAGTGTAAAAAACCAGCG	[9800]
<pre>scaffold39565_10.7_Plessonae</pre>	CGAAAAACATCCAGAGGAAGCTGAGTGTAAAAAACCAGCG <mark>GT</mark> AAGCTCTCCTTGGCTCCTTCCCCGCAAGCTTTACATAATAATAACAGACATTTTTA	[9800]
U40452_Pshqipericus		[9900]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTTGCAGTGGTGCAGAATACAGGGATCAAGAGTATGTAACACACATTGTGCAGGGTTTAACAGTATGTAACACACGGAGTCCAGGGTCAAGAGTTTGTA	[9900]
U40452_Pshqipericus		[10000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAGCACAAAGTGCAGGGGGTCAAGGGATCATAATGACAAATTGCAGGGGTTAAAAGTGCATAGCACAGTATTTAGGGGTCAGGAATACATAGCGTACAGG	[10000]
U40452_Pshqipericus		[10100]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGTATACACAGTAGGACCAAAAACAACCCTCTTCTCTCAGTACAAATGTCCCCCTCAGTACAAATTCTCCCCTAAATGCCCCCAGCAGTGGTGTATT	[10100]
U40452_Pshqipericus		[10200]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGGTTTTGTGCTGCCCTAGGCCTGACTAAACTTGTGATCCCCCTAATTTAAATATGACGCACCCCTTCCTGTAAAGGCCACACCTCTTCCTGTTTATGA	[10200]
U40452_Pshqipericus	hAT-Z_TS	[10300]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCGCCCTGTCATCTGTAAAC <mark>CACAGACCACAACCACAGTCTATCAGTGCCCATCAGTGTAGCCTCATCAGTTCCCAGAAGGTCAGCCTCAGCAGTGCCC</mark>	[10300]
U40452_Pshqipericus	(ATCAT)n	[10400]
<pre>scaffold39565_10.7_Plessonae</pre>	ATCATCATACCATCATATCATACCATATC	[10400]
U40452_Pshqipericus		[10500]
<pre>scaffold39565_10.7_Plessonae</pre>	AGCCCAATTTGCCCTATAAGACTGGCACTATACTAAAAGTGTGTAGCGCAAGCCGTCAGGGACTCTTTTTAGTGGGGGGGG	[10500]
U40452_Pshqipericus	hAT-32TSCPB	[10600]
<pre>scaffold39565_10.7_Plessonae</pre>	CGTGGTGCAGCTTT <mark>TTTGCTGCCCCCTGCAAAGTGCTGCCCTAGGCCTGGGCCTTGTTGGCCTAGGCCACGATAAAGCACTG</mark> GCCCCCAGTACAGATCT	[10600]
U40452_Pshqipericus		[10700]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCCCCAAAAATCCCTCCCAGTAGAGAGCTCTTTTTGCGCAAATCATCCTCAGCAAAGGATCCCCTTCCCCCCAAGTTTCTTTATTACAGACCTCCCCA	[10700]

U40452_Pshqipericus		[10800]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAATACAGACCTCTCTCCCCCAGTACAGACCTTTTTCTCCCCAGTACAGACCTCTCCCCCAGTACAGACCACAAGAGACCCCCCCC	[10800]
U40452_Pshqipericus		[10900]
<pre>scaffold39565_10.7_Plessonae</pre>	CACTTACCAGACCTCAGATACAATGGAAGGGCACAGTGAAGGGGTAGTCCTTGGCAGTGTCACTTGACACTGTGCATCAAAGCTAAGGAGGAGCTTACAG	[10900]
U40452_Pshqipericus		[11000]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCCCTCAGCTCTGTACATTCACTGTGTCTGAGGTTTCGATTAAGCCAATGTCTGCAGTGCAAATGGCGGAACATAAAAAAAA	[11000]
U40452_Pshqipericus		[11100]
<pre>scaffold39565_10.7_Plessonae</pre>	GAACCAGAGCCCTGAGCTAGTGACATCTCTGCACTTCACCACTGGGCAGCTGAAGACAACTTTGGGGATGTTTGTT	[11100]
U40452_Pshqipericus		[11200]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGGGACTATCTCCTAAAAATTGGGGACAAATAGCAACTTTAACTTAAAGTGTATTTAAAGTCCAAACTTTGGAAGAGCCTTCTCTGAGTTTGAAGGCTG	[11200]
U40452_Pshqipericus		[11300]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAGAGATCTGTCTCATCCTTACTGCATCTCCCTGCAGCAACCTACCACATCCTACCATGTCATTACTGCTCTCCCTGCAGCAACAAATCATTGGTATAT	[11300]
U40452_Pshqipericus	INTRON-3	[11400]
U40452_Pshqipericus scaffold39565_10.7_Plessonae	TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC	[11400] [11400]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC	[11400] [11400] [11500]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC	[11400] [11400] [11500] [11500]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC	[11400] [11400] [11500] [11500] [11600]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCCAGGAGGCACTTCAC GCAATCTTGCCTGTGCATCAGGCCTCGCTAAGTCCTCCTGGCCCAGGCCAGGATTTAAGAAAAAAAA	[11400] [11400] [11500] [11500] [11600] [11600]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC GCAATCTTGCCTGTGCATCAGGCCTCGCTAAGTCCTCCTGGCCCAGGCCAGGATTTAAGAAAAAAAA	[11400] [11400] [11500] [11500] [11600] [11600] [11700]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC GCAATCTTGCCTGTGCATCAGGCCTCGCTAAGTCCTCCTGGCCCAGGCCAGGATTTAAGAAAAAAAA	[11400] [11400] [11500] [11500] [11600] [11600] [11700] [11700]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC GCAATCTTGCCTGTGCATCAGGCCTCGCTAAGTCCTCCTGGCCCAGGCCAGGATTTAAGAAAAAAAA	[11400] [11400] [11500] [11500] [11600] [11600] [11700] [11700] [11800]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC GCAATCTTGCCTGTGCATCAGGCCTCGCTAAGTCCTCCTGGCCCAGGCCAGGATTTAAGAAAAAAAA	[11400] [11400] [11500] [11500] [11600] [11600] [11700] [11700] [11800]
U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus scaffold39565_10.7_Plessonae U40452_Pshqipericus	INTRON-3 TCAATTTCTTTTAATTAAATAGTTTCTTACTGTGTTTTTTGCTTTCTTGTAATGTTCTCCATGAGTTCCCAAACTTCTTTATTTTCCCTGCCAGCTTC TACCAGTTTTCTAGAATAGACTGTGCATACCTACCCATGCGTGAGATTTCACAAAATCCTCCTGTAATTTCTATGTAACACATCCCAGGAGGCACTTCAC GCAATCTTGCCTGTGCATCAGGCCTCGCTAAGTCCTCCTGGCCCAGGCCAGGATTTAAGAAAAAAAA	[11400] [11400] [11500] [11500] [11600] [11600] [11700] [11700] [11800] [11800] [11900]

U40452_Pshqipericus		[12000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAATAGATTGCGACATGTGCTTTCATATAGTCATTGCAGGGTTAAATTACAATTGCCCCCCTTCCCCACCTAACATTAGCCCTTAACCATGGGCCACATT	[12000]
U40452_Pshqipericus		[12100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TCCTTAAAAAATCCCCAACTGCATGTGTGTGTGTTAGACCCCTTTCACATGGTAAGGACTCCTTTGAGAGCAGGAGGAGCAGGGGATCAGTCTGCTGATCAGAGCCATG}$	[12100]
U40452_Pshqipericus		[12200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATCTGCTCTATGGGAGTAAATGGACTCCGCTGTCTGTTTACACCAGCCGGTCCATGGAGGTGAATAGACCATCAGGGGAACATGATCGTACCCTTCATGT}$	[12200]
U40452_Pshqipericus	ATTGAACTGTATCATGACATAGTTTGCAAAGAAGAAGAAGAAGACATAGAC	[12300]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAAGGGGCCTAAAACTGCCTTCTTTAATTGTGTGTGTGT	[12300]
U40452_Pshqipericus	CAGCTCTACCCCTGGACTACAGAGTGTTGTGGCAAAGCGGAGGCAGAGAGGACCAAGTGCTTTTATGAGCATAGAGAAGTTCGAGTTGAGGAATATAAGA	[12400]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGCTCTACCCCTGGACTACAGAGTGTTGTGGCAAAGCGGAGGAAGAGAGGACCAAGTGCTTTTATGAGCATAGAGAAGTTAAAGTTGAGGAATATAAGA	[12400]
U40452_Pshqipericus	TACCAAATATTGAAGAATCCTGCAAAGAACACAAAGAGCACCCACAACGTGCCTTCTCTTA	[12500]
<pre>scaffold39565_10.7_Plessonae</pre>	TACCAAATATTGAAGAATCCTGCAAAGAACACCAAGAGCACCCACAACGTGCCTTCTCTTA <mark>GT</mark> GAGTTTATTTTAGTATTTTATATATACCATTGTACTT	[12500]
U40452_Pshqipericus		[12600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTAAAGCCCAATTGCCCACCACGTGTTATGATTTGTCATGATTTGACTTTTTTTGGAAATGCCTAGAAATAACTTTGTCCTATCATGGAAGCCTAATAACTTTGTCATGGAAATGCCTAGGAAGCCTAATAACTTTGTCATGGAAGCCTAATAACTTTGTCATGGAAATGCCTAGGAAGCCTAATAACTTTGTCATGGAAGCCTAATAACTTTGTGAAATGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAATAACTGCCTAGAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGCCTAGAATAACTGCCTAGGAAGCCTAGAATAACTGCCTAGGAAGCCTAGAATAACTGCCTAGGAAGAATAACTGCCTAGAATAACTGCCTAGGAAGCCTAATAACTGCCTAGGAAGAATAACTGCCTAGGAAGAATAACTGCCTAGAATAACTGCCTAGAATAACTGCCTAGAATAACTGCCTAGAAGAATAACTGCCTAGAAGAATAACTGCCTAGAAGAATAACTGCCTAGAAGAATAACTGCCTAGAAGAATAACTGCCTAGAATAACTGCCTAGAAGAATAACTGCCTAGAATAACTGCCTAGAAGAATAACTGCCTAGAAGAATAACAATAACTGCCTAGAAATAACTGCCTAGAAATAACAATAACTGAAGAAGAAGAAGAAGAAGAAGAAGAAGAAGAAGAAGAA$	[12600]
U40452_Pshqipericus		[12700]
<pre>scaffold39565_10.7_Plessonae</pre>	AATATGTATGGTGCCATGATTTGCCACTAGAAAAGGAAATTTATGGTGTGTATTGATACAATTTTCCTAAATGTCCAATTTAA <mark>TTTTCACCTTCCTCTGG</mark>	[12700]
U40452_Pshqipericus	CR1-2 XT	[12800]
<pre>scaffold39565_10.7_Plessonae</pre>	ATCAACTGTGGGTATAGGATTGTGTATATGGGATTGTATGATTGTTTTTT	[12800]
U40452_Pshqipericus		[12900]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTAACTATGT AAGTCTCATACTTTTTTAACCTCTTTACAGCCCATTCAGACTGGCGCGACACGACTTTGTGGGGGCAACCTGTGAAGTCGCGGTGCATAG	[12900]
U40452_Pshqipericus		[13000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt cttgcactgcgacttcggcgcgacttgcacggacttcatataattgaagtcaatgcaagtcgcctccaggtcgcctcaaagtaatacaggagcctttttc}$	[13000]
U40452_Pshqipericus		[13100]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAGTCGGAGCGACTTGTGTCGCTCCTATTAGAATGGGTCCTGTACTACTCAATGGCGAGCGA	[13100]

U40452_Pshqipericus		[13200]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGAGTTATGTATCCCTTCATGTTATGATTTGGGGCTGTTTTTTGTTTTATTTCTAAAAAAATAATCTTTTCTCATAGTGATGTCTGCCTTTATATTACA	[13200]
U40452_Pshqipericus		[13300]
<pre>scaffold39565_10.7_Plessonae</pre>	GTGGTCTTTGCTGGCAGAATTGAGTAACAGCCACACCTGTCATCACAAATGTCCTTACACAGACCTCTATTATAGAGAAGGTTAGGAGGTGTGTCTGACC	[13300]
U40452_Pshqipericus	INTRON-4	[13400]
<pre>scaffold39565_10.7_Plessonae</pre>	ATATCTGTGTGAATTACTGTGAGCACTGGGTCATTTCTAGTGCCTATTGGACAACACAG <mark>TTCTTCTTAAACTGGGCATGGATGGTTAGGCACTGATGTTG</mark>	[13400]
U40452_Pshqipericus		[13500]
<pre>scaffold39565_10.7_Plessonae</pre>	GGAGAGAAGGCCTGGCTTGCAGTCTGTACTCTAATTTATCCCAAAGGTGTTCTATCGGGTTGAGGTCAGGACTGTGCATGGCAGTCAAGTTCCTCCACCC	[13500]
U40452_Pshqipericus	TS1 ER3	[13600]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAACACATTAATCCATGTCTTTATGAACCTTGCTTTGTGCAATGGTGCACAGTCATGTTGGAACAGAAAGGAGCCAACCTCAAACTGTTCCCACAAAGT	[13600]
U40452_Pshqipericus		[13700]
<pre>scaffold39565_10.7_Plessonae</pre>	TGAAAACATGAAACTGTCCAAAATTTCTTGGTATGCTGATGCCTTAAGAGTTCCCTTCACTGGAACTAAGGAGCCAAGCCCAACCCCTGAAAAACAACCC	[13700]
U40452_Pshqipericus		[13800]
<pre>scaffold39565_10.7_Plessonae</pre>	CACACCATAATCCCCCCTCCACCAAATGATTTGGACCAGTGCCCAAAGCAAGGTCCATAAAGACATGGATGAGTGAG	[13800]
U40452_Pshqipericus		[13900]
<pre>scaffold39565_10.7_Plessonae</pre>	TCCTGCACAGAGCCCTGACCTCAACCCGATAGAACACCTTTGGGATGAAATAGAGCGGAGACTGCAAGCTAGGCCTCTCCAAAATCAGTGTCTGACCTCA	[13900]
U40452_Pshqipericus	TST FR3	[14000]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAATCCGCTTCTGGAAGAATGGTCAAACATTCCCATAGACACTCCTAAACCTTGTGGACAGCCTTCCCAGAAAAGTTGAAGCTGTTATAGCTGCAAAGG	[14000]
U40452_Pshqipericus		[14100]
<pre>scaffold39565_10.7_Plessonae</pre>	GCCAACTCAATATTGAACCCTACGCTCTAAGACTGGGATGCCATTAAAATTAATT	[14100]
U40452_Pshqipericus		[14200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTGAACCTGACCTAAGACAGGAAGTGTGTTACTGGTTGGATCACCGGGTGAAAGTAAGGGAAAAAAAA	[14200]
U40452_Pshqipericus		[14300]
<pre>scaffold39565_10.7_Plessonae</pre>	ATCTAAGGATCGCTAAGCTACAATATATATATATATATAT	[14300]

U40452_Pshqipericus		[14400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt cttttacactaaatggtggaagtatacctgtgatcatgtcaaacataccataccatgcgcaggaaaatttgaaaaaaaa$	[14400]
U40452_Pshqipericus		[14500]
<pre>scaffold39565_10.7_Plessonae</pre>	ACATGTTCAGCTCACTTTGCTTTTCTTCATACTTTAGTATATCAGGCTGGCAGGATTTACACTTGCATTTATTT	[14500]
U40452_Pshqipericus		[14600]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAATTCACCATTACAATGTATAGAACTTTGACAAAAGTCAGGATTTGCGTTTTTTATAATTCGCAAGGAACATGTTAATAGGCTAAAATGAGAGAGA	[14600]
U40452_Pshqipericus		[14700]
<pre>scaffold39565_10.7_Plessonae</pre>	ATATAACAAACATGTTGCACACTTGCATTTTTATATGTGACTTTAGAATTCATCTATCACCGACTTGATCACAATAAAGTCTTACTTGACTGCTTTTGGC	[14700]
U40452_Pshqipericus		[14800]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTGCTGTACTAACCAGTAACAAAGTCCCTATACAGCATGTTTGTGTGGCATTATGCTCAGTACATTCATT	[14800]
U40452_Pshqipericus	CTACCTCTCCAATATTGCTAAAAGACATTCAAAACTCTATCCACCAGCTGTTCTGGGTTTTGCTATACAATACAATGAAATCACTACTGAA	[14900]
<pre>scaffold39565_10.7_Plessonae</pre>	ACATTTCAGCTACCTCTCCAATATTGCTAAAAGACATTCAAAACTCTATCCACCAGCTGTTCTGGGTTTTGCTTTCCAATACAATGAAATCACTACTGAA	[14900]
U40452_Pshqipericus	TGTTGTGCTGCAGAAGACAAAGCCAAGTGCTTTGGTGAACGG	[15000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTTGTGCTGCAGAAGACAAAGCCAAGTGCTTTGGTGAACGG <mark>GT</mark> ATGCCATTAAACTTCTTGAAATTAAACAATGTTATTGCACATGTAGGCTTCCTTC	[15000]
U40452_Pshqipericus		[15100]
<pre>scaffold39565_10.7_Plessonae</pre>	TACGGGGCGATTACATTTGTGTCCCATGTGTGTTTACCTGCATGGGGATACACAGATGTTTCTGCACCCTGTGCCGAGTAATCCCATTCTTGTCTATTGG	[15100]
U40452_Pshqipericus		[15200]
<pre>scaffold39565_10.7_Plessonae</pre>	ACACAGCGGCTGCATGGACACAGCTGCCATATGACACCAGTGGGTGCGGGTCCCCAAATGCACGTGCACGGTTATCACATAGGGAGCAGCGCTGTGTCCG	[15200]
U40452_Pshqipericus		[15300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCAGTTGCTGCGCCCCAAGACAGCTATGGGTCTGCTGGCACGGGGGTTTGCATGCA	[15300]
U40452_Pshqipericus		[15400]
<pre>scaffold39565_10.7_Plessonae</pre>	GGACGTGGATGTTATGGCCTGGTGTGAACTAAGCCTTAAAGTGTATGTCTGAGAAAAATAATTAAAAAAAA	[15400]
U40452_Pshqipericus		[15500]
<pre>scaffold39565_10.7_Plessonae</pre>	ACAATTGCACATGTTTTATTATTTTAAAAAAAATGCATCAAGTACAGAAAATGGCTTGTGGATTGCCAGAAACCCAGTGATTTCCTAATATCCTGTTGTT	[15500]

U40452_Pshqipericus		[15600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGCTGACAGCACAATGCTTCTGCTGCATTGAATTCAGTGGCAGTGTTGTCAGTCCTGCCCAATTCTCTTCTGCTGGATTACTAAACTCCAGTAGTCCGCA}$	[15600]
U40452_Pshqipericus		[15700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGCAGAACAAATAGTACTGTTTTGGATTTCAGCATAACAAAGGCCTTGGGCTGTCATCTGCAACAGTAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTGTGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTGTGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTGAAGTTGACACTTTGGTGAACGGGTATGCCATGAAGTTGACACTGAAGTTGACACTGAAGTTGACACGGGTATGCCATGAAGTTGACACTGAAGTTGACACGGGTATGCCATGAAGTTGACAGTGAAGTGGCTGTGAAGTGGAAGTGGCATGAAGTTGACAGTGAAGGTGAAGTGAAGTGAAGTGACGGGTATGCCATGAAGTGACGGTGTGAAGTGACGGGTGTGTGAAGTTGACACTGAAGTGACGGGTAGGGCTGTGAAGTGAAGTTGACACTTGGGGTGACGGGTGGCGAGTGGCGGTGGCGGGTGGCGGGGGGGG$	[15700]
U40452_Pshqipericus		[15800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AACTTCTTGAAATTAAACAATGTTATTGCACATGTAGGCTTCCTTC$	[15800]
U40452_Pshqipericus		[15900]
<pre>scaffold39565_10.7_Plessonae</pre>	CCATAGGTGCATACTGCTCTAAATCTTAGTATTGATCGGTGCCTCCTATGCCTCCCAGATTGCTTTTGCCTGCGTTTAGATGTTCCTACTTTATTGTAGT	[15900]
U40452_Pshqipericus		[16000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TGTAAACGCTTACATATACCTAGTGAAGTGAACAGGCTGATACACAGCGATGAAACACATCCTCCTACATAAGTTGTACCTGTTATCTGCCACCCTCTCT$	[16000]
U40452_Pshqipericus		[16100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCTACAGCCATTTAAAGTCAGAGCTTTAGGAGACAGGGTGGGGAACTGACATCACACACA$	[16100]
U40452_Pshqipericus		[16200]
<pre>scaffold39565_10.7_Plessonae</pre>	CCTGAGTGGAGGGAATGGACACACCTCAGGGAAACATGCACAGCTGAGGCTGTCAATCACCCGCTGTGTGCTGGAGGGAAGGAA	[16200]
U40452_Pshqipericus		[16300]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGGCATCACATACCTCCTAACCTTTTGAGATAGAAGCGAGGGACACCTATTAGCAAAAGTGTGTAGTCATAGGACACACCCCTGCCACACCCAATTAA}$	[16300]
U40452_Pshqipericus		[16400]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:constraint} AGGAGAATTATACAAAAAAAAAAAAAAAAAAAAAAAAAA$	[16400]
U40452_Pshqipericus	INTRON-5	[16500]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAATTTAGAAATTGGATAAAGGTTTAGCACTGGGAAACCCTTTTTGATAGATA	[16500]
U40452_Pshqipericus		[16600]
<pre>scaffold39565_10.7_Plessonae</pre>	GACAAAGGAGGAGGAAAAAGGGACATAGGGACTTTGTTCCAAATCAGGGAGAGTCCCTCAAATTAGGGACAATTGGGAGCTATGGCATTAGCATAAACTG	[16600]
U40452_Pshqipericus		[16700]
<pre>scaffold39565_10.7_Plessonae</pre>	TGAAATGTAACTCATGCAGATAGCAGAAAAAGGAGAGAGA	[16700]

U40452_Pshqipericus		[16800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt CATTTTTTATTTCAGAGGTTTAAATCCACTTTGCATTTGCGTGCG$	[16800]
U40452_Pshqipericus		[16900]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAACACAGCTAATCTGACTTCACCTAAATGCACGTAAAAGCACAGTTTGAATGGCCAATTAAAAACTCATTACATGCATTTACTGTTTATGCATTTAGA	[16900]
U40452_Pshqipericus		[17000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAGTCCTCTGAATTATATATAAATGCAGGTTTGACACTGAATGAGAACAAAATGCATAATAAGTTTTATTACCTCAAAAGAAAAACTATTACAGGGAATCT	[17000]
U40452_Pshqipericus	/T]n	[17100]
<pre>scaffold39565_10.7_Plessonae</pre>	ACAAAGGTTTGGGGACAA <mark>TTTCTTTTTTTTTTTTTTTTT</mark>	[17100]
U40452_Pshqipericus		[17200]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTGAAACCTCCACTTCCTTTGAATGTGATACAAAATAATAATATATGTAAATACTCTAAATTCAACACAAAATCAATATAAATTTTCAGTTGCAGCTGT	[17200]
U40452_Pshqipericus		[17300]
<pre>scaffold39565_10.7_Plessonae</pre>	GCGCTCCTTATTGATTTCAGCACACTAAAAAACAAAAATCAATTGTATCCTGTATAGTGCAGTGCAACAGAACAATTATGTGATAAATTCCTGGAAATCAA	[17300]
U40452_Pshqipericus		[17400]
<pre>scaffold39565_10.7_Plessonae</pre>	ATATAACTCAATATTAATTTGTGAAAAATAGAACCAGTCCTTTCAGAACACCATTATATAAATAA	[17400]
U40452_Pshqipericus		[17500]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTCTTCAATCGATGGTATTGTGAAAGGACTAGTTCTATTTTTCACAAATAAAT	[17500]
U40452_Pshqipericus		[17600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGCGCTGCACTATACAGGTTCCAATTATTTTTCTATATGTATACAATCGTTTTACATATAAATATTTAATACGTGGCAAGCGACTCCCAGAATCCAATG$	[17600]
U40452_Pshqipericus		[17700]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTTAGCTGTAGATAGTCTGGCAGTAGCTTATGTTTTCTCACGCTTTGTTTAAAACCTTGGGCAAAGACATCTGATAAGGGATGAGTCACTGCAAAGGCT	[17700]
U40452_Pshqipericus		[17800]
<pre>scaffold39565_10.7_Plessonae</pre>	CGTTTGTCTTTCAGCTTGAGCTAGGCAGCACTGTAAACAACAGATAAATACAAGCAGTGGATATAGTATAATATGCAAAAATATACCTTCTGCTATTT	[17800]
U40452_Pshqipericus	ATGCCCCAGGTAAAA	[17900]
<pre>scaffold39565_10.7_Plessonae</pre>	CTTTATGTGTCTGCTTTGTGCTTGTATATGTGTATATATA	[17900]

EXON-6

U40452_Pshqipericus	AAACTAACAAACTATCTTGAGGATAAACATAAACAGAAATGCCGTGTTCTTAAAGAATTTCCAGAAAGAGTTTCTCAAGCACT	[18000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAACTAACAAACTATGTTGAGGATAAACAGAAACAGAAATGCCGTGTTATTAAAGAATTTCCAGAAAGAGTTTCTCAAGCACT <mark>GT</mark> AAGTATCTATTCCCA	[18000]
U40452_Pshqipericus		[18100]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:action} ATCCAGGAAATATATGTAGGGGAAATGTGGAGAGGCTTTATTTTTATCTTTAGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCCCCTTACCGCTCGCCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCGTGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCTCGCCCCATTTCCATCTTTGACAGCTCAATCTCGTGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCTCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCCCCATTTCCATCTTTGACAGCTCAATCTCTGTGCGCCTTACCGCCCCATTTCCATCTTTGACAGCTCAATCTTGGCGCCTTACCGCCCCATTTCCATCTTTGACAGCTCAATCTTGGCGCCTTACCGCCCCATTTCCATCTTTGACAGCTCAATCTTGGCGCCTTACCGCCCCATTTCCATCTTTGACAGCCCCATTTGCGCGCCCATTTCCATCTTTGACAGCGCCCATTTGCGCGCCCTGGCCCCATTTCCATCTTTGACAGCCCCCATTTGGCGCCCCCATTTTGACAGCCCCATTTGCGCGCCCCATTTGCGCGCCCCCCATTTGCGCGCCCCATTTGCGCGCCCCATTGCGCGCCCCCATTGCGCCCCCCCC$	[18100]
U40452_Pshqipericus		[18200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGGAGCATGCAGTGAGCGCGCTCTCAGCACAGAAACCTATGCTGTCCATAGAGGGATATGTGGGCCATGTGGGTGTTAATACCCATCTTTGGGCCACTG	[18200]
U40452_Pshqipericus		[18300]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCATATAACTATATACATGCCAGAGTACCTTCATCACTATGTTGCTATAAATTTACCCTGACCTCTATTCTTTTGGTTGG	[18300]
U40452_Pshqipericus		[18400]
<pre>scaffold39565_10.7_Plessonae</pre>	TGACCTGGGAGGGACAGGGATGGGATGGGGGGTTGCCCCTGGGGATCGGATGGATGGGGGGGGGG	[18400]
U40452_Pshqipericus		[18500]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTGATACTTCTTCCTTAAAATCCCTGTCAGGTTATTTCTTATGTCTGTGTCCAATCGGAAAGAATTACCTTTAGTTTCTGCTCTGAAGACGCAACAGGA	[18500]
U40452_Pshqipericus		[18600]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTAAGAGGAAATTTCTCCAAAGTGAAGGGAATTGCCCCTGAGACAGTTTTCACAGAAACATTTGTTCAGTGGCATCTTAAGAGCATTATAGGCCCCCGG	[18600]
U40452_Pshqipericus		[18700]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAATGCAGTACACTGGGGGCCCCGCCTGCAAAATCACCCACAAGAATAAACATGCTAATTATCATTAAGGCTATATTTATT	[18700]
U40452_Pshqipericus		[18800]
<pre>scaffold39565_10.7_Plessonae</pre>	AATGACAGCTTCCAAGTTCCGCGGCTGAGTTCGGTTAACACACAC	[18800]
U40452_Pshqipericus		[18900]
<pre>scaffold39565_10.7_Plessonae</pre>	GACAGGCATTCTGCAGTTGTGGGCTCCCCCTCCACGGCATATGTGGCCTGACAGCAACGCAGGGCCGACAGGCACTTGCCACCTGCTAAGTTCGGTGGCG	[18900]
U40452_Pshqipericus	INTRON-6	[19000]
<pre>scaffold39565_10.7_Plessonae</pre>	GCATGCACATGGGAATCAGGACTAATGCTTGGCTGGGGCTGAGCCGCTGGGCAGGTGGGTCCCCCGGGCAAGTGGGGCCCCCGGGCAACTGCCCAGCAT	[19000]
U40452_Pshqipericus		[19100]
<pre>scaffold39565_10.7_Plessonae</pre>	GCCCATTCTGTAAGATGGCCCTGCATTTGTTCCCATTTAAAAATTTACCTTTGACTCTTGTTTTGGTGACAGCTGTAGAATTTTGGATTTTCTATCACTT	[19100]

U40452_Pshqipericus		[19200]
<pre>scaffold39565_10.7_Plessonae</pre>	TCATGCATGACAAATAGAGACAATGATGCTGATTTACTAAAGGAGCTGATCATTTTTCCACATTTCTTTATGTGAATATTATCTCAGCTATCCAATCAC	[19200]
U40452_Pshqipericus		[19300]
<pre>scaffold39565_10.7_Plessonae</pre>	ATATAAAGCAAATTCCTATTTATCATGATTGGATAAATGAAGTGAGTAATTACATTTTATTGTGACAAATCTCAACTTCTTAAGGGACCATTCACACACC	[19300]
U40452_Pshqipericus		[19400]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTCAGTGCAGTGCATTGCGCTAACACGCATGTTACTGCAACACACGTTAACACACCTTACTTGCAGTGCAGTGCAATTCATTTTTAATGGCAGCCCAA	[19400]
U40452_Pshqipericus		[19500]
<pre>scaffold39565_10.7_Plessonae</pre>	CACATCAAGGCAACACAGTGTTGCAGAGATCCACCATGTTTTGTTATGTGCGACCGTTCATTGTGGGGGCTCTGAGTTGCCAATATTGGTGAATGTCCGAA	[19500]
U40452_Pshqipericus		[19600]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTGTGGATGTGTCCGTATGTTGGCAGCCCATATATGGGTGAAAAAAGTGCTTTAACACAACGCACTAGGGCACCAAGGTCTAAATGGGGAGACAAAC	[19600]
U40452_Pshqipericus		[19700]
<pre>scaffold39565_10.7_Plessonae</pre>	GACAATAAAAGCTTAACAGGGGTTTCTGACTGCATCAAAAGCAAACAAA	[19700]
U40452_Pshqipericus		[19800]
<pre>scaffold39565_10.7_Plessonae</pre>	ATAATAACACAGTAATAAAAATGTTCTTTTATCAACTATTTCAACATTGATATTATTATAATGTTATTTCTGCTTACGAGTACAAGCCTTTTTTCGATG	[19800]
U40452_Pshqipericus	AACACTAGTTCAAGTATCCCAAAGGTTTGGTAATGCAAAAATATGACGATGTGGAAAAAAGTCACCATTGAAAT	[19900]
<pre>scaffold39565_10.7_Plessonae</pre>	CGGTAACTGCTTCTTTCTCATTTATCAGAACACTAGTTCAAGTCTCCCAAAGGTTTGGTAAGGCAAAAATTTGAAGATGTGGAAAAAGTCACCATTGAAAT	[19900]
U40452_Pshqipericus	TGCACATTTGAATGAAGATTGTTGCAAAGGTGATGCCGTGGAATGTATGATTGAAAGG	[20000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCACATTTGAATGAAGATTGTTGCAAAGGAGATGCCGTGGAATGTATGATTGAAAGG <mark>GT</mark> AAGCCTGTGCACCAAACATTTTAATATCTCCTTCATCACA	[20000]
U40452_Pshqipericus		[20100]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTGGGCCTTCTTTACACTTGTGGTTAAGTGGTGGTAAAAATAGGTGGTTGACCTGTGTTACTATCACCCAAAGGCAGCTGTGGGGGGTGAAGCAGGGGGG	[20100]
U40452_Pshqipericus		[20200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGGGTTAATACATATGCTTCGGCAAAAATGATTTCCCCTGTTGTGTGGGCTGTATGGCAAACTAATGCAAAATATTCAAGTGGATGGGGCTGCGCCAC	[20200]
U40452_Pshqipericus		[20300]
<pre>scaffold39565_10.7_Plessonae</pre>	AACCATTAGGCTTACAGTTGTGGCACAGTATGTCCATTGAAAAAAGTCCATTCAACTGT <mark>AAAAAAAAAA</mark>	[20300]

U40452_Pshqipericus		[20400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt CTGCCACCCCTTGCCACTCTGTAAAAAGCGACCCTCCTCAAAATGCTTTTCACAAATAGCCACAAGTGTACATGAGTCCTAAAACTTTTACCTAATACA$	[20400]
U40452_Pshqipericus		[20500]
<pre>scaffold39565_10.7_Plessonae</pre>	GTAAGTAATTTGACATCTGATAAGGATACATGCCTATATCTCTCAAGCCTGTTTTTTTT	[20500]
U40452_Pshqipericus	ATGGAAGCAACGGAACATATTTGTTTAGCTAAGGAAAATTATCAAGCAAACTAAGTGACTGTTG	[20600]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTTATCATGCATATGTATTTTATATCTTTGAC	[20600]
U40452_Pshqipericus	TGCTAAAGGTGTTTTAGAACGAACACCCTGTATTCTTGCCCTACCAAACGAGGAACCTGACTTGCCCATAGAACTGAAAGAATATTATGAAGATGAACAT	[20700]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCTAAAGGTGTTTTAGAACGAACACCCTGTATTCTTGCCCTACCAAACGAGGAACCTGACTTGCCCCTGGAACTGAAAGAATATTATGAAGATGAACAT	[20700]
U40452_Pshqipericus	GTTTGTGAGAACTATCAAAAAGACAAGAGAAAATACCTTGCCCA	[20800]
<pre>scaffold39565_10.7_Plessonae</pre>	GTGTGTGAGAAATATCAAAAAGAAAAGAGAAAAATACCTTGCCCA <mark>GT</mark> AAGTATACCTTTTAATGTTTATACAATTGCCATCAAGTTGGGTGTCCTCCGAAC	[20800]
U40452_Pshqipericus		[20900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt tctaccataacccaaactctgaacttttgcctttttgtactccaatgagcttcgggctggaaactaaccttacatcgtttcagatctgtcagtgactact$	[20900]
U40452_Pshqipericus		[21000]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAATATTACTTATATTATTCTCACAGTATTGCAAATGTGTGCTATATCAATCTTTTCATTTATTT	[21000]
U40452_Pshqipericus		[21100]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAAAAATGATTGATTTTTTTTTTTCTGAAGTTAAAAAGTCTTAATACTGTTTTATATACTATAGCTA <mark>TATTTATTTATTTATTTATTTATTTATT</mark> CGAAAA	[21100]
U40452_Pshqipericus		[21200]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAATGAAAAGAAAAAGAGCAAACAGAACAAACAATGCAGGCTCCAGATAAAAAATTGCCATAGTCATATTATAGTGTGAATACGTATGCGCTGCTATTT	[21200]
U40452_Pshqipericus	(AGATAA)n	[21300]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAGAACGTGATACAAACAATCTTAGTATCTATACTAATAATCAAGGGAC <mark>AGAAAAAAGAAAGAAAGAAAAAAAAAA</mark>	[21300]
U40452_Pshqipericus		[21400]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAACTGTTATAATAGTGACCTCTAACACTAAGATACAACACTAATAGACAAAAAAAA	[21400]
U40452_Pshqipericus		[21500]
<pre>scaffold39565_10.7_Plessonae</pre>	AATTCATCTAATAAAATATTCAATTGTGCAATACACCATTCATGTGATGAAACATTCAGTAAGGATGGTGGAACCATACAAATTAAAAAGGGAAAAAAAA	[21500]

U40452_Pshqipericus		[21600]
<pre>scaffold39565_10.7_Plessonae</pre>	ACAAAGTCCAAAAAATGTGCTCGTAAGTTCAATATCAACGTGTATGTGGTCAATCCACAAAAAGATTGTAATCATTCAT	[21600]
U40452_Pshqipericus		[21700]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAAAAATTGTGATGATAGGACAGAACCACCACCAGCAGGTGAAATCACTGCTCACCTTACAGAAAGACCAATTGGTCAAAACACGCTTAATCACCAGAT	[21700]
U40452_Pshqipericus		[21800]
<pre>scaffold39565_10.7_Plessonae</pre>	AGCTCAAGGGAACTGATCTCTTCCACTGCAATGGTCTCCCTCGCATGAATCACCAGATAAATCCACCGATCAATCTCTGTGGGTTTTCACTTCCAATTTTT	[21800]
U40452_Pshqipericus		[21900]
<pre>scaffold39565_10.7_Plessonae</pre>	GTAAATCAAGTTTCCAGAAGGTTCTGCAGAAGGAAGAAGTGACGCAGCAGAGATAAGGAATCACACATGGCTCCCGCCGGGTTTCAAACTTTGGGGCTGC	[21900]
U40452_Pshqipericus	KibiDr1	[22000]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTTCAAGAGAGGTTTCTTGGGTCCA <mark>TCCTGAAGCGTTTCCTTTTTTTCCATTTTTAATTTGTATGGTTCCACCATACTTTCTGAATGTTTCATCACAT</mark>	[22000]
U40452_Pshqipericus		[22100]
<pre>scaffold39565_10.7_Plessonae</pre>	GAATGGTGTATTGCACAACTGATTATTTGGATTAATTGTTTTAAGCATATATAGGGATTGTTTATGAACAAGAGCACTGATTTTTTGTCTATTATT	[22100]
U40452_Pshqipericus		[22200]
<pre>scaffold39565_10.7_Plessonae</pre>	GCCATAGTTATATTGACAAACAATCACAGTCCACCACTGAAAATTAACATAGTTTACCCTCATAGCATGCCCACTGTACTACAGAATAGGGCCAACATAT	[22200]
U40452_Pshqipericus		[22300]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTATATAAACAAGAAGTTATTCATTACTGACCATATTGACAACAATATTAAATAAGTAGTCTGTCCACATCCATTGGGGAAAGCCCAGAGGTATCTAA	[22300]
U40452_Pshqipericus		[22400]
<pre>scaffold39565_10.7_Plessonae</pre>	CCACAGGGGGCCATAATTTCTTAAACTTGCCAGGGTGTCCTCTGTGTTGAAAAATATACTTTTCCAACCGTAGGGTATCCCCCATTTGGGTAGTCCATTCC	[22400]
U40452_Pshqipericus		[22500]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGCCATAGGAGGGTCAGCCGCCTTTCAATGCCTGAATATAAGCTTATGGGCCTGGAAGAGACAATTGCCTGTTTAGGATTATCCTCAATTGGGAGGTCA	[22500]
U40452_Pshqipericus		[22600]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCAGAATATCTAATAAACATGGTTTATTGTCTGCCTGGATGTTGACCTGGAATACCCTATTAATGGTATTAAGGATACCTGTCCAGTAGAGATGTAACT	[22600]
U40452_Pshqipericus		[22700]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGCACAGTGTCACTGTCTATATTTTTAATATATATATACTTATACACCATTTTTGTTTTCCCATGGACACCTTTAACAAAGGATACCCAGAGCATTATGAGTT	[22700]

U40452_Pshqipericus		[22800]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTAAAGGGGGACACGGCAGAATTTGTTTACTTATGAATACTTGTTGAGAGATGAGCACTATGTAATGATTTGGCGTCTCTCTC	[22800]
U40452_Pshqipericus		[22900]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGGAGGCAGAACCTTGATGCCCTGGGCTCACAGTAAGTGGGATGAATGA	[22900]
U40452_Pshqipericus	Helitron-1_DR	[23000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAGACAACACTAGAGATCGGGTGGATATACAGTATCTCACAAAAGTGAGTACACCCCTCACATTTTTGTAAATATTTTATTATATCTTTTCATGTGACAG	[23000]
U40452_Pshqipericus		[23100]
<pre>scaffold39565_10.7_Plessonae</pre>	CACTGAAGAAATTACAATTTGCTACAATGTAAAGTAGTGAGTG	[23100]
U40452_Pshqipericus		[23200]
<pre>scaffold39565_10.7_Plessonae</pre>	CATTAATGTCTAAACCGCTGGCAACAAAAGTGAGTGTACCCCTAAGGTAAAATTGGGCCCAAAGTGTA <mark>AAAATTTTGTGTGGGCCACCATTATTTTCCAGC</mark>	[23200]
U40452_Pshqipericus		[23300]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTGCCTTAACCTTCTTGGGCATAGAGTTCACCAGAGCTTCTCAGGTTGCCACTGGAGTCCTCTTCCACTCCATGACGACATCACGGAGCTGGTGGA	[23300]
U40452_Pshqipericus		[23400]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTTAGAGACCTTGCGCTCCTTCAACTTCCGTTTGAGGATGCCCTACAGATGCTCAATAGGGTTTAGGTCTGGAGACATACTTGGCCAATCCATCACCTT	[23400]
U40452_Pshqipericus	<i>INTRON-8</i>	[23500]
<pre>scaffold39565_10.7_Plessonae</pre>	TATCCTCAGCTTCTTTAGCAAAGCAGTGGTCATCTTGGAGGTGTGTTTGGGGTCGTTATCATATTGGAATACTGCCCTGAAGCCAGAGTCCCTGAAGGGA	[23500]
U40452_Pshqipericus		[23600]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGAATCATGCTCTGCTTCAGTATGTCACAGTACTTGTTGGCATTCATGGTTCCCTCAAGGAACTGTAGCTCCCCAGTGACGGCAGCACTCATGCAGCCC	[23600]
U40452_Pshqipericus	TCIDR1	[23700]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGACCATGACACTACCATCACCATGCTTGACTGTAGGCAAGACAAATTTGTCTTTGTACTCCACACCTGGTTGCCGCCACACACGCTTGACACCATCTG	[23700]
U40452_Pshqipericus		[23800]
<pre>scaffold39565_10.7_Plessonae</pre>	AACCAAATAAGTTTATGTTGGTCTCATCAGACTACAGTACATGGTTCCAGTAATCCATGTCCTTAGTCTGCCTATCTTCAGCAAAACTTTTGCGGGCTTT	[23800]
U40452_Pshqipericus		[23900]
<pre>scaffold39565_10.7_Plessonae</pre>	CTTATGCATCATCTTAAGAAGAGGTTTCCTCTTGGGACGACAGCCATGCAGACCAATTTGATGCAGTGTGCGGGGTATGGTCTGAGCACTGACAGGCTGA	[23900]

U40452_Pshqipericus		[24000]
<pre>scaffold39565_10.7_Plessonae</pre>	C?CAGCAATGCTGGCAGCACTCATACGTCTATTTCCCAAAGACAAACTCTGGATATGATGCTGAGCACGTGCACTTGAATTCTTTGGTCGACCATGGCGA	[24000]
U40452_Pshqipericus		[24100]
<pre>scaffold39565_10.7_Plessonae</pre>	AGCCTGTTCTGAGTGGAAACTGTCCTGTTAAACCACTGTATGGCTTTGGCCACCTCGCTGCAACTCTGTTTTAGGGTCTTGGCAATCTTCTTATTGCCTA	[24100]
U40452_Pshqipericus		[24200]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCCATCTTTATGTAAAGCAACAATTCTTTTTTCAGATTCTCAGAGAGTTTATTGCCATGAGGTGCCATGTTGAACTTCCAGTGACCAGTATGAGCAAA	[24200]
U40452_Pshqipericus		[24300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGAGAGCGATAACACCAAATTTTACACATCTGCTCCCCATGCACGCCACTAACGGGTCAAATGATACCAGGGAGGG	[24300]
U40452_Pshqipericus		[24400]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTGAGGAGTTATTTTGAGGAGACAGCAAATATACACTGTTAAACAAGCTGTACACTCACT	[24400]
U40452_Pshqipericus	Helitron=1-DR	[24500]
<pre>scaffold39565_10.7_Plessonae</pre>	TCACATGAAAAGATATAATAAAAATATTTTCAAAAATGTGAGGGGTGTACACACTTTTGTGAGATACTGTATAT	[24500]
U40452_Pshqipericus		[24600]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAAAAACACACAGTTCAGAACTGGTCCCATCCAAGAATAGCGAGATGCAGTGGCGGCTGGTGCATTGGAGGCGCCGGACACATAGAGTTCAATGGGGTT	[24600]
U40452_Pshqipericus		[24700]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTTTTCTGAAGCACCTGATTAGAGCCAGAGGCTCTAATAGGCTTCAGAAAAGGGTGGGT	[24700]
U40452_Pshqipericus		[24800]
<pre>scaffold39565_10.7_Plessonae</pre>	GTGACAATAGCGAATGAAAATTCGCTATTGTTACACGGATCCTCCTCTTGGCAGATCAGGAAGCGGGTCCGAGACCCGTTTTCTGATTGTCCGAAAGGCG	[24800]
U40452_Pshqipericus	GA-rich	[24900]
<pre>scaffold39565_10.7_Plessonae</pre>	AACCCATGCCATTGGCCAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG	[24900]
U40452_Pshqipericus		[25000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAGTGCACTGGTCACTCTGCAGTGAAGGGGGCACAATGGTGATTCCTGATGACTCTTGGGATGTATGACATCATTTTGGCCTATGCCAGAAACTAGAAAA	[25000]
U40452_Pshqipericus		[25100]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGCTAAGTTTGAAAGAAGTAAATATAATCTACTTTCCTGTCTATCTA	[25100]

U40452_Pshqipericus		[25200]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTTCCACTTTAAGTGACTGCATGTTTTTTTTTTTTTAAATGATCAAAGTGTTAAAAGTGCATATTTAACGTTCTGGTTATGTCACTTTAAACAGTCATTTGCA	[25200]
U40452_Pshqipericus	CTTTACGCATGATTATTCCAGAAGTCACCAGGAATCATCTCCTCAATCGTGTTTACGAGTTTCTAGAGGATTTGAAATGCTGCTGGA	[25300]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTGTTTCTCCAGCTTTACGCATGATTATTCCAGAAGTCACCAGGAATCATCTCCTCAATCTTGTTTACGAGTTTCTAGAGGATTTGAAATGCTGCTGGA	[25300]
U40452_Pshqipericus	AAAGTGCTGTGCTAGTGCAAACTCTGCTGAGTGTCTCAAGGATGCC	[25400]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAGTGCTGTGCTAGTGCAAACTCTGCTGAGTGTCTCAAGGATGCCGTATGTAT	[25400]
U40452_Pshqipericus	CCTAAATTGTTAGAAGCTGCACTAAAAGAAAATGAAGAGAAATTCTAAAACAGAACTGTGG	[25500]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTTTTTCATTTAAAGTATTTATTTTATTTTCATTTC	[25500]
U40452_Pshqipericus	TGCCTTGGAGAAGCTTGGGTTTAATGATTTTTATATTCA	[25600]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCCTTGGAGAAGCTTGGGTTTATTGATTTTTATGTTCA <mark>GT</mark> AAGTTAAGCGTGTAGCATTTACAATTTTGAACATTTTATCAAAAGTCATATAACATAAG	[25600]
U40452_Pshqipericus		[25700]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAGGTTTCTCTGAAGGAATTACTAAATTTGGTAGAAAACCAAACTAAGGATTGTATAGTGTAATGTAATTGTTTAGTGCTTTT	[25700]
U40452_Pshqipericus	hAT-2_ET	[25800]
<pre>scaffold39565_10.7_Plessonae</pre>	CGGGAACCAGAGCCGGTTTCCACATCACAGGACTCGCAGCCAGTTCACACTGCCCTATGCAAACCGCA	[25800]
U40452_Pshqipericus		[25900]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGATCAGTT <mark>TGCATATTGCAGTGCAAACTGTCAAATGGTGCAGGAATCGGGTCGCATTAGGTGCAATCTGATTCCAGTGCAGACCAAAATAAGGGTCC</mark>	[25900]
U40452_Pshqipericus	hAT-2 ET	[26000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCACCGTTTTGGTCCAGACGTGATGCAAATTCAGCCATATAACCTGTATGGCTGAATTTGCATTGCACAGA	[26000]
U40452_Pshqipericus		[26100]
<pre>scaffold39565_10.7_Plessonae</pre>	GTGCGAATCACATACGACGTCTGGCATCGCGCTTGTGCCAACTAGCACTTAAAGCTGAACTTTGGGAAAACAGCACAATTACTTGTGTTTAGATATATTT	[26100]
U40452_Pshqipericus		[26200]
<pre>scaffold39565_10.7_Plessonae</pre>	CCATGCATTTAAAGCACTCAGAGTTTCTATTCTGCCACTGGACCCACAGATTTTCATATTTTTTAACAAGCTAAACTGAATACAATCAAAAGCAAGTAA	[26200]
U40452_Pshqipericus		[26300]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGCTTAGTGTGCATTATCCTATTTACTGTAAAATCATTACATGAATTTACAAACTTTTAGAAATGTTCTTTAACCACTTGCTTACTAGACACATATACC	[26300]

U40452_Pshqipericus		[26400]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCTTCCTGCCCAGGCCAATTTTCAATTGACAATTCCGCGGTCATGCTACACGACACCTATATGAAATTTTTATCATTTTTATCTTTTCACACAAATAGAGCCAATTCCGCGGTCATGCTACACGACACCTATATGAAATTTTTATCATTTTTATCTTTTCACACAAATAGAGCCAATTCCGCGGTCATGCTACACGACACCTATATGAAATTTTTATCATTTTTATCTTTTTCACACAAATAGAGCCAATAGAGCCAATTCCGCGGTCATGCTACACGACACCTATATGAAATTTTTATCATTTTTATCTTTTTCACACAAATAGAGCCAATAGAGCCAATTCCGCGGTCATGCTACACGACACCTATATGAAATTTTTATCATTTTTATCTTTTTTCACACAAATAGAGCCAATAGAGCCAATTGAAATTTTTTTT	[26400]
U40452_Pshqipericus	A-rich	[26500]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTCTTTTGGTGGTATTTAATCACCGCTGGGTTTTTTATTTTTATGAT <mark>AAACAAACAAACAAACAAAAAATTTAGAAAAAAAAAA</mark>	[26500]
U40452_Pshqipericus		[26600]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAAATAGGTAATTTTTCTCCTTCATTG <mark>ATGTGCGCTGATAAGGCTGCACTGATGGGCACTGATAGGCTGCACTGATGGATACTGATAAGGCGGCACTGA</mark>	[26600]
U40452_Pshqipericus	hAT-2 OG	[26700]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGGTACTGATAGGTGGCACTGATGGGCACTGATGAGGCATGACTGGTGGGCACTGATTAGGCTGCACTGATAGGCACAAGTAGATGGCACTGATAGGTG	[26700]
U40452_Pshqipericus	hAT-2 TS	[26800]
<pre>scaffold39565_10.7_Plessonae</pre>	ACACTTAAAGGCACTGATAGGTGGCACTGATAGGCGGGACTGATGGGCACTGATAGGTTGCACTGATGGGCACTGATTGGCAGCACTGATAGG	[26800]
U40452_Pshqipericus		[26900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGTGGGCACTCATTAGCAGCACTGGTGGCCACTGTGTGGGACTGATTTCCCTCTTACATAAGCCGGTAAGCGGCTTTATTCTTTTCTCTTCATGCTGTCA}{\tt GGTGGGCACTCATTAGCAGCACTGGTGGCCACTGTGTGGGGACTGATTTCCCTCTTACATAAGCCGGTAAGCGGCTTTATTCTTTTCTCTTCATGCTGTCAGCACTGATTTCCCTCTTACATAAGCCGGTAAGCGGCTTTATTCTTTTCTCTTCATGCTGTCAGCACTGATTTCCCTCTTACATAAGCCGGTAAGCGGCTTTATTCTTTCT$	[26900]
U40452_Pshqipericus		[27000]
<pre>scaffold39565_10.7_Plessonae</pre>	GCGTGGGGGGAAAAGAGGGGGTGCTGGCTGGTGGGGGGGGGG	[27000]
U40452_Pshqipericus		[27100]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGAGGACATCCATGTACACCCTTTCTGCATTTTATGCCATCTTTCGGCTATAGTGCGGGCACCAAGTGGTTAAAGGCATACATTTATATATTCCGTGC	[27100]
U40452_Pshqipericus		[27200]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCTGTGAAAGAGGCCTTACATTTTCAATGTGTGGCTTTCCAGTAGCCATTTGGGTTCCTGCCATGGCACAAAAACATAGTGGTAGGGTTCTAGATTGTT	[27200]
U40452_Pshqipericus		[27300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGTGTATGTATAGTAATGTAATTGGATTTGTTTTGTGATAAATTGCAAGGGGTCTTCAAGGGGTGCAGGGGTGCCAGGGGGTGCAGGGGGTGCGAGGGGGGGTGGGGGGGGGG	[27300]
U40452_Pshqipericus	INTRON-10	[27400]
<pre>scaffold39565_10.7_Plessonae</pre>	AATGCACTTTTCTAAAAACTCCAATTAATATAAAAAAGTGCGTAAACTTTTTAACCCTCCTCTTATGTTAGAATTCTGTTACTATCCATGATGTTAGGGTC	[27400]
U40452_Pshqipericus		[27500]
<pre>scaffold39565_10.7_Plessonae</pre>	GGTTTTGACCCGTGTTTTAAATCAGCCATAAAATACCCTCTGAATAATTATTTAT	[27500]

U40452_Pshqipericus		[27600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TATCCTTGAAAAGATTGGTTTTAATATTTTTGGTGTGACCCCTTAGACCTTTCTTT$	[27600]
U40452_Pshqipericus		[27700]
<pre>scaffold39565_10.7_Plessonae</pre>	AACCAGCAACTCTGAAATGTATTGGTCCTATGTTTCTTTACTTTATTTA	[27700]
U40452_Pshqipericus		[27800]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGGGTTGAGAGGTGATTGGCTGTCAGTGTTACTAAGCAGAGAGAG	[27800]
U40452_Pshqipericus		[27900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt CTGGGTCAAAATTGACCCTAACACCATGAACGTCTTAATTTTCACCACAGTATTTTATAATTTAGTGAAAAGGATTATTTTTCTATTACATTGTTTAAATT$	[27900]
U40452_Pshqipericus		[28000]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAGGTTCCTGACAAAGTAAAACAATCTTGATGCAATAAACAAATTTATGTGATACTTATAGACATTCAAAAACCTGAAAACGGGTCGGTTCTGACCCTAA	[28000]
U40452_Pshqipericus		[28100]
<pre>scaffold39565_10.7_Plessonae</pre>	CACAAGAGGAGGGTTAAAGACCCCTCATATAATACAGGCAGG	[28100]
U40452_Pshqipericus		[28200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt cttacatgagtattttctttaatttcccctcactgggccaccacttccactattttagtacatattagcagctattagtttgttt$	[28200]
U40452_Pshqipericus		[28300]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAATGCCTTAAATAAATAATTACAAGAGAGGGGGATAGGGCTCAAACCCTCATTTTGGTTCAAGGGCTGTGAAATTGTGAGGGTAGGCTGAGACTGGTT	[28300]
U40452_Pshqipericus		[28400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf CATTGTTAGCAAAGAATCTGCTAGAATCTGTGAGTGATCCTCACCAACAAATCCAGAAAGAGTTTTTAAACCCTTCCAGAAATAATTTCCAAACCCTTCT$	[28400]
U40452_Pshqipericus		[28500]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCTCCCTTTACTTGCTGTCCTGTGTGATGCCAGTCAAATAAACAGAAAGGGAGAAATATGAAACAGATTGAAAGTTGGGAGAAATCTCATGGCAAGAAA	[28500]
U40452_Pshqipericus		[28600]
<pre>scaffold39565_10.7_Plessonae</pre>	AATTTGACAGATGTTCTAACCCACCCATCTTCCCGATATCCATAACTGAAGGTTCTAACCCTTCTATCCTCTAGTCAAAGCTAGGGGGGAGAAAAAAAGTT	[28600]
U40452_Pshqipericus		[28700]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAGCAGGAGTTGGCATTTTAAGTTGGTTGAAAATATACAACACTTAAATAATGATCTTTGGTCCAGTTGGCCATTAACACTTTGCTAAGAACCGGTTTT	[28700]

U40452_Pshqipericus	GCTGCTTGTTAGATACTTTGGAAAGATGCCTCAGGTAACAGCCCAAACACTTGTTGAATTAACTGGTAGAATGGCCAAAATTGGAGTC	[28800]
<pre>scaffold39565_10.7_Plessonae</pre>	CCTTTTTTCCAGGCTGCTTGTTAGATACTTTGGAAAGATGCCTCAGGTACCAGCCCAAACGCTTGTTGAATTAACTGGTAGAATGGCCAAAATTGGAGTC	[28800]
U40452_Pshqipericus	TACTGCTGTGGCTTGCCTGACAACAAGAAGCAACCCTGTGCTGAAGAAAAG	[28900]
<pre>scaffold39565_10.7_Plessonae</pre>	TACTGCTGTGGCTTGCCTGACAACAAGAAGCAAACCTGTGCTGAAGAAAAG <mark>GT</mark> AAGCTCTTCTTGTTTTCTATCCTTTCAATAGATTTTTATTGATTTTC	[28900]
U40452_Pshqipericus		[29000]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAATAACAAAGGGATAACAACACTCCAACATGGAGGATAACATAGGGATTGGTACAGTAATAACAAAAGGGAGAAAGAA	[29000]
U40452_Pshqipericus		[29100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTTCTGTATAATATCTAGTATTGGGCAGATTTATTTCCTAAACTTGAAACACAATGAAATATTCTGGGTAGGATATATTGGATATAGGACATATATTGAA$	[29100]
U40452_Pshqipericus		[29200]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTGATGGTTCTAAAGTCTAAATGTTTTTTACTGTAATGCATTCCCTGCATTAAGGTAAAAAATGTCCCACTACTTTTTGTGCCCCCACCCA	[29200]
U40452_Pshqipericus		[29300]
<pre>scaffold39565_10.7_Plessonae</pre>	GACTCCTCCACCGGTCCAGCACTGTCCTATCTGCTCCTTCTTCCTGGTCTCACAGGAGACTCTGAGAGCTGCGGCAGCCATATGCCGCTGTCAGTCA	[29300]
U40452_Pshqipericus		[29400]
<pre>scaffold39565_10.7_Plessonae</pre>	CCTGTAAGGGGGTACAGGGGGGGGGG	[29400]
U40452_Pshqipericus		[29500]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGGGGGACCGCCAATGAAAAGGTTAGCCACTGCTTTATGCACTGTTGTTGCACAGAGAAGGCAAGTATAACCCCAAGTATAACCCCTTTCTTT	[29500]
U40452_Pshqipericus		[29600]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAAAAGAAAAAACCTTTAATATTACTTTAAATTAACATGGCAGCTAATATACAGCATCTGTCTTTTTGGCTACAG <mark>CTCTATGCAAAAATCCTTTCATTG</mark>	[29600]
U40452_Pshqipericus		[29700]
<pre>scaffold39565_10.7_Plessonae</pre>	CAATTTTTTTTTTGGTTTCCTTGCATAAACTGCAAGTTTCATATCCCCCCAACCCCCCACAAAATTTTAATGAGATTTAGGGCTTTGACTTTGCCAATC	[29700]
U40452_Pshqipericus		[29800]
<pre>scaffold39565_10.7_Plessonae</pre>	TATAACCTTTTATTTATTTTTTGAGCCATTCCTTGATGAACTTGCTAGTGATCTGAATCATTG	[29800]
U40452_Pshqipericus		[29900]
<pre>scaffold39565_10.7_Plessonae</pre>	CCTAAAGTGGTTCTAAAGGCAGAATTTTTTTTTTTTTTT	[29900]

U40452_Pshqipericus		[30000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AGTCACGGCCAAACAACCCTATTTTTGATTTGTCAGTGCACCACACATTAGGGTAATTTACTAAAACTAGAGAGTGCAAAATCTGATGCAGCGGTGCATG$	[30000]
U40452_Pshqipericus		[30100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTGGTCAATCAGCTTCTAACTTCAGCTTCTTCAATTAAGCGTTGATAAAAAAAA$	[30100]
U40452_Pshqipericus		[30200]
<pre>scaffold39565_10.7_Plessonae</pre>	ATCAATCAACCCCATTATATAATATGACAGCTTAAATGCCCACCGAAGACCAAATAAAATAAAAGGTATAAGCTATCCCAAGCGCTTAAAATGAGGACCA	[30200]
U40452_Pshqipericus		[30300]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCACCAATAATTATATTGGCTATGCAGCTACTTAGCTTAAAAATATGGAAAAAAA <mark>ATATATATATAATAGATAGA</mark>	[30300]
U40452_Pshqipericus		[30400]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGTAGCGCAAACTTGTCACAAAGCAAGTGCCAAAAGTACTGTGCCAAACTGCACAAAGTCCAAAATGCAATAAATGTCACATAGGCAGCAGGAAAAGCT	[30400]
U40452_Pshqipericus	INTRON-11	[30500]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGAAATGACAAACAGTCCATATTCCTTCACTCTATCTCCTGGGATGACAATATACAGAGACGTGTTTCTCAAAAGGATCACCACCGCCACCGTGCTC	[30500]
U40452_Pshqipericus		[30600]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAGGGGGGAGGAGAATTACCCACTCACCAGATGCTGTGGCCCAACCCACTTTCGTGTTTGAGCATTCAGGCTTATAATGATGGTCACTGGTCCAGAATA	[30600]
U40452_Pshqipericus		[30700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TCCACAATGGGAATCACACCAGCACAGTCAAAAGAAGCTCAGCAGCATATGGAGAGAGA$	[30700]
U40452_Pshqipericus		[30800]
<pre>scaffold39565_10.7_Plessonae</pre>	TATTAAATAAAACATAAGGTAGTACACTCACATTTCTTGTATGTCATCAAACACAAATAACAGTCACAAATAGGAGAGGAATGCCCTTCGGGAGATGATG	[30800]
U40452_Pshqipericus		[30900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt tccaggctccacgccgtggacgccgcgcccttccgggtggtccaaactcctcttacaaaacctggttacaccgcctggtcacgccccgacacgtttcgtga}$	[30900]
U40452_Pshqipericus		[31000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTTCCACTTCTTCCCCGGGAGCGTGACCACGGCGGCGTTGGCGGTAACCTTAAATACCCATGCCCAACACCCCACAGGGCGGGGCCTGGATTGACAACATC}$	[31000]
U40452_Pshqipericus		[31100]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCCATTGGTGTTTACTCCGGTGTATTGACCAATAACAGACTAGATTTACAGGATCACTTCCTCTCAATCTACTAATCTCCATGGTAAAACACCCCTATGA	[31100]

U40452_Pshqipericus		[31200]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCGGAAATAGGGTGGAGTCGAGGTGGATACCTGTAGAATACTGGCCAATAGCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGAATACTGGCCAATAGCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGAATACTGGCCAATAGCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGAATACTGGCCAATAGCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGAATACTGGCCAATAGCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGAATACTGGCCAATAGCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGACAATGACCGGGCTGAGACAACACAATGACCCTCTATCTCGCCGCTATTCACTGTAGACAATGACCCTCTATCTCGCCGCTGAGACAACACAATGACCCTCTATCTCGCCGCTGAGACAACACAATGACCCTCTATCTCGCCGCTGAGACAACACAATGACCCTCTATCTCGCCGCTGAGACAACACAATGACCACAATGACCACAATGACCCTCTATCTCGCCGCTGAGACAACACAATGACCACAATGACCACAATGACCACAATGACCACAATGACCCTCTATCTCGCCGCTGAGACAACACAATGACACAATGACCACAATGACACAATGACACAATGACACAATGACCACAATGACCACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATACTGACAATACTGACAATACTGACAATACTGACAATACTGACAATACTGACAAATGACACAATGACACAAATGACACAATGACACAATGACACAAATGACACAATGACACAATGACACAATGACACAATGACACAATGACACAATACTGACACAATACACAATGACACAATGACACAAATGACACAATGACACAATGACACAATGACACAATACACAATGACACAATACACAATGACAAATACACAAATGACACAATACACAATACACAATACACAAATACACAATACACAAATGACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACACAAATACAAAATACAAAATACAAATACAAATACAAAATACAAAATACAAAATACAAATAAAAAA	[31200]
U40452_Pshqipericus		[31300]
<pre>scaffold39565_10.7_Plessonae</pre>	ATAGGCCGTCCTACAACCCGGAAGTATAACACATTGGAAGTTAATTACACCCTATGCCAAAGGCAAAATAACACCAAACCCTCAACGACCAATGGACCGA	[31300]
U40452_Pshqipericus		[31400]
<pre>scaffold39565_10.7_Plessonae</pre>	TGATGCACAAAACAAAGTATGAACGATCACCATCCATCGCAACGCAACGAACGAACGAACGAACGAACGAACGAACGAACGAACGAACGAACGAACGAACGAACGACACCACCCACCCACCCACCCACCCACCCACCCACCCACCCACCCCACCCCCCCCCC	[31400]
U40452_Pshqipericus		[31500]
<pre>scaffold39565_10.7_Plessonae</pre>	TGATTAGCCGCCAACACCCCCCCTAGCCCGGACAATAATACAGAGGCCCCGGATGGACCCCCCCGGACATATACAGAGGCCCCGGACGATATACAGAGGCCCCGGGACGATATACAGAGGCCCCGGGACGATATACAGAGGCCCCGGGCCCGGACGACATACAGGGCCCGGGCCGGAGGGCCCGGGGCCCGGGGCCCGGGGCCCGGGGGCCCGGGGGGCCCGGGGGGGGGG	[31500]
U40452_Pshqipericus		[31600]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGAAAGATCAAGGAGTTTTCATGTGTACATAGTACACTAAGCACAGCAGCAGCGCCATCTACCAATGGACTAGTGGTACAATGTAAGGAAAAATTTGGTAA	[31600]
U40452_Pshqipericus		[31700]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTGTATGGAAATTGGTTTAATCCATGTACCAGAAAATATTAATAATAATAATAAACAAAATCATGTCTCACCAGAAAAAAGAACACGAATCTACTGGAAACT	[31700]
U40452_Pshqipericus	(AAATAT)n	[31800]
<pre>scaffold39565_10.7_Plessonae</pre>	AATAAGCA <mark>AAATTAAAATATAAAATATAAAATATAAAATAAA</mark>	[31800]
U40452_Pshqipericus		[31900]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAGTTTAGGGACATAATTGTGTGCGCGGACACTTTGGAGGATCATGGTCCTCCCAACGCACATCATCAGAAAATTAAAAAATGTAGAATTGGAATTGGA	[31900]
U40452_Pshqipericus		[32000]
<pre>scaffold39565_10.7_Plessonae</pre>	AACTAAAAATCAATAATCACTAATAAAAACAATTTAAATCCAACTCTATATTTAATCCAACAGGGCTCAAAGATTGGGTCTGGAATATCCATTTCGTTTCGC	[32000]
U40452_Pshqipericus		[32100]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTTTGACAATTCCCTGACCCAATTCGCACCCCTCCACGGGGGTGTCACCCGCTCAACACCCCAGAACTGTAGACAACTGGGGTCCCGTTGGTGGTGTGA	[32100]
U40452_Pshqipericus		[32200]
<pre>scaffold39565_10.7_Plessonae</pre>	GCGGAAATGGGCTGGGACACTGTGCTCTTTTTACCCTTGATAATATTGCTGATGTGCTCCCCAACTCTAATTGTCAAAGTACGTTTAGTTCGTCCTATA	[32200]
U40452_Pshqipericus		[32300]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAATGATTTTGCAGGGACACATTAAAGCATACACCACATGTGGTGGTGTCACATGTGAAATGCTTAATGTCAAAGTCCTTGTCAATTGCTGCAACAT	[32300]

U40452_Pshqipericus		[32400]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAAGGACTCATGTCTCCTCAGATTTTGGGGGGGTGTAAGAACATGACTTACATCTTCCACATTTGTGAAACCCTCGTCTGCCCGGAAAGAAGAAGAAGTGTC	[32400]
U40452_Pshqipericus		[32500]
<pre>scaffold39565_10.7_Plessonae</pre>	CGGGGGGGTTAAGGATCTTCTTCACCACTCTGTCCCCAAAACTCGGGGCCCTACGATAAATGAAGGATGGGACATCAGGCAACACAGGCCCCAATACGGCGCAACACAGGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGACGCCCCAACACAGGCCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCCCCAATACGGCGCCCCCAATACGGCGCCCCAACGGCCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCCCCCAATACGGCGCCCCAACGGCCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCGCCCCAATACGGCCCCCAATACGGCCCCCAATACGGCGCCCCCAATACGGCGCCCCAATACGGCCCCCAATACGGCCCCCAATACGGCCCCCAATACGGCGCCCCAATACGGGCCCCCAATACGGCGCCCCAACGGCCCCCAACGGCCCCCCCAATACGGGCCCCCAACGGCCCCCACGGCCCCCAACGGCCCCCAACGGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCAACGGCCCCCC	[32500]
U40452_Pshqipericus		[32600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt tccatctgcaagatgttccaatgggttttgaagatggcttctacttttttgtggtcctcacaatagtcagacaagaagctccattgtttatcgtcagatc}$	[32600]
U40452_Pshqipericus		[32700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt CATTTGTGGGTCTCGTCACATTCTGCAAGCATGCAGCTCGAGGGATATCCAAAATATTTTCAATCTCTTTCTCAAGAGATTGTTCGTCATATCCCTTTTG}$	[32700]
U40452_Pshqipericus		[32800]
<pre>scaffold39565_10.7_Plessonae</pre>	GATAAACCTCTCCTTAAGAACATTAGTTTGGGCAATATAGTCTTCTTCAAATGTACAGTTCCGTCATACTCGCATGAACTGGCCACGC	[32800]
U40452_Pshqipericus		[32900]
<pre>scaffold39565_10.7_Plessonae</pre>	AGACCTGGTCTTTTCCAATCTGTTTAATGACAAACTGTAGGCTTGCTT	[32900]
U40452_Pshqipericus		[33000]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAATCTCTTTTTCCAAATTGTAGATGTTTGCAGTTTGCCACCAACTGTTACAAGAGTTACCTGCAGATCCTGCAGTTAAATGGGTTCTTGGAGACTTAT	[33000]
U40452_Pshqipericus	<i>Tc</i> 1- 13_Xt	[33100]
<pre>scaffold39565_10.7_Plessonae</pre>	AGCATCAAACAGTAGACTCTTGGGCTAACTTTGTCATCCCTGACCAAGCAGTTGTTCGAAATCTACGCCACTCATAGATTTTTTTT	[33100]
U40452_Pshqipericus		[33200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGACTAATTCACACAATTTGACCATTTTTTTACATTTTTTGTCACGCTCATAGGCATCAACAGCCTTCTTTTTAAAGGCCTTAGAGAGCTCTAGTGAT	[33200]
U40452_Pshqipericus		[33300]
<pre>scaffold39565_10.7_Plessonae</pre>	GTTGGCATAATAGCACCATACGTCAGTAGCAAAGTAAACACCAGACCCTAGATTCCTGAATTTCATCTAACAGTTTCCACCCAC	[33300]
U40452_Pshqipericus		[33400]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTCTAATCATTGACACCTAATCTGGATCACTCAGTTCTAATGTTATGGATTTGAAGTAGTGCAAAACGTAAAGATGTACTTGA	[33400]
U40452_Pshqipericus		[33500]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTAGTTTATGACAATGAATGAATTTAAATTTTAGATATCATTTATCCAAGTAATCTTTATATGTAGATTGAAAAGTAAATTCTATTTTTTATTATAA	[33500]
U40452_Pshqipericus	C	[33600]
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<pre>scaffold39565_10.7_Plessonae</pre>	CCATATGCTTGCTTTGAACACACCATTTTTTCATTTAGAAATCCTGTTTAAAATATCCTCACTAAACCTCTCAATGTTTTCTAATATTTTACATCACAGC	[33600]
U40452_Pshqipericus	TGGATATTCTTTTGGGAGAGATGTGTGAAAGGGAAAAGAAAACATTCATAAATGACAATGTTCACCATTGTTGCGTCGACTCATATGCTAATAGAAGACC	[33700]
<pre>scaffold39565_10.7_Plessonae</pre>		[33700]
U40452_Pshqipericus	ATGCTTCACTAAACTTGGACCTTATGCTAATTATGAAGCACCAGTATGGGATGAAAGTAAGCTCCACTTTACTGCCGACATGTGTAAAGGTTCAGCAGAT	[33800]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGCTTCACTAAACTTGAACCTTATGCTAAATATGAAGCACCAGTATGGGATGAAAGTAAGCTCCACTTTACTGCCGACATGTGTAAAGGTTCAGCAGAT	[33800]
U40452_Pshqipericus	GACCAGCTAAAGACGAAACTGGT	[33900]
<pre>scaffold39565_10.7_Plessonae</pre>	GACCAGCTAAAGACGAAACTGGTGTAAGTTTTGGAAAAACAGCCATGAGAGGCTTTAAAGATTGTCACATAACACAATGCATGC	[33900]
U40452_Pshqipericus		[34000]
<pre>scaffold39565_10.7_Plessonae</pre>	ATATGGGTTAACTTGCAATTTTATGGTGTGTATTTTACATGTCTTGTTTTCTTTGTTTCCTGTTTAAACTGTTGTCCTTCCT	[34000]
U40452_Pshqipericus		[34100]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGCACCCTGCCTATTTTTTTTTTTTTTTAATTGCGCGATGCAGGTTATTGGTTTGAACTACCACTGAAAACAATAGTATTTTACATGTCCTGTTCTATGTT	[34100]
U40452_Pshqipericus		[34200]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGATATGAGAAATGTATTCGATTTATCAGGGTTTATTATTAATGAAAGTGCTGCCTAAAAATGTCACCAGGTCCAATTTTTACTGCGTTAAACTC	[34200]
U40452_Pshqipericus		[34300]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTCAAAAAGCAGCATGGGTATTAATGCAACACATAGTGTAAATGGGGCCTTCAGAAGTGTTGTATAAGATTTAAACCCCTGTTAGGTTTGCTGTATCTC	[34300]
U40452_Pshqipericus		[34400]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTTAAGAGATTCAACCTCTGTTAGGAATTCATCTGTGGCTTCCTATGGTGGCACATGGAGAAGAGGAGAAACCCAGAGTGCCAGAGGGGGGACCGGAGC	[34400]
U40452_Pshqipericus	(T)n	[34500]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAGGAGGATTGGGGCTGCTCTGTGCAAAACCATTGCACAGAGCAGGTACGTATGATGTG <mark>TTTTATTTTTTTTTT</mark>	[34500]
U40452_Pshqipericus	<i>INTRON</i> _12	[34600]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTAAAGCAAAGCCTGCAGTGGCAGACCAGGTAAGTCAAGCTTCCAATGCAAATCTGGAGTAAGACTGAAGAGAGAAAATATATGATTGGAGCACCACGT	[34600]
U40452_Pshqipericus		[34700]
<pre>scaffold39565_10.7_Plessonae</pre>	GGATGTATCAAAGTCCACAGATAACAAATTTATTTCCTTAAAAGGTTAGACAAAAAGTATGTAAACAATGGATGCATTTTAGGGGGCTAAGCCCTACCCTC	[34700]

U40452_Pshqipericus		[34800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCATCAAGTGTTATGGTTGTACATGGACCAAAAAAAAATGAATTCTTTGTCCTTTAAGTAGACAGAGGATCTGGGTGATACCTAAGACTGGGTTCATATA$	[34800]
U40452_Pshqipericus		[34900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TATGCAAATTTCCCCGCATCCAATTCGCATAACAGGAGAGTGTGACCTGCTCTCAATGGAGCAGGTACACACAGCTCCAAGGTGGCCACGGTCCACATTG}$	[34900]
U40452_Pshqipericus		[35000]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAAGGGCCTGTGCATCTTTGGTTCCGATTCAAGTGCAAATTCAGGCCAGAATACAGACCTGATTTGCACCTGAATCGGCGAACAGGGATACACCGGACC	[35000]
U40452_Pshqipericus		[35100]
<pre>scaffold39565_10.7_Plessonae</pre>	CCATGCTGTGAACAGCATATATAAACCCAGCCTCAAGGTTGCATTGTGTTCTGTGTTGGCTTGCTGGCAGCTAGATTTTGCAGTGACCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGAGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGGAGAAGCCGTGGAGAAGCCGTGGAGAAGCGAGAAGCCGTGGAGAAGCCGTGGGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCCGTGGGAGAAGCGAGAAGCCGTGGGAGAAGCGAGAAGCCGTGGGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGAAGCGAGGAG	[35100]
U40452_Pshqipericus		[35200]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTAACCCTCCTCTGCCCTGAGGGCTGGTTCACACCAGATGCAGTCCAGTGAATTTTTATTCTGCCTCAAAAAACACAAGCACAGTGTTTAACATGGATGATGATGGATG	[35200]
U40452_Pshqipericus		[35300]
<pre>scaffold39565_10.7_Plessonae</pre>	CAATAGCCCTACTTTACACCAGTGCAGTGCGGTCCAGTGCAGGCAAGAAAAGTAGAACATGTTAAATTTTTTCTGTACAGAATGCCTCTGGAACACAGCA	[35300]
U40452_Pshqipericus		[35400]
<pre>scaffold39565_10.7_Plessonae</pre>	AACACATCACAAATGCCCTGGAATGCATGAAAAAACTTGCATGCA	[35400]
U40452_Pshqipericus		[35500]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTTTGGATTTAGGAGATGCTCATTGATCCCATAAAAACCACACTCTACATCATACCTTTCAACTAGAAGTATTTATT	[35500]
U40452_Pshqipericus		[35600]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGTAACCTTGCAACCTTAAATTGAAACATACATTATGGGTGCATGAAGTCATAAATTTAATTTCTTAACATTGTTTAAAATTATATATA	[35600]
U40452_Pshqipericus	-ATTGTTGGTTGAATTTTTGAAAATGAAGCCTACCTGTGGGAAAGAAA	[35700]
<pre>scaffold39565_10.7_Plessonae</pre>	GGTTGTTGGTTGAATTTTTGAAAATGAAGCCTACCTGTGGGAAAGAAA	[35700]
U40452_Pshqipericus	AGCAGAGAACCAGCAGGCATGCTTTGATGAGAAG	[35800]
<pre>scaffold39565_10.7_Plessonae</pre>	AGCAGAGGACCACCAGGCATGCTTTGATGAGAAG <mark>GT</mark> ATGTAATGTGACTGCTGTGCAATGTGTATAATGCCAACTACAATGGAACACGTGCTTCAGTGAA	[35800]
U40452_Pshqipericus		[35900]
<pre>scaffold39565_10.7_Plessonae</pre>	GGTGTATATTTGCGTTTTGTCTATATCTGTTGTTCATGCAAAAAAGGTTCTGTTTTGCAAAGGAAGAACTGTTAGCTAAATATAAATTGTGGATGTTCTG	[35900]

U40452_Pshqipericus		[36000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATTCCCATATCTACAGTGTTGTCACGAGCAGCTAATCATTTTAAGTGTTATTCAATCAGCATTATCACTCTTACAGTAGATAGTGCTTACAATAATTGGT$	[36000]
U40452_Pshqipericus		[36100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATAAATCCAATGTTATTAGCGATCCGAAGTAAAAGTTAGGGTTTCTTAATGTATAACTAGGGACAAAACTTTTTTTGTTTTGGATAGAGGGGAGAGGGGAGAGGGG$	[36100]
U40452_Pshqipericus		[36200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATTAGAACACCTGTCAGTTTTTATTGCTTTATCATTGAAAGTGAAAGTAATATAAAATCCCAAATTTTGGGTTGTCCCCAGAAAAGTAATTGAGGGGAAA$	[36200]
U40452_Pshqipericus		[36300]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ACTTCCAATGGGGACACTAGTTCTGGTGACCTGGGGGTCCCCAAGGAATTCCCTTAATTTGCAGGGATTTGCTCCCACTTCCTGTTTTGGCTATGGGACACTAGTTCGGGACACTAGTTCGGGACACTAGTTCGGGACACTAGTTCGGGACACTAGTTCGGGGGTCCCCAAGGAATTCCCTTAATTTGCAGGGATTTGCTCCCACTTCCTGTTTTGGCTATGGGACACTAGGGACACTAGTTCGGGACACTAGTTGGGGACACTAGTTGGGGACACTAGTTGGGGACACTAGTTGGGGACACTAGTTGGGGGGTCCCCAAGGAATTCCCTTAATTTGCAGGGATTTGCTCCCACTTCGTTTTGGGCTATGGGACACTAGGGACACTAGTTGGGGGACTTGGCTCCCACTTGGGGACACTAGGGACTTGGCGGGGGGGG$	[36300]
U40452_Pshqipericus		[36400]
<pre>scaffold39565_10.7_Plessonae</pre>	GGAAGTGAATCCCCAATGGGGGAAAAAAAAAA	[36400]
U40452_Pshqipericus		[36500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf AATGTTTTGCCTATAGTTCTACTTTAAGTCATGGGAGAGCTGCTTACCATGCATG$	[36500]
U40452_Pshqipericus		[36600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCAGCTTGGGGAACAATTGAGGTATTGGTGCAGGTGGATGCTTCTTATCTAGGAGGGGTTCATTAGTGGGGGCAATGAGAGGGGCCAAGAAAGGGGCTGTA}$	[36600]
U40452_Pshqipericus		[36700]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGGTGTCATAAAAATGGTTAGATTAATAGAGGTATAGACACATGTAATATATTAAAAACCAAAGTTTATTACACAAAAACTTATTCTTATAAAAACAGAA	[36700]
U40452_Pshqipericus		[36800]
<pre>scaffold39565_10.7_Plessonae</pre>	TACCCATACAATAGAATGAAGCTTCACTCTGAGGTCAACTTCTGCTCCAGCTAAACAGGTTAGGTTGTCCAGTGTTACCTTGAGCTATATGTTGCTGTCTGT	[36800]
U40452_Pshqipericus	<i>INTRON-<u>1</u>3</i>	[36900]
<pre>scaffold39565_10.7_Plessonae</pre>	CACTATGAACACTTTCCTTTTTCATTTTTGCACACCTGTTTATTGTTGGCTGGGCATCTGGCCACTGATGCACCTGTTTGAGTCCACCTTGACCTGGATCCACCTGGATCCACCTGGATCGACCTGGATCCACCTGGATCGACCTGGATCGACCTGGATCGACCTGGATCGACCTGGATCGACTGGATCGACCTGGATCGACCTGGATCGACCTGGATCGACCTGGATCGACTGGATCGACCTGGATCGACTGGATCGACCTGGATCGACCTGGATCGACTGGATGGA	[36900]
U40452_Pshqipericus		[37000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTTGGTCCAATGGGTCTATAAGAGTGGCATATCCCCCATGGTTGCAGTGTGAGGTTGTTTTTTTT$	[37000]
U40452_Pshqipericus		[37100]
<pre>scaffold39565_10.7_Plessonae</pre>	AATATTGCTAAATACAGCCTTCTATAATAGGAGGGTTCATTATCATTTGGGACAAGCTTTCTAATGCATCTATTTGATGCTCCTGCCTTCTCTTAGAGAC	[37100]

U40452_Pshqipericus		[37200]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGGGATTATTGGCCAACTAGGTACAAAATGGCAAGATATGTCTGCTAGATCTTCGTGACTATTTGTTTCCTAAATGTCAACTATACTCAGATTAAAGTC	[37200]
U40452_Pshqipericus		[37300]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAGGTCACCATTTAAAAAAAACATATTAATGTACATCTTTTTTTAGGAACAAAATGTACAAATACAGTATTTTCTTCGACAGGAGCCTGTAGAGCATTG	[37300]
U40452_Pshqipericus		[37400]
<pre>scaffold39565_10.7_Plessonae</pre>	CACCCGAGAACAATGGGTGCAATGTCCAGCTTCAGCAGACACTGCCTAAAGCTCTCTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTCTTAGCTGTAGGCAGACAGTTCTTAGCTGTAGGCAGACAGTTCTTAGCTGTAGGCAGACAGTTCTTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTCTTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTCTTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTTCTTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTTCTTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTAGGCAGACAGTTTCTTGCCATACCACTACATGGGCAGACAGTTTCTTAGCTGTGTGGCAGACAGTTTCTTGGCAGACAGTTTCTTGGCAGACAGTTTCTTGGCAGACAGTTGCAGACAGTTGCAGACAGTGCAGACAGTTGCAGACAGTGCAGACAGTTGCAGACAGTGCAGACAGTTGCAGACAGTTGGGCAGACAGTTTCTTGGCAGAGACAGTTGCAGAGACAGTTGCAGACAGTTGCAGACAGTGCAGACAGTTGCAGACAGTGGCAGACAGTTGCAGAGAGAG	[37400]
U40452_Pshqipericus		[37500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TGTGAATGAACTATGAGCACGTTCATCAGCATGCTCAGAGTTTCCTGATACTACAAGTCCGTTTGTTGTAGTGGAATATGGGACATGTATTTTGTTCATA$	[37500]
U40452_Pshqipericus		[37600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TATTCCTGCTATTTTCAAAATATGACAGTGGGTGCAGGGAGAAGAACCCCCTCCTACTGTCACTGGGTGAAGTGGGTAAGGGGAGAATGCGTAATAG$	[37600]
U40452_Pshqipericus		[37700]
<pre>scaffold39565_10.7_Plessonae</pre>	TAACATGTTACACCCTAATAACGGGTGTAATATGTTACTAATGGTGAACTTATCCATTAACCCACACTAATTAAAATGCTACACAAACGGTATAGTGAAC	[37700]
U40452_Pshqipericus		[37800]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTTTTCTTGATGGCCTATCTAATAAAAATGTACTATATAACAATAAATA	[37800]
U40452_Pshqipericus		[37900]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTCTAGCTGCTTGATAATGCATTTGATGAATGCACACTGGGCCTAATTTACTAAAGATGTGAAATGAAGTTAAGTAATCTTGGTTACCTTTTTAAGGAG	[37900]
U40452_Pshqipericus		[38000]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTCCACCCAAAACTAATCTCCAGTTTTTAGTAAATGCTAAAAAGGTACGCCAACTATCCTTTTCTTGGGAACACTGTTGGTGAGCTATATGCACTGGA	[38000]
U40452_Pshqipericus		[38100]
<pre>scaffold39565_10.7_Plessonae</pre>	GTTCCTGGATCTGCACACCTCTTAAATATGAGGAGGTCTTGCTTTGCTTGGGTTTTTAATGTACTTTATACTGTGTGATGGAGCATGAAGAGGAAAG	[38100]
U40452_Pshqipericus		[38200]
<pre>scaffold39565_10.7_Plessonae</pre>	GAGTTAACATTTTTGGCAGGCAAATCTTACTGCCAAGGTATGTAAGATATGCAATAAACCAGGGGGTTCACATACCCAAACTTTATTTA	[38200]
U40452_Pshqipericus		[38300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCATTTTATTGAATGTTTGACATGAACACAGGAATATCCCATCATACTTTAGTAAATCACAGTTGTCTGTGCATCTTTCAAATACTTGATGCTAAATAG	[38300]

U40452_Pshqipericus		[38400]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTATCTAACGAATATTGGGTTATATGCACAAAAAGAGTGCTTATTAGTTAG	[38400]
U40452_Pshqipericus		[38500]
<pre>scaffold39565_10.7_Plessonae</pre>	AATACCCACTGCTTGGTAGTGCAACTAGCAAGTTATGTTAGATGCCTTATAGGCTGTTGGGTGAAGATGACTTTTATTAAAAGTGACTGAAATGTTTTG	[38500]
U40452_Pshqipericus	AAAGGAGGCCTTCATGAAATAATTAAGGACCA <mark>TTGACATGGAACAG</mark>	[38600]
<pre>scaffold39565_10.7_Plessonae</pre>	TATCTTCTTTAATAATAAATGAATTCTGTTATTTGTGTATATTTT <mark>AG</mark> AAAGGAGGCCTTCATGAAATAATTAAGGACCA <mark>TTGACATGGAACAG</mark> GTAAGTC	[38600]
U40452_Pshqipericus	STOP CODON	[38700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGCAGTGTGCATGAATGGTATATGTATTTGTATTTCTACCTTGTAGACATAAATAA$	[38700]
U40452_Pshqipericus		[38800]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTTTACATATACCATACCTGTGAAAGCTGAAAATCATTGAAGTAGACACCGCGATTATAGTGATTTCTGCTAGTTTTGGGGTCCTGTGCTGAATGGCTG	[38800]
U40452_Pshqipericus		[38900]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCTGCTGCATTGTGAACGCAGGATGTCGCCCTCTCAGCACAAGCACATTTTAGAGACCACTTTGCATTCTCCCAGTGCATGCA	[38900]
U40452_Pshqipericus		[39000]
<pre>scaffold39565_10.7_Plessonae</pre>	GATGAGCTAGGTAGGAGGGTCAGGGACGTCACAATTTCCACCTGCTTCATGCTTTGCATTCACTGAGAAAATGTAAAGCATTCTCTGAATGGCGCCCATG	[39000]
U40452_Pshqipericus		[39100]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAAACGGGCAACCTTATGTGTTCAGAAAGCTGCAGGGCAGCTGCTCAGCATGGAGCCCAGAAGCTACAGTGTCTTGGTCCATCTGCAGTGCCCCTCTGT	[39100]
U40452_Pshqipericus		[39200]
<pre>scaffold39565_10.7_Plessonae</pre>	ACCCCATCTAATTCTGGCTGCTTACTGCAGAGTGCTGAGTTCCACTTCAATCGAATACCTTTTTAAACTTGGTCTGTTTATAATGTACATGTTAAAGCA	[39200]
U40452_Pshqipericus		[39300]
<pre>scaffold39565_10.7_Plessonae</pre>	GAACTTAACTCACTAAAACAAGATTTTCCTATTTTGTCCCTCTTCCCTTCTTACAAGGATGCTAAAAAATGTATTATAATTTTTTACCTTTGTGTT	[39300]
U40452_Pshqipericus		[39400]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTTTTTGTGCTTGTACTTTCTGGTTTCCAGCCTAGGCCTGGCACAGCTGTCTACTGGAATCACTGTATCACAGTTATCCCAGGAGGCAGTGTGTT	[39400]
U40452_Pshqipericus	1NTRON-14	[39500]
<pre>scaffold39565_10.7_Plessonae</pre>	ACCAAGCTGCAACTTTGCTGTGCATGTGCGCACATGCGGTGAATTTCCGCGCATGTGCACAAATGCTAGAGGGTGCTAAAGCGCCCATCTCCCAGTAGTT	[39500]

U40452_Pshqipericus		[39600]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCGGCATTTGTGCACATGTGAGGAACTCCGCTGCATGTGCACAGATGGACTGCCAGTCTTTGCCAGCTCTCAAATACCTGGAAGAGAAAAGTTTAGATG	[39600]
U40452_Pshqipericus		[39700]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCGCAGTATGTAAGAGCAACCTAGGAAAAACAGCACAAAACGAATTAATAATGATACATTTTATTCAGTGCACTCTTTTAGGAAGGGTGTTTAAGGGGAAAAACGAATTAATAATGATACATTTTATTCAGTGCACTCTTTTAGGAAGGGTGTTTAAGGGGAAAAACGAATTAATAATGATACATTTTATTCAGTGCACTCTTTTAGGAAGGGTGTTTAAGGGGAAAAAAAA	[39700]
U40452_Pshqipericus		[39800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TATTTTAGCTGGCATTATAAGTACACATTACTCATCTTCTTTTACCTTTCATATGTACATAATTTTTTTT$	[39800]
U40452_Pshqipericus		[39900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATTCTTTCAGCTGTGTTATGATTGGCATGCTAATCGCAAAGCATGTGTTCTCCTGTTCCTGACACAGGAGGGGAGACAAACCAAACAAA$	[39900]
U40452_Pshqipericus		[40000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGGGGTTTCTCAATTTGTTGCATTCCAGCAACTAATCAGATTCAAACTGCACTTGAAAATTACTGTTGAACCTCCTCTACCACATATCATATGTAGGG	[40000]
U40452_Pshqipericus		[40100]
<pre>scaffold39565_10.7_Plessonae</pre>	ACCAAAGTTCCTCAAACAATGTCAAGCTTCCTTGACCTTGGTACTTTAAGTCTAATGGACCTGACTTATTTAT	[40100]
U40452_Pshqipericus	GCCAAAAGCTGATAGAACAAGACTGTGTGTGAGCCACAGGGTTAAACCATCATCAACTGGTCACTGATACATGTCTCTG	[40200]
<pre>scaffold39565_10.7_Plessonae</pre>	GAACTTTCCTTTCTTCAAACAGGCCAAAAGCTGATAGAACAAGACTGTGTTGAGCCACTAGGTTAAACCATCATCAACTGGTCACTGATACATGTCTCTG	[40200]
U40452_Pshqipericus	AAGACTCAAGTAGAAAAAAAAAAAATCATTTTACTGCAAAAATAAAAAAAA	[40300]
<pre>scaffold39565_10.7_Plessonae</pre>	AAGACTCAAGTAGAAAAAAAAAAAATATTTACTGCAAAAATAAAAAAAA	[40300]
U40452_Pshqipericus		[40400]
<pre>scaffold39565_10.7_Plessonae</pre>	TATAACTGATACTATACAACATAACATAACATAATGAAGAAGTTAAAGCCTCATTTAAAGACTCCGTTTTATAGTCAAATATATGTTTATTCAA	[40400]
U40452_Pshqipericus		[40500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GCTTCACCATATCAAAATGATATAAGAGCCAGTAGTTTGTCAAATGTGAAGGAGTAAATATGGGATCCAACAAGGACAGGGTACCCAACCATATACAAAA$	[40500]
U40452_Pshqipericus	A rich	[40600]
<pre>scaffold39565_10.7_Plessonae</pre>	AATCCCCCAATAAACCGGCATATATTAAACTGG <mark>AAATGATAACAGACAAGAAAACAAAAAGAAAAG</mark>	[40600]
U40452_Pshqipericus		[40700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCTCTGCCCTATGGAAGCAAAAAAGGGAGGATTGACATCCAAGGGCCAGGCCTCAAAGCAATTATAACAATTGTAGAGTAGGTGTATCCCAAGAGATAG$	[40700]

U40452_Pshqipericus		[40800]
<pre>scaffold39565_10.7_Plessonae</pre>	ACAGGGGAGAAGAGGAGAAAGAGTAGACAAAGTACGAGAAGCAAAGAAGAAGAGAGAAAAGATGGTGTGGGGTCCCGAGGGAATCTCGAGGCCAGGAATGAGG	[40800]
U40452_Pshqipericus		[40900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGGGAATAAAAGGAGTAGAATCCTCTTGGATAAGGGGAAGCCCAACACATGATAACTAGAGGGGGCCCAAGGAGTATCCTGGAGCTCCATCTAATAAAC}$	[40900]
U40452_Pshqipericus		[41000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCCGAGGAAGAGGAGAAAAATGTATCCTATCCTGTAAGATACAAGCTGCTTAAAAGAATCTGATGAAAAAATGTATCCAGACATACCAGAGTGGGGT	[41000]
U40452_Pshqipericus		[41100]
<pre>scaffold39565_10.7_Plessonae</pre>	GAATTAAAGACTTTAAACTTTTGATGTTCCAGAATCATACACACATGTTAAAAAAAAAA	[41100]
U40452_Pshqipericus		[41200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTTACATAGGTTCATCTTGTAGCACTGTAACTATCCATATACTATACCTTTGGGATCCCTCTAGAAAATCATCTGGAAATCAATGCAGTGGGCATCACTA}$	[41200]
U40452_Pshqipericus		[41300]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt CTCCACCACTACTCCATTGATCTCTGTTAGCTCATTCCACGTTAGCTCAGTTTCGTACAAAGCAGGTAGTCTGTGAACAAATTACTTTAAGCTGTCAGGT$	[41300]
U40452_Pshqipericus		[41400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTTAAAATATGTATATTTAGCCCCTACACTGCACCTAGTGAATAAACAGCATGGAAACTTAGCTTTGAACTAATATTGAAATGGCCAAAATAATACAATG$	[41400]
U40452_Pshqipericus		[41500]
<pre>scaffold39565_10.7_Plessonae</pre>	TATAGGATATTTAAATACCCCTATTAGAGGTGCCTTTGTCTTGCGATGTTGACATCTTAAAACAAAATGGGCCTAATGAATG	[41500]
U40452_Pshqipericus		[41600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AAATGTTCTTGTTGTCCAAAGTAATCAACTAGCTCTAATGCTTGGGCTATAAACATGACAGTTGTTAACTGATTGCAATTGCTAACTAGAACACTTTTTT$	[41600]
U40452_Pshqipericus		[41700]
<pre>scaffold39565_10.7_Plessonae</pre>	TATATAAGAAACCAAATATAAAGTGTCAATCTATACTGAAATATGGAACCTATAAAATTTTGAAGGCCAGGCAATATTTATGAAAGGCATAATCTTAACC	[41700]
U40452_Pshqipericus		[41800]
<pre>scaffold39565_10.7_Plessonae</pre>	AATTGGACATGCACTTCAAGCTACACCTGAAATTTTCAAGGGTACTTCCAGTTGGTAAAGTAAATTCACTATATCAAATTAAAAAAGCAAACAGATATTT	[41800]
U40452_Pshqipericus		[41900]
<pre>scaffold39565_10.7_Plessonae</pre>	GTCATGAATTTGCTAGTTCTTTCCCAGGTACAGAGGCGGTCAGGCACACAGCAACACTTCACTCATTCTGGCTCAATATTCAGTCTAGATTCTAGAAGAA	[41900]

U40452_Pshqipericus		[42000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCCTTTTATATACTGAGGGGGGGGGG	[42000]
U40452_Pshqipericus		[42100]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGCAGTAGCAGTTACAAGGACATATTGACTAGTACAATCTGAGGATGCACAGCTGAAGACTATATAGTAGTAAGTA	[42100]
U40452_Pshqipericus		[42200]
<pre>scaffold39565_10.7_Plessonae</pre>	AATCAACGGATGTACAGCTGGAGACCAGGCAGGAGTATATAGCCTACAGATAGTCATAGACTTGGACAGTCTATGGGTGCACAGCTGAAGACCATGCTGG	[42200]
U40452_Pshqipericus		[42300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTAAGCACAGCACCCCAGGCATGGCAAGGCCACAAAGCCAGGCAGCAGCAGGCAGGCAGGCAGGGGAGGGGGGGGGG	[42300]
U40452_Pshqipericus		[42400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGGGATCAGCCTCCTGGAGCTCACTGATCAATTCCCCACTGGGAAAGAGATGATCTACCCTGAACTCCAGCCTATTTAACAGGAACCCATCCACACAGGG}$	[42400]
U40452_Pshqipericus		[42500]
<pre>scaffold39565_10.7_Plessonae</pre>	GCATATCTACCACGAGGCAAACAACTAGGCAGAAGGAACACTATAAACATGCCCAGACTGGCAGTAGGGTACACTGACCATTTACTAGGTGGGCCGGGCT	[42500]
U40452_Pshqipericus		[42600]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAAGGCCACATGTCCTACCAAGAACTTTCTTCCCCCAGCTCCCCTTTCACTCAC	[42600]
U40452_Pshqipericus		[42700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf ACCAGTCAGAGGTTACAGCACTCTGCAGAGCCCTGTTCACTGTATCCAATGAGGACGCTGCTGACATCTCCATCCTACCAGGTGTGCCTGGCTCT}$	[42700]
U40452_Pshqipericus		[42800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt CTGTACTGTGTGTGTGTGTCAGCGAGTGCTCAAAGGAGCATGAAGAAGGTGGCGGAGAAAGGGGGGGG$	[42800]
U40452_Pshqipericus		[42900]
<pre>scaffold39565_10.7_Plessonae</pre>	GCACTACTCTGGTTCACTTCCCCTCCTGGCAGCAGACAGA	[42900]
U40452_Pshqipericus		[43000]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:carged} ATGCAGGTGCCATGATTCATGGGAACTGCATTTCTTGAAGGGGGCGTTGCACCATACATA$	[43000]
U40452_Pshqipericus		[43100]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCAACTAGTAGTAGGAGAAGAGCAGCACTTTCCTGGGCAAGTTACCAGAGATCGGGGTTGTGCCACATGGGAAGAAGAAGGAATGCAGACCTTTGCTGTC	[43100]

U40452_Pshqipericus		[43200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGGTCTGGGGCCCCACTACAAGGGAGAGAGAGAGAGAACTGATGAGAGAGCTGAGCCAGAGCCATGGAGGACCATATGACATGTCTACATCAGGGCTACATCA	[43200]
U40452_Pshqipericus		[43300]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTGTTCTGGCACTGAAGCTGGGAGGTCTGTTGCCCTGTCTGT	[43300]
U40452_Pshqipericus		[43400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TACCCTGCATACCAGGAATGGTGAGGGGAACTACTGAGCTTAACCCCTTGCTGCCCGCATTTAAGTTAATGCTAACAAAGAACTGCCCCTTACTTA$	[43400]
U40452_Pshqipericus		[43500]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCATCAGCCACCCTTTGTGTACAGTTTTAAAGGGACAGCTACATGCACATCGCCTATAGAGAGGGCCCCAACAAAGCCAGCAAATACTGACAGTTAGGCTAGGCTAGGAGAGGCCCCAACAAAGCCAGCAAATACTGACAGTTAGGGCTAGGCTAGGAGAGGCCCCAACAAAGCCAGCAAATACTGACAGTTAGGGCTAGGCTAGGAGAGGCCCCAACAAAGCCAGCAAATACTGACAGTTAGGGCAGGCCCCAACAAAGCCAGCAAATACTGACAGTTAGGGCAGGCCCCAACAAAGCCAGCAAATACTGACAGGTAGGCCAGGAGGCCCCAACAAAGCCAGCAAATACTGACAGGTAGGCCAGGCAGG	[43500]
U40452_Pshqipericus		[43600]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGGTACTTTGCAGTCACTTTCAGGGGTATCCCTTTGTTAGGAACAATGTCAAAGGAAAAGATCCATAAGCTACACATCATGAGCCAGGCCCATGTGCA	[43600]
U40452_Pshqipericus		[43700]
<pre>scaffold39565_10.7_Plessonae</pre>	TGAAATGAATGAACTGCAACCCTGGCCCCTGGGGGGTACTCTTATGGTTCTGGGGGGGGGG	[43700]
U40452_Pshqipericus		[43800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCCAAGGGATGGAGGGAAACACGTTCGGGCGTGGCCTGGCTGG$	[43800]
U40452_Pshqipericus		[43900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TGTGCAGCTTATGGATCTTTTCCTTTAACATTGCTGCAGCTTGGAGCTGCGACAAACTCTCTGTCCTTGCCTGCGCTCACAGCTGTCTAGGTACCCTTAGGTACCTTAGGTACCTTAGGTACCTTAGGTACCCTTAGGTACCTTAGGTACCTTAGGTACCTTAGGTACCTTAGGTACCTTAGGTACCCTTAGGTACCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCCTTAGGTACCTGCGCTGCGACCTAGGTACCTTAGGTACCCTTAGGTACCCTTAGGTACCTGCACGTAGCTGCGACGTAGCTGCGACCTAGGTACCTGCCTG$	[43900]
U40452_Pshqipericus		[44000]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTGGGGAAGAGAAATTTCTGCTGTTTCCAAGGAGCTAGTCTACAGTTTACTAATGAGTTTGAGGCCTGTATGGTACAGTTGAGAATGCCAGAGGCAGAT	[44000]
U40452_Pshqipericus	(CATTG)n	[44100]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAACAAGTTCCAGCCCGCCATCACCTGAGCCTGAAGTAGCCATTGCATTGTACTCATTGCATTGCATTGCATTCTGGGATCCTTCCT	[44100]
U40452_Pshqipericus		[44200]
<pre>scaffold39565_10.7_Plessonae</pre>	CACGGATGTAGGAAAGCTGGTGTAGACCTTTATTAAGAATGGTTCCATACCCTATTACGA <mark>TTGGTACAAAAAATTGCGATTCTTCTCTGGGAAACCTCTT</mark>	[44200]
U40452_Pshqipericus		[44300]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTCTCATAGAGGAAGATGACTTTGACATCTGGGTCGACCAAGCTGTGCAGGCTGTTGAAGAGTGGAAAGGGGTTGACACTGTAAAGAGACAGAGAATAG	[44300]

	Gvpsv-24-1 XT	
U40452_Pshqipericus		[44400]
<pre>scaffold39565_10.7_Plessonae</pre>	TCGAAAGTCTATGGGCATCTGCTTCAGATGTCATCTGAAACTCACAAGGTTGCAGTGCTGCAGGATGTCTGGAAGCTCATCAAGAGTATGGAAGAGGAGA	[44400]
U40452_Pshqipericus		[44500]
<pre>scaffold39565_10.7_Plessonae</pre>	AAGCTGGGCAGAACGAATCTACTTATTCAAGCACACCTTTCAGGGAGAGAAAAGGTTGTCGCCTCAAACAAA	[44500]
U40452_Pshqipericus		[44600]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:constructed} {\tt ATCTTCTAGATCAGAAACTCCAAAACAAGAGAGACCACAGTAACTTGTTGACCTACTCCAAAATGATGAAAATATTCTGGGAAGAAGTGGCTTGGATCACGGC$	[44600]
U40452_Pshqipericus		[44700]
<pre>scaffold39565_10.7_Plessonae</pre>	CAAGAAAATCCAACACTTCTTAGGGACATGGGCAGGGCGAGAGTGGAGCTTCTGTGCACTGTACAGGCCATCCTGGGTGCACCAGAAGGGAAGAA	[44700]
U40452_Pshqipericus		[44800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AGCCATGTTTAATATTACCAGAATCCCTGACAAGCTAATAGGCCCTGTGTACACCCTCCCAATAAAGGGCTGCATGCGCAGGGTGCACTCAGGGACACTT}$	[44800]
U40452_Pshqipericus		[44900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGAATAGTCATTGCCAAGGGTTACCGTGAGACTTTTACTTTACGCCATTAATGCTTTCGCGGTTGATCTAGTTTTTACTAATGCTGTGTTTAATGGTGT$	[44900]
U40452_Pshqipericus		[45000]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAGCCTTATTGCGATGTAGGTAGGTGGGGGGGGGGGGG	[45000]
U40452_Pshqipericus		[45100]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCATTTACCCTTCAATAATACACCTTATCCTATTAATTCTCAAGTTGCCTGTGGCCTATTTTCTGTGCATTCTTGTAGTTTGGTAACGTTTAAGTT	[45100]
U40452_Pshqipericus		[45200]
<pre>scaffold39565_10.7_Plessonae</pre>	ACACAGATGAGAGCGTCCTTCAGTCTATAACCAGGTGTATCAGTGGGGACTGAGCAGTGTAAGGAGTCGTGGCAGAACAGAGGACCCCTCAAACCAAGCA	[45200]
U40452_Pshqipericus		[45300]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTCCTGTGGGGGTAGCGCTACATTAGAATAAAATTCCTAACCCGGAGATATTGAAACAGTTCTGTACAGAGAGCTCATACTTATCATGAAAAAACTAAA	[45300]
U40452_Pshqipericus		[45400]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGGTTCATGCCCTCTGAATTAAATTGTATATAGGGGGGGTATTATTTGATTAAATCCAGTGTTTACATGACTATGTATG	[45400]
U40452_Pshqipericus		[45500]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCAGTAAGATGAGATAACCAGAGTACTTAGTGTTGTCCCAAACTTTCAGGGAGTGAGGTATATTTGGGGGTTTGCAGTCTAGGGGTTCTAATTCCATGGC	[45500]

U40452_Pshqipericus		[45600]
<pre>scaffold39565_10.7_Plessonae</pre>	CTAATAGAACTCTAAGGGTTTATATTTCCATTGCCTCAATGCATGC	[45600]
U40452_Pshqipericus		[45700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGCCAGATAGTACAACCTACTTTAAAAAGTTTTGAATACCTAAACCAGCCTTTTTTCAAACGATGCAGGGACATTTTTCCTTCC$	[45700]
U40452_Pshqipericus		[45800]
<pre>scaffold39565_10.7_Plessonae</pre>	GCAGGGGTATCTCTTTATTTTAGTGACAGGGCCAATTATTTTACCACATTGATGCTGGGGCTTTTCTTTC	[45800]
U40452_Pshqipericus		[45900]
<pre>scaffold39565_10.7_Plessonae</pre>	AGAATGTGTACTTATCATGTTGCACATGCATTGTTGTATGTTTTCCTTTAAATAATAATATCATCCTTTTACTCTGTTACTCTGTAGCTAGGGAATGTAG	[45900]
U40452_Pshqipericus		[46000]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAGTCTCAGCATTCTTTTCATGCTTATCCTCTATTTACCACATATACTGTATTTTGCTGGAGCTAAACACACTATGATTTTTTTAAATGTCTATCTGT	[46000]
U40452_Pshqipericus		[46100]
<pre>scaffold39565_10.7_Plessonae</pre>	TATGATCTTCACAAGTTTTTTGATTTAGGTTTTCAACAATCTGCACAATGTAATTTGGATGAAGTCACTATTAGGCAAGCTACTAGACTGCAAGTTAACC	[46100]
U40452_Pshqipericus		[46200]
<pre>scaffold39565_10.7_Plessonae</pre>	ACTTAAGGATCGCCCTATAGCAGATATCCTGCACTTCTGTCTAGGAGGGGGCGCACACGCCACTGGATGGCTGCTTTTGCTGTGATCATTCACAGCA	[46200]
U40452_Pshqipericus		[46300]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAGTCAATTGGCATGTGCCCGCTGACATCCATTACCCGGCACTGGCTGCCCGCCGATCAGGAAGAGAGACACAGAACGGAACTCTGCCTATGTAAGCAAGG	[46300]
U40452_Pshqipericus		[46400]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGAGCTCCATTCTGACGGGGTAATCAATGAATTTGGTGCCCCTGCAAAGCAGGGAATAAAATATCTCACTTCCCTTAGTAAAAGCAGCAAGCA	[46400]
U40452_Pshqipericus		[46500]
<pre>scaffold39565_10.7_Plessonae</pre>	ACAAAAATACTGGTTAGGCACACAGTTAACCTTTTGATCGCACTAGACGTTTAACTCCTAACTCATTAGTACAGTGACAGTGCATATTTTTAGCACTGAT	[46500]
U40452_Pshqipericus		[46600]
<pre>scaffold39565_10.7_Plessonae</pre>	CACTTTATTAGTGTCATTGGTTCCCACAAAGTGTCAAAAGTGTCAGTTTGTGTCTGATTGCCCGCTGCAATATCCCCAAATCCCCACCATTACTAGTAAAA	[46600]
U40452_Pshqipericus		[46700]
<pre>scaffold39565_10.7_Plessonae</pre>	AAAAAAAACTTTTGTGCAAACCAATTAATATATGCCTTTTGGGATTTTTGTTTACCAAAAAGGTGTAGCAGAATACATGTTGGCTTTAATTTTTGAAGAA	[46700]

U40452_Pshqipericus		[46800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATTCTTATTTTTCAATTTTGTTATTGGATATACCTAGTTACATAGTTAGT$	[46800]
U40452_Pshqipericus	(A)n	[46900]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAAGTTAAAAATATTGCTTTTTTTATTTTTTTTTTTTT	[46900]
U40452_Pshqipericus		[47000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTGATCAAAATACCACCAAAAGAAAGCTTTATTTGTGGGAACAGTGTTGCATGACTGCAAAATTGTCAGTTAAAGTAACACAGTTTTTTTT$	[47000]
U40452_Pshqipericus		[47100]
<pre>scaffold39565_10.7_Plessonae</pre>	CCGTTTTGTGTGTGAAACTCACTCAAAAGGCTACACCCGGTGCATAAAAAAAGTGCACACAAAACGTGGGAAACACAAAAGTGCACAAGGCGTACACAAAAGTGCACAAAGTGCACAAAGTGCACAAAGTGCACAAAGTGCACAAAGTGCACAAAAAAAA	[47100]
U40452_Pshqipericus		[47200]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGCCTTCTCCGAAGAGGGAAGGCCCTAACCCTGATCCCCGGGGCGCAAGGCTCCTGACAGAGACTGAGGTACTTACT	[47200]
U40452_Pshqipericus		[47300]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGCAGACCCCGTTCTCTGGGAGGACTGCCGAAACAGCCCACCCCAAGTACTAGGTGGGGCTCCCCCGAAAGGAGACCCCATGGGAGAGGGCGAACCTAG	[47300]
U40452_Pshqipericus		[47400]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGAACTCCATATCCACCCCCTCCCAAGACTTTCAGGGCAGGCCGGAGGACCTACGCCATACAAGTCCTGCTGTGAAAGTATACACCGCCACATAAAAA	[47400]
U40452_Pshqipericus		[47500]
<pre>scaffold39565_10.7_Plessonae</pre>	AGGACAAACTGTGAACACACACACACACAGAAAAGAGTGGGGAAGGGGAAGTGGAGAAGTGTGGGGTAGTGTCAAAGTGCAAAACCTTGACAGGCCCCCCGG	[47500]
U40452_Pshqipericus		[47600]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGGAATAAAAATTCCTCTGGTGCCAGCCAAAAGGCCTGTCCCAAGAGGCAGGTGCGAAAAAAGTGTGTATGTGGTGCAGTGACCAAGGTGCCAAAAAC	[47600]
U40452_Pshqipericus		[47700]
<pre>scaffold39565_10.7_Plessonae</pre>	TCAAAGATGAACTGCGCCTCACCAGTGAGAATTACGGCACCCCTGAGCTGCCACCCCAGGGGCACATACACCACAGGCTTTTTTTGGTCCCAGCCTTCGG	[47700]
U40452_Pshqipericus		[47800]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCAGACCAGTGCAGCCTATCCCTACCAGCAGACATCGGAACCTGAACAGCTTAGGGAGAGCCTACCTTCAACTGGGGACAAGCCTTTTGCAAGGCTAGA	[47800]
U40452_Pshqipericus		[47900]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCAACCCAGGAGGAAAAGCCCTCCCATCGCCATCCCAGTTCCTCAGCCAATTACATCCAGGAGCAATACTAACCGGCCTCCAGGACAGACCGGCTTTCT	[47900]

U40452_Pshqipericus		[48000]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCCCCCTTCAAACTGGTAAGAACAGCTTAACCCATTCAAATCAGAAGGCGGCGGACCCTACATGCTTTCCCGGATGACCTACTCCCCATACCCCAGTGA	[48000]
U40452_Pshqipericus		[48100]
<pre>scaffold39565_10.7_Plessonae</pre>	CGTCCAGGCGCTCCGGGTTTTGGTCATACTGGTACTAGCTAAGGTCCATACCAAAGGACAGAACAGATACTACACTTGATCTTAGCCAAAAGGCAGAGA	[48100]
U40452_Pshqipericus		[48200]
<pre>scaffold39565_10.7_Plessonae</pre>	AGCGTTACTTTCTTTTGCAGGCCAAAAAAACGCTAAACCGCTGTCTTGGAATTAGCAGCAGAAGAGATAAAACCTTGAAGGCATGTATATTGGTAGGAAT	[48200]
U40452_Pshqipericus		[48300]
<pre>scaffold39565_10.7_Plessonae</pre>	GTAACATTTTAATTCCTAGCATGAATGTCAATATAAAGATAGAGTGAGATACTGGACATGTGTAAAGTGTGTATCCATGAGGGGGTAATGTACGCCACCAT	[48300]
U40452_Pshqipericus		[48400]
<pre>scaffold39565_10.7_Plessonae</pre>	GTCATCAACTTTAGGTTGTGTTGAGTGTGTGTGTGTAAGTCACCCAGTGTACTTTAATTACAATTTCCAGCATGCCACACTACTCTTTGTAACTGTAAATTTA	[48400]
U40452_Pshqipericus		[48500]
<pre>scaffold39565_10.7_Plessonae</pre>	GTTATGTGTCCCAGTAATATTTTTCCTAACCCTGTAGAGCTACAAGTTCCAGCCATTACACAGTCTGTACCAACAATCACATTCTGTGCCACATATTTTA	[48500]
U40452_Pshqipericus		[48600]
<pre>scaffold39565_10.7_Plessonae</pre>	GACTATTAGCCACCAAAGAAAAGAAAATCCCACAAAAACACGCATCATTAATCACAAAAATATATAT	[48600]
U40452_Pshqipericus	/ATIn	[48700]
<pre>scaffold39565_10.7_Plessonae</pre>	ΑΑΑΑΑΑΑC ΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑΤΑ	[48700]
U40452_Pshqipericus		[48800]
<pre>scaffold39565_10.7_Plessonae</pre>	TATATCTATATAAATTTTCCTTTTGTTCACCAATGTTCATATAGATTGGACAAATTAGGTTTTGTCTAATTCTACAGTTAATCAATGTTGTGTTTTAACTG	[48800]
U40452_Pshqipericus		[48900]
<pre>scaffold39565_10.7_Plessonae</pre>	CTTATTTCTTCGTATATGTGAAATTCAATACACTTATTAAATAAA	[48900]
U40452_Pshqipericus		[49000]
<pre>scaffold39565_10.7_Plessonae</pre>	CAACAAACAAGCACCTTTTATAATCAATAAATACATTAGCTTACAGTCCTTAAAAAAACACTTTATAGAGAGAG	[49000]
U40452_Pshqipericus		[49100]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTATGGTGGGTTAAACCAATATTGTTGGATGGAGAATGTGAAAAACTGGTGACAGCAAAAGGGTTAGGATGACAAGGTTTGCATTTGGAGTGTATA	[49100]

U40452_Pshqipericus		[49200]
<pre>scaffold39565_10.7_Plessonae</pre>	ATCTTTAAAAATCTCTATAACAATACAATACAAAAAAAA	[49200]
U40452_Pshqipericus		[49300]
<pre>scaffold39565_10.7_Plessonae</pre>	CATTAAAATCCAAAATGAATCAAACAATAATAATGTAGGCCAGCGTTTCTAAAAAATTTTATAGTCGGGGCATCCTTTAGAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGTACGCAAAATCTCAAAGAATCTTATAGTCGGGGCATCCTTTAGAAGTACGCAAAATCTCAAAGAATCTCAAAGTACGGGGCATCCTTTAGAAGTACGCAAAATCTCAAAGAATCTTATAGTCGGGGCATCCTTTAGAAGTACGCAAAATCTCAAAGAATCTCAAAGTACGGAAATCTCAAAGAATCTCAAAGTACGGGGCATCCTTTAGAAGTACGCAAAATCTCAAAGAATCTCAAAGTACGGGGCATCCTTTAGAAGTACGCAAAATCTCAAAGAATGTAGGCAAGAAGAATCTCAAAGAATCTCAAAGAATCTCAAAGAATCTCAAAGAATGAAGAATGAAGAATCTCAAAGAATGAAGAATCTCAAAGAATTTTATAGTCGGGGCATCCTTTAGAAGAAGAATCTCAAAGAATCTCAAAGAATTTTATGTCGGGGGCATCCTTTAGAAGAAGAATCTCAAAGAATTTTATGTCGGGGGCATCCTTTAGAAGAAGAAGAAGAAGAAGAAGAAGAAATGAAGAAG	[49300]
U40452_Pshqipericus		[49400]
<pre>scaffold39565_10.7_Plessonae</pre>	CACCCCCGTGTCTAAGCTGTACAAAATCACAAAATATTACTTATCAATGAGAAACCTGAGCCTGGAATGATGCACAGCCTTGATGAAGTTAACTTCACCTTAAGCTGAAGGTGAAGCTGAAGCTGAAGGCTGAAGGCTGGAATGAAGCTGAAGGCTGGAAGGTGAAGGGAAGGGAAGGGAAGGGAAGGGAAGGGAAGGGAAGGGAAGGGAAGGGG	[49400]
U40452_Pshqipericus		[49500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGTTCCCACATCAGAGGTCCCCTCATCATATCACAACCGCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGACCTTTTCTTATCAGAGGTCCCCTTCATGTGGGAGTCCCCTTTAAATCAGAGAGTCCCCTTTAAATCAGAGGTCCCCTTTAAATCAGAGGTCCCCTTTAAATCAGAGGTCCCCTTTAAATCAGAGGTCCCCTTTAAATCAGAGACCTTTTTCTTATCAGAGGTCCCCTTTAAATCAGAGACCTTTTTTTT$	[49500]
U40452_Pshqipericus		[49600]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCCCCATTGCCATCAGATTGGAGTACCCCAATGACAGAGTACCCCATAACATCAGTGTCTCCCCTTCACCACAGAGTTACCCCATCATAGAGCCCCATTAGCAGGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGCAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCCATTAGAGCCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCCATTAGAGCCCCCATTAGAGCCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGCCCCATTAGAGGCCCCATTAGAGCGCCCCATTAGAGGCCCCATTAGCAGGTAGCGCCCCATTAGCAGGAGTAGCGCCCCATTAGCAGGAGGAGGTAGCCCCATTAGCAGGAGGTAGCGCCCCATTAGCAGGAGGTAGCCCCATTAGCAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGA	[49600]
U40452_Pshqipericus		[49700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ACCGTGCCCCCACCCTCGGTGGTAACCCAAACGGCACCCCAGTGGAGAAAAACTGATGTAGACATTTCTTTTCTTTTAATAAACTATGTTTTAACCTAT}$	[49700]
U40452_Pshqipericus		[49800]
<pre>scaffold39565_10.7_Plessonae</pre>	TACATAGATTTTAAGAATTAATACATTACTAAATAAATCACACTGCTTAGTTCTGCATCCTGAGGTACATGTGTTGTTTTTAATATAATATTCAAAGTCT	[49800]
U40452_Pshqipericus		[49900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCCAACTCTTATCTTCTATGGGACTCTGCTAATCAACAGTGCAGCCTCTGTAGTGACTTGAAGTGGTTGTAGAATCTTAACTCTATGCTGTTGAAATAT}$	[49900]
U40452_Pshqipericus		[50000]
<pre>scaffold39565_10.7_Plessonae</pre>	TTACTTTTGCTTCTCTTTATAAAATGTATTTAGACAGCTAAACCGTAGTCTCAGTTTTTTAACTGTTTTCATATTCAGGTATTGAAGCATTATAGACTT	[50000]
U40452_Pshqipericus		[50100]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCATAGATGTACCTAGAAATTAAAATAAATGAATGAATG	[50100]
U40452_Pshqipericus		[50200]
<pre>scaffold39565_10.7_Plessonae</pre>	TACATCATTACCAGTACACTGCTCATATTTAACAAACTCCATATGCCGAAAGTCTGTAGTAAAACTGTGGCCAATTTATGGACATTCAAATATGTGCACA	[50200]
U40452_Pshqipericus		[50300]
<pre>scaffold39565_10.7_Plessonae</pre>	TACTTAACAAACAATAAAAAAAAAATGTTAAGGCCTCGTGCCCTAAAGGCTTAAAATATACTGGGCATAAAATTTCAGTTGTCATCTAAGCACCAATGCT	[50300]

U40452_Pshqipericus		[50400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt tctgtaccatggtagtcacccacgcatacatgtatgaggcttttcccaaacttctgcattcagaaaatatgatgtaccacacggtaacagccaatgcaca$	[50400]
U40452_Pshqipericus		[50500]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAGCATCGTGTATAGGCATAGGGGCTGCCCGGGGCTGGGGGGGGGG	[50500]
U40452_Pshqipericus		[50600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATGCTCAGGTTATATTCAGGCACTCACTGTCTGAGAATATATAAAAAAATCTTTAACAACTATAGTCCAGCCACAGGTGCACTTTAAGTGTAAAACATGA$	[50600]
U40452_Pshqipericus		[50700]
<pre>scaffold39565_10.7_Plessonae</pre>	AAACTGGATTGCTATACTTTATTGTCAAATAAACTGCTGGTACTTACT	[50700]
U40452_Pshqipericus		[50800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TATTGTTTCTAAAAGTGGAAGACTAAACTTAATGTATTGTGCTACCTGCCAAAAGAGGAAAACAAAAAGCATATTTGAAATCTTTAAATAGCTAAGAAGTGAAAGTGGAAGACTAAAAAGCATATTTGAAATCTTTAAATAGCTAAGAAGTGAAAAGAAAAAAAA$	[50800]
U40452_Pshqipericus		[50900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AAGAACTGATGTACCAGTAAGTACTTACATCACTGCTAATCTCCTCTGCCTCTGTCCATGAGGAACATTCTTGCTATGCTTCCTGAAGGACCTTGTAA}$	[50900]
U40452_Pshqipericus		[51000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GAGTCGCTCCCATAAGGGGGCACTCTGTGTCCTTCAGCTTTTTCTTGTGAAAGGTAAATATATTTTAGAATTTTTATGTATATTAT$	[51000]
U40452_Pshqipericus		[51100]
<pre>scaffold39565_10.7_Plessonae</pre>	AACAATGCACCAAAAAGCCTACACCATGTATCAAACCACATGCACTGAATTAATT	[51100]
U40452_Pshqipericus	CR1_1a_Xt	[51200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGTAAAGTTGAATAAAGACACTAGCCCATCCAGTTCGACC	[51200]
U40452_Pshqipericus		[51300]
<pre>scaffold39565_10.7_Plessonae</pre>	TCACTAAGATGCACATCCAAGAGTTTTTTTAAACTATTAATTTTTCCCGCTAACACTATTGATTG	[51300]
U40452_Pshqipericus		[51400]
<pre>scaffold39565_10.7_Plessonae</pre>	ТААААААССТССТАСАСАGCTTAAGGTTCAACCACTTCTCCTCCAATTTTAGTGAGTG	[51400]
U40452_Pshqipericus		[51500]
scaffold39565 10.7 P. lessonae	CCTATGCTAGAATCACCATTGAGGTATCTGTATATCGTTATCATATCTCCTCTCAAGCGTCTCTTCTCCAAGGAGAATACGTTTAGTGCTTGCAGTCGTT	[51500]

U40452_Pshqipericus		[51600]
<pre>scaffold39565_10.7_Plessonae</pre>	CATTGTAACTAAGGTCCTCCAATTGCGTTATTCATTTTGTCACCCTTCTCGGACTCTCTCCAGTTTCAGCACATCATTCCTGAGGATTGGTGACCAGAA	[51600]
U40452_Pshqipericus		[51700]
<pre>scaffold39565_10.7_Plessonae</pre>	CTGAACGGCATACTCCAGATGTGGCCCCACCAGAGTTTTATAAATTGGCAGGATATTTATCTCTGGAGGTAATTCCCTTTTTAATGCATGATAACATTCT	[51700]
U40452_Pshqipericus		[51800]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTAGCTTTGCTTGCTGCAGCTTGACATTGCATGGCATTGCTGAGTCATCCACTATGACCCCCAGATCTTTTTCCTCCTGAATTCCCCCTAGAGATTCT	[51800]
U40452_Pshqipericus		[51900]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCCCTAGAGGGAAGCTTGCGCTTATATTTTAGCCCCCAAGTGTATTACTTTACATTTTTAACATTAAACCTCATTTGCCATGTAGTTGCCCACCCCTT	[51900]
U40452_Pshqipericus		[52000]
<pre>scaffold39565_10.7_Plessonae</pre>	ТАТТТТАТТААGGTCTTCTTTCTATATCCTGCTGTGAAGTTATTGCCCTACATAGTTTTGTATGATCAGCAAAAACTAAGATTGAGCTATTTATCCCAAC	[52000]
U40452_Pshqipericus		[52100]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCTATATCATTTATGAATAAATTAAACAGAATCGGTCCCAAGACAGATCCTTGGGGTATTCCACCTACCACTCCAGACCAGTCTGAGTACACACTATTT	[52100]
U40452_Pshqipericus		[52200]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGACCACCCTTTGGACGTGCCCCTGTAGCCAGTTTTCTATACATAAACAAAC	[52200]
U40452_Pshqipericus		[52300]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGGGACTGTATCGAATGCTTTTGCAAAATCCAGATACACCACATCCACAGGCCTTCCTT	[52300]
U40452_Pshqipericus		[52400]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTGGTTTGTAAAGCAATGCTGATTACTACTAATTTACTGTTTTCATTGGCTAATTATTGTATATAGTCTCCTATCATCCCCCCAAAAGTTTACATACT	[52400]
U40452_Pshqipericus	CR1-2_XT	[52500]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTGATGTTAGCCTGTAGCTTCCAGGAATAGAATAATTATCTCTTAACTTTTTAGATATTGGTACCAGGTTGGCCTTTCACCAATATAACTGGTACCATC	[52500]
U40452_Pshqipericus		[52600]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTGTCAATATGCTGTCCTTAAAAATTAGTCTGGCTTTCACCTGACTGA	[52600]
U40452_Pshqipericus		[52700]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTTAAGCTTTTCTAATTTTCTCTTCTGTTAGCTACAAGGGTACATTTTGAGATGAATTAATGGTACAGTCTTTCCTGATTACCAAATATATAACCCCCC	[52700]

U40452_Pshqipericus		[52800]
<pre>scaffold39565_10.7_Plessonae</pre>	CCTCCTTCCATTGTAAAAACTGAGAAGAAGAATGCATTTAGTACAGTAGCCTTCTGCTTGCCATTTGTAACCAACTTCCCTTCATCTTCCTTTATGGGGC	[52800]
U40452_Pshqipericus		[52900]
<pre>scaffold39565_10.7_Plessonae</pre>	TAATGTGCTCTGAGATTGTTTTTTACTGTTAATATAAAAATGTCTTGGAATTTTCTTTTTACTCTTCTGCTATGTGCCTCTCGTAGTCTATTTTGGCC	[52900]
U40452_Pshqipericus		[53000]
<pre>scaffold39565_10.7_Plessonae</pre>	ACCTTGATTGTGTTCTTACATATCTTATTGCATTCCTTGTAGTGTTGGAATGCTGACAAACACCATTCAACTTTATATTTTTTAAAAGGCCATTTTCTTC	[53000]
U40452_Pshqipericus		[53100]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTTTAATATGACTTTTTACTTTAAAGTTTAGACACCCAGGTTAACCCTCACTCTTTTATGTTTATTGCCTATTGGAATAA	[53100]
U40452_Pshqipericus		[53200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ctttttttttttttttttttttttttttttttttttt$	[53200]
U40452_Pshqipericus		[53300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCTTATCTGAAGGTACACCCACGCGGGCCGATAAATATCGACCCATGCTGATCCCCCCCATCAGATCATGTTTAATGTTCTTAAAGTACTCTCATT	[53300]
U40452_Pshqipericus		[53400]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTCCTCAGTGTTTCAACTTTCTAGGACTTGGTCCTATTTAATACATTGTAGCATTGAATGCAATTTATAAAAGTTGGCTCTTTTGAAATTTAGTGTTCC	[53400]
U40452_Pshqipericus		[53500]
<pre>scaffold39565_10.7_Plessonae</pre>	TGTACTACCCTCCTATCCCCTTTTCCTATGATGTATGCTGAATGTAATAATCCTGTGATTGCTAGTTCCCAATTTGTCCTAAATTTCCACATGTGTGACC	[53500]
U40452_Pshqipericus		[53600]
<pre>scaffold39565_10.7_Plessonae</pre>	AGATCTGTGTTATTTGTAATTAATAGGTTTAGCAAAGCATTCTTTGTAGCCAAGGCACCCCCAATTGGAACATGAAGTTGTCCTGTAAGGTATTTAAGA	[53600]
U40452_Pshqipericus	CR1-2_XT	[53700]
<pre>scaffold39565_10.7_Plessonae</pre>	AATGACGGGCCTTAAATGAGTGAGCTGTTCCTTCTGCCCAGTCAATATTTGGATCATTGAAGTCCTCAATTATGATGACATTGCCCTGCCTTGCCACTAT	[53700]
U40452_Pshqipericus		[53800]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCTAACTGTGATAGGAGGTCCAATTCCTCCAAATCCTTCAGGTTAGGAGGCTTGTAGCATACACACATTATTAATTTCCCACTTTTTCCCCCCATTTGA	[53800]
U40452_Pshqipericus		[53900]
scaffold39565 10.7 P. lessonae	AGTTCCACTCACAGGAATTCTACCTCCCTTGCTCCAATTGCAATGTCATCCCTCATGTTCACTAGCAAATCATTCCTGATATACAGGCACACCCCCT	[53900]

U40452_Pshqipericus		[54000]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCCATCTAGCCTCCGATACAAGGAATACTCCTGGAGAGTTGCCAGCCA	[54000]
U40452_Pshqipericus		[54100]
<pre>scaffold39565_10.7_Plessonae</pre>	AATTAAAAAACAGCTGACAATACTGCAGTTGACATTAAGGGGTTGATTTACTAAAAGTGGAAAGTGCAAAATCTGGTTCAACTCTGCATAGAAACCAAAACA	[54100]
U40452_Pshqipericus		[54200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GCTTCCAGGTTTTTTTTTTTTTTGTCAAAGCCTAATTGAACAAGATGAAATTAGAAACTGATTGGCTACCATGCACTCTCCAGTTTTAGTAAATCAACCCCT$	[54200]
U40452_Pshqipericus	(ATATACA)n	[54300]
<pre>scaffold39565_10.7_Plessonae</pre>	AAATGTATCCCCAAAT <mark>ATATATAAAATACCTATAACCAATATACAGTATAC</mark> TACGCAATATACCTATTTTAGCAAAATATTAATTCCCCTATATGCAAAAC	[54300]
U40452_Pshqipericus	Ā-rich	[54400]
<pre>scaffold39565_10.7_Plessonae</pre>	CTAATTT <mark>ATCAGAATCAATTATAATACATCAAAATGAAAATCAAA</mark> GCAGAGGAGACAGTACATGGTGTATAAAGGCATGGATTGCAAGTGTGGTGTTCCA	[54400]
U40452_Pshqipericus		[54500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AATACACAATATACACACAGATTTGAGTAAATGGTGGGCTGGATATAACATGTTGTAATTTATATATCACTGACATTTATATTTTTGGGATTTGTGTTTTTTTT$	[54500]
U40452_Pshqipericus		[54600]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTTTTTAAATTATCATTGAGAGTTTTTGCAAGTGTGGTGTTAAAAATACAAATGATGCACACGGAATTAAGTAAATGGTGGACTGGATACAG <mark>TATATT</mark>	[54600]
U40452_Pshqipericus	(TATATTA)n	[54700]
<pre>scaffold39565_10.7_Plessonae</pre>	ATATTTTATATACTCAGACATTTTGTTTTCATTTGTGATTTATTT	[54700]
U40452_Pshqipericus		[54800]
<pre>scaffold39565_10.7_Plessonae</pre>	TCACAGTTCGATGTAGCTAGGGCCAATGAGACACAGGTATCAAGACTTTCTATATGGTACACCGGGGAACCTATTTGATCACAATTTTGTATGACCCTTG	[54800]
U40452_Pshqipericus		[54900]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGCTGATTATCACTTTTCCTATCCCAAAACGTGTTGCTAACAGTTCTCCCCCTTGTACAGATTGGACCTGAGTGTCAGTGGAGTTCTAGTATAATGATA	[54900]
U40452_Pshqipericus		[55000]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCAGCATTACATAGACAGTAGTTTCCTTTAGCTGGTTTACTGTGTACAATTTTCTCAGTAGAACAGGATATACTG TATGCACAGTATCTCACAAAAGTG	[55000]
U40452_Pshqipericus	Helitron-1 DR	[55100]
<pre>scaffold39565_10.7_Plessonae</pre>	AGTACAGCCCTCACATTTTTGTAAATATTTTATTATATCTTTTTATGTGACAACACTGAAGAAATGA	[55100]

U40452_Pshqipericus		[55200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GCTTGTATAGCAGTGTATATTTGCTGTCCCCTTAAAAAAAA$	[55200]
U40452_Pshqipericus		[55300]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf G}{\sf A}{\sf A}{\sf A}{\sf A}{\sf T}{\sf G}{\sf T}{\sf G}{\sf A}{\sf C}{\sf T}{\sf G}{\sf G}{\sf A}{\sf G}{\sf G}{\sf G}{\sf G}{\sf G}{\sf G}{\sf G}{\sf G$	[55300]
U40452_Pshqipericus		[55400]
<pre>scaffold39565_10.7_Plessonae</pre>	TGGTTATTGGAAGTT <mark>TGAAAAAAATAATTGTTGCTCTAGATAAAGATTGCCTAGGCTGTAAGAAGAATGCCAAGACCTTGAAACTGAGCTGCAGCTTAGT</mark>	[55400]
U40452_Pshqipericus		[55500]
<pre>scaffold39565_10.7_Plessonae</pre>	GGCCAAGACCACAGAGGGTTTAACAGGACAGGTTCCAATCAGAACAGGCCTCTCCATGGTCGACCAAAGAAGTTGAGTGCACGTGCTCAGCGTCATATC	[55500]
U40452_Pshqipericus		[55600]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGATGTTGTCTTTGGAAAATAAACATATGAGTACTGCCAGCATTGCTGCAGAGGTTGAAGGAGTGGGGTGTCAGCCTGTCAGTGCTCAGACCATATGCC	[55600]
U40452_Pshqipericus		[55700]
<pre>scaffold39565_10.7_Plessonae</pre>	GCACACTGCATCAAATTGGTCTGCAAGAAAGCCTGCAAACAGTTTGCTGAAGACAAGCAGACTAAGAACATGGATTACTGGAACCATCTCCTATGATCTG	[55700]
U40452_Pshqipericus		[55800]
<pre>scaffold39565_10.7_Plessonae</pre>	ATGAGACCAAGATAAACTTATTTGGTTCAGATGATGTCAAGCGTGTGTGGCGGCAACCAGGTGAGGAGTACAAAGACCAGTGTGTCTTCCCTACAGTCAA	[55800]
U40452_Pshqipericus		[55900]
<pre>scaffold39565_10.7_Plessonae</pre>	GCATGGTGGTGGGAGTGTCCGGATCTGGGGCTGTATAAGTGCTGCTGGCACTGGGAAGCCACAGTTCATTAAGGGAACCATGAATGCCAACATGTACTGT	[55900]
U40452_Pshqipericus		[56000]
<pre>scaffold39565_10.7_Plessonae</pre>	GACATACTGAAGCAAAGCATGATTCCCTCACTTCGGAGACTGGGCTCCGGGGCAGTATTCCAACATGATAATGACTCAAAAACACCCTTCAAGACAACCA	[56000]
U40452_Pshqipericus		[56100]
<pre>scaffold39565_10.7_Plessonae</pre>	CTGCCTTGCTAAAGAAGCTGAGGGTAAAGGTGATGGACTGGCCAAGCATGTCTCCAGACCTAAAATCTATTGAGTATCTGTGGGGATCCTCAAAGGGTAG	[56100]
U40452_Pshqipericus		[56200]
<pre>scaffold39565_10.7_Plessonae</pre>	GTGGAGGAGTGCAAGGTCTCTAACATCCACCAGCTCCGTGATGTCGTCATGAAGGAGTAGAAGTAGACTCAAGTGGCAACCTGTGAAGTTCTGGTGAACT	[56200]
U40452_Pshqipericus		[56300]
<pre>scaffold39565_10.7_Plessonae</pre>	CCATGCCCGAGAGGGTTAAGGCAGTGCTGGAAAATAATGGTGGCCACAGAAAATAGTGACACTTTGGGCCCCAATTTGGACATTTCACTTAGGGGTGTAC	[56300]

U40452_Pshqipericus		[56400]
<pre>scaffold39565_10.7_Plessonae</pre>	TCACTTTTGTTTCCAGCAGTTTAGACATTAATGGCTGTGTGTG	[56400]
U40452_Pshqipericus	Helitron-1_DR	[56500]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTACATTGTAGCAAAGTGTA <mark>ATTTCTTCAGTGTTGTCACATGAAAAGATATAATAAAATATTTTCAAAAATGTGAGGGGTGTACACACAC</mark>	[56500]
U40452_Pshqipericus		[56600]
<pre>scaffold39565_10.7_Plessonae</pre>	ATACTGTATAT TCAGTGCAACATTCCACAGCTTTGGAATAAAAAACACACAGTTCAGAACTGGTCCCATCCAAGGATAGCGAGATGCAGTGGCGGCTGGT	[56600]
U40452_Pshqipericus		[56700]
<pre>scaffold39565_10.7_Plessonae</pre>	GCATTGGAGGCGCCGGATGCATAGGGGTTCATGGGGTTTTTTTTTT	[56700]
U40452_Pshqipericus		[56800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt tccggggcgcagagcactgcgccctagcccactcatttgtgtgacaatagcgaatgaaaattcattgttacacggatcctcctcctggcagatcagg}$	[56800]
U40452_Pshqipericus		[56900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AAGCGGGTCCGAGACCCATTTCCTGATTGGCCGAAAGGCGAACCCATGCCATTGGCCAAGGAGGAGGAGGAGGAGGACTTGAGGAGACGCAGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGACGCGGGAGGA$	[56900]
U40452_Pshqipericus		[57000]
<pre>scaffold39565_10.7_Plessonae</pre>	GGGGAGGGGGAGGCACCGCAGGGGGGGGGG	[57000]
U40452_Pshqipericus		[57100]
<pre>scaffold39565_10.7_Plessonae</pre>	GTCACTCTGCATTGAAGGGGGGCACGCTGGTCTCT??????????	[57100]
U40452_Pshqipericus		[57200]
<pre>scaffold39565_10.7_Plessonae</pre>	***************************************	[57200]
U40452_Pshqipericus		[57300]
<pre>scaffold39565_10.7_Plessonae</pre>	***************************************	[57300]
U40452_Pshqipericus		[57400]
<pre>scaffold39565_10.7_Plessonae</pre>	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	[57400]
U40452_Pshqipericus		[57500]
<pre>scaffold39565_10.7_Plessonae</pre>	??????????????????????????????????????	[57500]

U40452_Pshqipericus		[57600]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:construct} AGCCGTCACTGGCGAGATGGAGAAGGGGGGGTCTCGGCTCCTCCAGGTGTTACAGAGTGGTGTCCTTCCATTGATGAGTGTTTTAAACCCCTTCTTATGTGAGTGTCCTTCCATTGATGAGTGTTTTAAACCCCTTCTTATGTGAGTGTGTCCTTCCATTGATGAGTGTTTTAAACCCCTTCTTATGTGAGTGTGTCCTTCCATTGATGAGTGTTTTAAACCCCTTCTTATGTGAGTGTGTCCTTCCATTGATGAGTGTTTTAAACCCCTTCTTATGTGAGTGTGTGT$	[57600]
U40452_Pshqipericus		[57700]
<pre>scaffold39565_10.7_Plessonae</pre>	GAGCTGCCTACTATGTGTGAATAAATGTTTCATCTTAATACTAGACTGAGATTGTGCTTTTCTCCTTTCTTCATGGTCTTTATTGCAGTGATTACAT	[57700]
U40452_Pshqipericus		[57800]
<pre>scaffold39565_10.7_Plessonae</pre>	CCAGCTCTCAGGAACGAGCAGCCATGAGTGGACATTGGACTTTATTGTCTATAAATATTTCCTCAAGAGACTTTTGTTCTTTGGTGCGCCGTTCTCTTTGTTTTGTTCTTTGTTCTTTGTTTTGTTCTTTGTTCTTTGTTTTGTTTTGTTTTTGTTTTGTTTTGTTTTGTTTTT	[57800]
U40452_Pshqipericus		[57900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTTCTCACCCCACATGCACCCCTCAGTATAGGCCCACACTACTCCTTCATCTTGCCTTCTTTTCATCTCACACACA$	[57900]
U40452_Pshqipericus		[58000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AATAAAACTCTTCCGATAGTAACCCCTTCTTACAGACTGGATCATTTGGAGAGAAGGATGACGCCATCAACCCAGTGTAAGCTTCAAAACCACCAAAACATCAAAAACCACCAAAAACAAAAAA$	[58000]
U40452_Pshqipericus		[58100]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAGCTTACAGGAGCAGGAGGGAGGGGGAAGGGGGGACATTGGGGGGACATTGGGGGGGAGGGGGGGGGG	[58100]
U40452_Pshqipericus		[58200]
<pre>scaffold39565_10.7_Plessonae</pre>	$\label{eq:construct} ATGAAATTGCCAATACCAGCTTTTATCAGTTTTACAGACAG$	[58200]
U40452_Pshqipericus		[58300]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCATTTAAATAATCAACACATACTGTGCAACTCACTGTCTTTAGAGCTCTTCTATCCTACTTACT	[58300]
U40452_Pshqipericus		[58400]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCTGCTGTTAGTCCTTGCAATAACATGCCGTGCCTCGCTCACAGCCTCTCTGTGGCAATGCTCCACAATTCCACAAGAGCATGAGCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCTCAATTGGGAATTCCACAAGAGCATGAGCCAATGGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATGCTCAAGAGCATGAGCCTCAATTGGGAATTGCGGAATGCTCAAGAGCATGAGCCTCAATTGGGAATTGCGGAATGCTCAAGAGCATGAGCCTCAATTGGGAATTGCGGAATGCTCAATTGGGAATTGCGGAATGCTCAAGAGCATGAGCATGAGCCAATTGGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATTGCGGAATGCGGGAATGCGGGAGGCGGGCG	[58400]
U40452_Pshqipericus		[58500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGCGATGCCAGACAGGGTTGCCATAGTAATCACAATGAAAATAAGGCTTGGCTACAAACTCACTGTGAATGCATGAGATTACGGCAGTGAGAATTAGAA$	[58500]
U40452_Pshqipericus		[58600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AGGGCGCGGTGAAAATGGAAGATAATACTCAACATGGCACCGGCTTCAGATGAGGAGGAACAAATTACAAACTGCAGCAAATACAAGAGACATCACAAGGACATCACAAGGAGAAATACAAAGAGACATCACAAGGAGAAATACAAAGAGAAATAAAAATAGAAAATAAAAAAAA$	[58600]
U40452_Pshqipericus	Penelope-5-XT	[58700]
<pre>scaffold39565_10.7_Plessonae</pre>	GGTTACAACTATCCCTAAAGAAGCTTGGGGCTCCTAAACACCATCGGTTACCCATAACATGGTGTCTTAACATATTTGCTGCAGTAGTTTGACAACTATT	[58700]

U40452_Pshqipericus		[58800]
<pre>scaffold39565_10.7_Plessonae</pre>	ACCTTAAATATTTAACACCAGCTCCAGGTGTCATGGTACCATGGTGAATGTATTATGGAATATTGGAACATGGTGGTTTTCTTGTTCAATATTTTATTAA	[58800]
U40452_Pshqipericus		[58900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf GATTCAAGCAAAGCAAAAGTCAGAAATACAATTATCTCTGGTTTGTAGAACTTAGCAAGGTGTATGCAACGCACCAAAAAAAA$	[58900]
U40452_Pshqipericus		[59000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATTGCATTGCTCAGGGCTTTTTTTCACTTGGAACTTGGTGGAACTTAGTTCCACCACCTCTGGCTCAGGCCCTCTGCTCCCCGCTCACACTATCACTTGA}$	[59000]
U40452_Pshqipericus		[59100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AAACCCAGAAGTCCAGCTTCTTGGTTTACAAGTGACAGCTTGTCTCCCAATGGCTGCCACAGATATGATCTCCTGAGCAGCTCTTATTGAGTGCAGGATG}$	[59100]
U40452_Pshqipericus		[59200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GGGACAAGGGGGATGCAGGTGCCGTGCGGATGCTGACACCCCCACTGGATAATCTCCCTGCAAGTGAGAGAGTCGGAGCCACCTACATTGTGGTCGTGGTGTGGTGGTGGTGGTGGTGGTGGTGGTG$	[59200]
U40452_Pshqipericus		[59300]
<pre>scaffold39565_10.7_Plessonae</pre>	TGAGTCCCCCTTTCTGGGAAAAAAAAAGCCCTGGCATATATATATATTTTTTTGTCGGGAAAAAAAAAA	[59300]
U40452_Pshqipericus		[59400]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATTCGTGACCTGACCTAGCAACCAAGAACAGAGTCTATAAAATGAAGGGGTCAGGTGTGATAAGCCGGTGTTCTAAAATTTAGTCGGACCCGTGAAAATT}$	[59400]
U40452_Pshqipericus		[59500]
<pre>scaffold39565_10.7_Plessonae</pre>	CCGGACCCTCATCACTTCCGGTCACTTCCGGTGATACTACGTCACCACCAGCTGGAGCGACAGCTGTTGCGAGCCCCGGTGCGCATGCGCACAAGAAATT	[59500]
U40452_Pshqipericus		[59600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCCAGGTCCGATGAGATTTTAAGACTTTGCTAACAGCTGACTGA$	[59600]
U40452_Pshqipericus		[59700]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf A} {\sf C} {\sf A} {\sf C} {\sf C} {\sf C} {\sf C} {\sf A} {\sf C} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf T} {\sf G} {\sf T} {\sf C} {\sf G} {\sf C} {\sf A} {\sf C} {\sf G$	[59700]
U40452_Pshqipericus		[59800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTCTGGAATTTTCAGGGGGCCGATGAAATTTTACAACACCGGCGCACACATCAACATCAGTGGTGAGCAACAAGAAGAAACGAAGGGAAGGAA$	[59800]
U40452_Pshqipericus		[59900]
<pre>scaffold39565_10.7_Plessonae</pre>	CTAAAACTGGAGTACACAGAATCTGGTGCAGCTGTGCATGGCGGATACCAGATTTTACTGTTAGCCTAGGTTTGCACTGCCGTAACTTCAAGGTCGCGTG	[59900]

U40452_Pshqipericus		[60000]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ACTTTCGTGTGACTTTGAATCTGGCATCCCTGTGTGACTTCATCGCGGCTTGCATGCA$	[60000]
U40452_Pshqipericus		[60100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ACCTAAAGTAGTGCAGGAACCTTTTCTAAGTCTGAGCGACTTGAGTCACACCGATTAGAACAGTTCCATTTCAGGTAATGGGGGGCGTGACTTTTCATGTG}$	[60100]
U40452_Pshqipericus		[60200]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf ACTTT}{\sf G}{\sf AACTCT}{\sf G}{\sf CACAT}{\sf G}{\sf ACCG}{\sf G}{\sf G}{\sf CACG}{\sf G}{\sf G}{\sf G}{\sf G}{\sf G}{\sf CT}{\sf AAA}{\sf A}{\sf G}{\sf CT}{\sf T}{\sf A}{\sf A}{\sf A}{\sf G}{\sf CT}{\sf G}{\sf A}{\sf G}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf G}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf G}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf G}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf A}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf CT}{\sf CT}{\sf A}{\sf CT}{\sf CT}{\sf CT}{\sf A}{\sf CT}{\sf C$	[60200]
U40452_Pshqipericus	A-rich	[60300]
<pre>scaffold39565_10.7_Plessonae</pre>	GCTGCACCAGATTTAGTGTGCTCTAGTTTTGGTAAATCAACACCCTAGTGTCAAGCGTGTAAAGAGAGAAACATAA	[60300]
U40452_Pshqipericus		[60400]
<pre>scaffold39565_10.7_Plessonae</pre>	TAAGGGGAAGG GGTTCTGGGAAAAAGAAGAGGAGGAGAGTAAAGTGGAGAACATGGTGCTTTTAAGAAAATTTAAGATCCCTGTCACATTCTTGCACAATCAA	[60400]
U40452_Pshqipericus		[60500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\sf CATTTGCCTTGACACATGACACCATGGTGATTGGATTATGACTACTGAGGTACCCGTGGGGCTATAACTGCTGAGGTATGTTGCCATCACTCATTAGAACCCATGACACCACCATGACACCATGACACCACCATGACACCACCATGACACCACCATGACACCACCACCACCACCACCACCACCACCACCACCACCA$	[60500]
U40452_Pshqipericus		[60600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AAGGAAGCCATTTGGACAGGTGCTGGAGAAGTCTGCAGCAGTGACTAACTA$	[60600]
U40452_Pshqipericus		[60700]
<pre>scaffold39565_10.7_Plessonae</pre>	GAGCTTTTCACACTATAACAAGTAACACTATTTGTGAGACTCATGCATG	[60700]
U40452_Pshqipericus		[60800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTCAATCTGTGAGAAAAAAAAAAAAAAAAAAAAAAAAAA$	[60800]
U40452_Pshqipericus		[60900]
<pre>scaffold39565_10.7_Plessonae</pre>	GCACACACACACCACTACCACCACCACCACCACCACCACCACCACCACCACCACCACCACCACCACCACCACCCCCCCCCC	[60900]
U40452_Pshqipericus		[61000]
<pre>scaffold39565_10.7_Plessonae</pre>	$A {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf A} {\sf A} {\sf A} {\sf A} {\sf A} {\sf A} {\sf G} {\sf A} {\sf$	[61000]
U40452_Pshqipericus		[61100]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TGCTCCTTTCTCACAGTGCCCTGCATTTCCTTCATTTTGGATATCATTCGGCTGTCAACGTGTCTGGTATGGCTTCTCTGTTCTTAACCGCTAGCAGAGAGAG$	[61100]

U40452_Pshqipericus		[61200]
<pre>scaffold39565_10.7_Plessonae</pre>	TACTAGCCTACATTATCCCATCATTACCAAGCCATTTTTCAGTTTCAATTATTGTGATACTTTGGCAGACAATTACTCTGTCATGAAACACTGTACAAAT	[61200]
U40452_Pshqipericus		[61300]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ATAAATTTTATGACTTTTTTGGAGCCAGATAAAGCTGATATTTTATCTTTTAGTATACATGGGAAAAAAGACAAACATATTAAAAAAAA$	[61300]
U40452_Pshqipericus		[61400]
<pre>scaffold39565_10.7_Plessonae</pre>	TTAATCTCCGTT AAAAAAAAAAAAAAAATGCCATTACAGTGTAAAATATCCCAAAAAATTACCCATTTTTGAAAGTAGACACCACAATGTGATCACGTTTCTTT	[61400]
U40452_Pshqipericus		[61500]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt TTGCCACAATTTTTTGGAAATTAAGAAATATGCTGGTATTAATGTAGTGAAGACATGAAGCTTGGTAATGGTCCACTAGATACAGTGCACTGGTAATAAA$	[61500]
U40452_Pshqipericus		[61600]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGAGGGAGGAGAGAGGGGGGGGGGGGGGGGGGGGGGG	[61600]
U40452_Pshqipericus		[61700]
<pre>scaffold39565_10.7_Plessonae</pre>	CAGTTACAAGCATGACTCCCATAGGCAAAGTCATGATTGGACTTGTAGTGCTGCAACAGCTGGTGGGTG	[61700]
U40452_Pshqipericus		[61800]
<pre>scaffold39565_10.7_Plessonae</pre>	A GTACATACAGTGGGAGGACTTAGGGGGGCTTAATGTGCAGATGCCCAGGATAAAGGAGGTGAAGGGTTAAAAAAACAGGTCACAGGAGGGAG	[61800]
U40452_Pshqipericus		[61900]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt GTAAAAAAAAAAAAAAAAAAAAACTTTATTGGAATTCGCTGCTGATGTGATATGTACAGTGATCTGGATCGAAAGAATGACTTATTAATTCATTAGTTGCTCTTCC$	[61900]
U40452_Pshqipericus		[62000]
<pre>scaffold39565_10.7_Plessonae</pre>	CTCTCCTACTTCCATTGTGAGAGAGAGGGGAGGGGCACACTACACTGTTTACATACA	[62000]
U40452_Pshqipericus		[62100]
<pre>scaffold39565_10.7_Plessonae</pre>	CCCAATGGTATAGAGATGCTTCAATTGGCTCTGTACGATTTCCATAGAAAGCCCAATGTTTTTTTT	[62100]
U40452_Pshqipericus	hAT-N33 CPB	[62200]
<pre>scaffold39565_10.7_Plessonae</pre>	TGCAGAAATAAGTGTCCACTACTTTTCTGTTTAAAGACATACAGCCAGGGCCGGTGCTTCCACTAGGCAAACTAGGCAGCCGCCTAGGGCGCATTGCCAC	[62200]
U40452_Pshqipericus		[62300]
<pre>scaffold39565_10.7_Plessonae</pre>	CTAGGGGCGCAGCCATGAGTCGGCACTCTCTCGCCATCATTAATGATCCTTGTGAGCTGAGGGGAGAGGCGGCAGTGGGCAAGGAAGAAGGGACAGAGGAAGAAGGGACAGAGGAAGAA	[62300]

U40452_Pshqipericus		[62400]
<pre>scaffold39565_10.7_Plessonae</pre>	AGATAATAGGTGTCTGGGGCAGTGGCAGAACTACCAGGCCTTGGCTTTCCACCGCTGTTTGGGGGGGG	[62400]
U40452_Pshqipericus		[62500]
<pre>scaffold39565_10.7_Plessonae</pre>	***************************************	[62500]
U40452_Pshqipericus		[62600]
<pre>scaffold39565_10.7_Plessonae</pre>	***************************************	[62600]
U40452_Pshqipericus		[62700]
<pre>scaffold39565_10.7_Plessonae</pre>	***************************************	[62700]
U40452_Pshqipericus		[62800]
<pre>scaffold39565_10.7_Plessonae</pre>	***************************************	[62800]
U40452_Pshqipericus	hAT-N33 - СРВ	[62900]
<pre>scaffold39565_10.7_Plessonae</pre>	??????????????????????????????????????	[62900]
U40452_Pshqipericus		[63000]
<pre>scaffold39565_10.7_Plessonae</pre>	GAGGTTAAGGTATATCTAAAGCCAAATCTTTTTCTTTTGTTTTTTGCCATCTGAAGAGGGGGGTATTCTGTCCACTTTCTATACTGCAGACTCAACAGGA	[63000]
U40452_Pshqipericus		[63100]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTTAGAGGATATCTTTCCAATTTGAGGGAAATCCCACTAACAGGGACACAGACAG	[63100]
U40452_Pshqipericus		[63200]
<pre>scaffold39565_10.7_Plessonae</pre>	ATTTTAAAAAGTTGTGTCCTTTGTTAGGATTTAAACACACATTTACTCTTCTAGTCTCGGTTTGCTTCTATATGTTCTTGTTATCTAATATTTCTTAAAG	[63200]
U40452_Pshqipericus		[63300]
<pre>scaffold39565_10.7_Plessonae</pre>	TTATTGTCAGTGTAGTTTTAAAGCAGTCTATTCTAACCATTTTGAGTAAGGTAATAAGTTGCAGCAGTAACATATATAT	[63300]
U40452_Pshqipericus		[63400]
<pre>scaffold39565_10.7_Plessonae</pre>	TTTGGACACACCTTCTCATTCAAAGAGTTTTCTTTATTTTCATGACTATGAAAATTGTAGAGTCACACTTCAAGAGGAAGTCACCTGAAATGTTTTTCAC	[63400]
U40452_Pshqipericus		[63500]
<pre>scaffold39565_10.7_Plessonae</pre>	TTCACAGGTGTGCCCTGTCAGGTTTAATAAGTGTGATTTCTTGCCTTATAAATGGGGTTGGGACCATCAGTTGTGTGCGCAGAAGTCAGGTGGATACAC	[63500]

U40452_Pshqipericus		[63600]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt AGCTGATAGTCCTACTCAATAGACTGTTAGAATTTGTATTATGGCAAGAAAAAAGCAGCTAAGTACAGGAAAAACGAGTGGCCATCATTACTTTAAGAAAT$	[63600]
U40452_Pshqipericus		[63700]
<pre>scaffold39565_10.7_Plessonae</pre>	GAAGGTCAGGTCGGAAAACTTGGAAAGTGCCCCAAGGGCACAAAAAAAAAA	[63700]
U40452_Pshqipericus		[63800]
<pre>scaffold39565_10.7_Plessonae</pre>	${\tt ACCGCCCCAGGAAAGGAAGACCAAGAGTCACCTCTGCTGCGGAGGATAAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTCACCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTAAACAGCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTAAACAGCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTAAACAGCAGCTCAGAATCGCAGGTAAACAGCAGCTCAGATAGTTCATCCGAGTAAATCGCAGCTCAGAAATCGCAGGTAAACAGCAGCTCAGAATAGTTCATCCGAGTAAACAGCAGCTCAGAATCGCAGGTAAACAGCAGCTCAGAATAGTTCATCCGAGTAAACAGTAGTAGTTCATCCGAGTAAACAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGAAATCGCAGCTCAGAAATCGCAGGTAAAACAGCAGCAGCAGCAGCAGCAGAAATCGCAGCAGAGTAGAGTAGTAGAGAAAGTAGAGTAGAAAGAA$	[63800]
U40452_Pshqipericus		[63900]
<pre>scaffold39565_10.7_Plessonae</pre>	TAGAGACCAGGTCAATGGCACACGGAGTTCTAGCAGCAGACACATCTCTAGAACAACTGTTAAGAGGAGACTGTGTGAATCAGGCCTTCGTGGTAGAATA	[63900]
U40452_Pshqipericus		[64000]
<pre>scaffold39565_10.7_Plessonae</pre>	TCTGCTAGGAAACCACTGCTAAAGAAAGGCAACAAGCAGAAGAAGAGACTTGTTTGGGCAAAAGAACACAAGGAATGGACATTGGACCAATGGAAATCTGTGT	[64000]
U40452_Pshqipericus	[64083]	
<pre>scaffold39565_10.7_Plessonae</pre>	TTTGGTCTAATGAGTCCAAATTTGAGACCTTTGGTTCCAACCACCGTGTCTTCGTGCGACGCAGAAAAGGTGAACGG [64083]	

	Species	1	2	3	4	5	6	7	8	9	10
			0,019	0,021	0,019	0,004	0,021	0,136	0,137	0,458	0,469
1	P. lessonae		±0,003	±0,003	±0,003	±0,001	±0,003	$\pm 0,008$	±0,008	±0,011	±0,010
		38		0,006	0,012	0,017	0,006	0,136	0,137	0,455	0,464
2	P. ridibundus	±6,065		±0,002	±0,003	±0,003	±0,002	$\pm 0,008$	±0,008	±0,011	±0,010
		40	11		0,014	0,020	0,010	0,139	0,140	0,459	0,468
3	P. cf.bedriagae	±6,123	±3,417		±0,003	±0,003	±0,002	$\pm 0,008$	±0,008	±0,011	±0,010
		39	23	27		0,019	0,012	0,138	0,139	0,458	0,467
4	P. shqipericus	±6,160	±4,955	±5,589		±0,003	±0,003	$\pm 0,008$	±0,008	±0,011	±0,010
		8	34	38	37		0,020	0,136	0,137	0,461	0,471
5	P. bergeri	±2,702	±5,910	±6,417	±6,446		±0,003	±0,007	±0,007	±0,011	±0,010
		37	11	17	21	34		0,142	0,143	0,451	0,462
6	P. epeiroticus	$\pm 5,849$	$\pm 3,060$	±3,755	±4,876	$\pm 5,849$		$\pm 0,009$	$\pm 0,008$	±0,012	±0,011
		268	268	268	270	265	247		0,003	0,477	0,481
7	R. chensinensis	$\pm 14,746$	$\pm 14,847$	$\pm 14,778$	$\pm 14,907$	$\pm 14,382$	±14,987		±0,001	±0,010	$\pm 0,010$
		270	270	$270 \pm$	272	267	249	5		0,479	0,483
8	R. kukunoris	$\pm 14,589$	±14,695	14,595	$\pm 14,745$	$\pm 14,220$	$\pm 14,739$	±2,284		±0,010	$\pm 0,010$
		887	881	868	884	886	770	927	931		0,123
9	X. laevis	±22,641	±22,564	±22,491	±22,634	±22,297	±21,711	±20,822	±20,882		$\pm 0,007$
		907	897	884	901	904	788	935	938	243	
10	S. tropicalis	±21,166	±20,953	±20,738	±21,124	±20,785	±20,667	±20,247	±20,203	±14,193	

Table 2.5. Calculation of the number of nucleotide differences (lower diagonal) and uncorrected p distance (upper diagonal) and their standart deviations among western Palearctic water frog species and outgroups on the basis of nucleotide sequence of the serum albumin protein coding exons.

		1	2	3	4	5	6	7	8	9	10
			0,048	0,058	0,045	0,011	0,055	0,237	0,24	0,624	0,636
1	P.lessonae		±0,008	±0,010	$\pm 0,008$	±0,004	±0,011	±0,016	±0,016	±0,020	±0,020
		29		0,016	0,02	0,047	0,003	0,233	0,237	0,622	0,629
2	P.ridibundus	±5,072		±0,005	±0,006	±0,009	±0,002	±0,016	±0,016	±0,020	±0,020
		27	12		0,034	0,056	0,020	0,251	$0,255 \pm$	0,616	0,629
3	P.shqipericus	±4,901	±3,419		±0,008	±0,009	±0,007	±0,017	0,018	±0,020	±0,020
		32	9	19		0,045	0,023	0,237	0,24	0,625	0,634
4	P.cf.bedriagae	±5,161	±2,823	±4,125		±0,009	±0,008	±0,016	±0,016	±0,020	±0,020
		6	26	25	31		0,053	0,241	0,244	0,616 ±	0,629
5	P.bergeri	±2,370	±4,807	±4,685	±5,108		±0,011	±0,017	±0,017	0,021	±0,021
		22	1	9	8	21		0,271	0,273	0,607	0,622
6	P.epeiroticus	±4,256	±0,956	±3,028	$\pm 2,648$	±4,211		±0,021	±0,024	±0,024	±0,024
		143	141	143	139	133	108		0,005	0,651	0,658
7	R.chensinensis	±9,859	±9,471	$\pm 9,500$	±9,720	±9,729	$\pm 8,829$		$\pm 0,003$	±0,019	±0,019
		145	143	145	141	135	109	3		0,652	0,658
8	R.kukunoris	±9,934	±9,557	±9,570	±9,810	±9,837	±8,832	±1,686		±0,019	±0,019
		376	375	377	340	340	242	393	394		0,179
9	X.laevis	±12,351	±12,232	$\pm 12,180$	$\pm 12,370$	$\pm 12,556$	±11,923	$\pm 11,458$	±11,452		±0,015
		384	380	383	348	348	248	398	398	109	
10	S.tropicalis	±12,166	±12,130	±12,126	±12,376	±12,557	±12,113	$\pm 11,370$	$\pm 11,370$	±9,163	

Table 2.6. Calculation of the number of aa differences (lower diagonal) and uncorrected p distance (upper diagonal) and their standart deviations among western Palearctic water frog species and outgroups on the basis of aa sequence of the serum albumin protein coding exons.

CHAPTER 3

MANUSCRIPT 2: DISCORDANCE BETWEEN MITOCHONDRIAL AND NUCLEAR PHYLOGEOGRAPHY IN WATER FROG (*PELOPHYLAX*) COMPLEX OF THE EASTERN MEDITERRANEAN REGION

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3.1. Introduction

Terrestrial environments in the eastern Mediterranean region, including Anatolia, has experienced high rates of crustal deformation for many millions of years. Significant events (cf. Steininger and Rögl, 1984) include: the closure of Tethys Sea linking the Mediterranean and the Indian Ocean as a result of uplift of the Arabian Platform; the isolation of the Paratethys Sea from the Global Ocean and its evolution into separate basins, including the modern Black and Caspian seas; the deformation of mountain ranges (e.g., the Pontides, Caucasus and Taurus); and the onset of the Messinian Salinity Crisis (MSC) caused by interruption of the marine connection between the Atlantic Ocean and the Mediterranean Sea. Seyrek et al. (2014) provided an up-to-date synthesis of the crustal deformation since the Late Miocene in a major part of the study region. In addition to geological processes, climatic changes modifying terrestrial environments include significant global cooling and the formation of ice sheets during the Pliocene and Pleistocene

periods (e.g., Ehlers and Gibbard, 2007) and an increase in topographic relief as a consequence of climate change through erosional unloading of the Earth's crust caused uplift (Westaway et al., 2009a) have also caused significant changes in terrestrial environments.

To study how such geological and climatic events have affected species distribution, diversity, and genomes, a variety of mitochondrial and nuclear genetic markers have been used. Although the genome of an organism is subjected to common historical processes, molecular markers do not respond in the same way, resulting in discordance in tree topologies, branch lengths due to heterogeneities (Edward, 2009), times of divergences and phylogeographical patterns among genes. This is not surprising because each type of marker has a specific characteristics such as their inheritance mechanism, effective population size, recombination and mutation rate (Avise, 1994; Ballard and Whitlock, 2004; Currat et al., 2008; Leache, 2009; Petit and Excoffier, 2009).

Many potential mechanisms have been suggested to explain observed discordant patterns in both tree topologies and phylogeographic patterns in natural populations; these include incomplete lineage sorting, gene duplication, horizontal gene transfer, recombination, introgression (Degnan and Rosenberg, 2009), adaptive processes (Plötner et al. 2008), and sex-biased asymmetries (Coyne and Orr, 2004; Maroja et al., 2009,). In this paper we discuss the four most common drivers: incomplete lineage sorting, introgression, adaptive processes and sexbiased asymmetries. Incomplete lineage sorting is the retention and stochastic sorting of ancestral polymorphism, whereby some alleles, more closely related in the gene tree than is expected given the species tree, suggest paraphyletic and misleading relationships among lineages and unpredictable biogeographic patterns among mtDNA and nuDNA markers (Funk and Omland, 2003; Ballard and Whitlock, 2004; Maddison and Knowles, 2006).

The next most frequently observed driver introgressive hybridization, has been defined as exchange of genes between distinct evolutionary lineages (Seehausen, 2004). It has obvious impact on adaptive radiations such as Darwin's finches and African cichlid fish (Freeland and Boag, 1999; Salzburger et al., 2002) and on the formation of new species as seen in *Heliconius* butterflies (Mavarez et al., 2006). In particular, in evolutionary lineages that have diverged for long periods of time and have then come into secondary contact, for example as a result of range expansion from glacial refugia following the last glacial event.

The mode of introgression and its interaction with adaptive and sex-biased processes generally results in asymmetric introgression, which can form the last pattern of biogeographic discordance between mitochondrial and nuclear markers (Toews and Brelsford, 2012). Sex-biased processes include male-biased dispersal, as in *Pelophylax ridibundus* (Holenweg Peter, 2001); female-biased, a fitness loss in hybrids particularly the heterogametic sex when interbreeding occurs between divergent species, for example in eastern Palearctic water frogs, P. nigromaculatus and P. plancyi (Liu et al., 2010); or selection on genes playing a role in mating behaviour, for instance in field crickets, Gryllus firmus and G. pennsyllvanicus (Maroja et al., 2009). In adaptive processes, selection can also favour one group of mitochondrial haplotypes over another in a particular geographic region due to fitness advantage of its own distinct nuclear background. For example, Plötner et al. (2008) showed that, in northern Europe, P.ridibundus with the P. lessonae-specific mt haplotype have a fitness advantage over those with *ridibundus* mitochondria, being less sensitive to oxygen deficiency. All these sex-biased and adaptive processes lead to asymmetric mitochondrial introgression, resulting in a discordant pattern among mtDNA and nuDNA markers.

Water frogs (genus *Pelopyhylax*) in the eastern Mediterranean region represent a genetically and phylogenetically diverse group, made up of lineages of both older origin such as the Middle or Upper Miocene and others with a more recent origin

in the Pliocene and Pleistocene (Plötner et al., 2009, Akın et al., 2010b). Although they are found in all types of freshwater environments, their skin physiology makes them highly sensitive to environmental changes caused by climate change, crustal deformation, and regional scale vertical motions that result in a loss or gain of surface waters needed for their survival and reproduction. Their populations tend to be highly structured genetically and to preserve signals of historical responses to geological and climatic changes (Akın et al., 2010b). They are thus an almost ideal group to study of the effect of geological and climatic processes on patterns of phylogeography and on the molecular evolution of protein coding (mitochondrial) and non-coding (nuclear) genes (Beerli et al, 1996, Plötner et al. 2010, Akın et al., 2010b).

Recent phylogeographic study of eastern Mediterranean water frogs (Akın et al., 2010b) revealed extensive genetic diversity in mtDNA, separated into six main haplogroups (MHGs). Three MHGs exist in Anatolia, the most widespread being distributed from western Anatolia to central Russia; the other two Anatolian MHGs are restricted exclusively to the areas west (the Cilician Plain) and east of the Amanos Mountains. The other three MHGs occur in the Levant, on Cyprus and in Europe. Estimates of divergence times of these groups showed that their separations were well correlated with geological events. How nuDNA reacted during these geological processes is, however, not known. Furthermore, it is not easy to identify any nuDNA marker that reflects true phylogenetic signals. In this study, the composite serum albumin intron-1+RanaCR1 (SAI-1+RanaCR1) was used as a nuDNA marker. It supports some aspects previously identified using mtDNA, including the sister group relationship between P. perezi and P. saharicus, and their joint sister group relationship to other western Palearctic water frogs, and the monophyly of the *P. lessonae* and the Anatolian populations (Plötner et al., 2009). It also indicates that the Anatolian water frogs consist of two distinct groups, and frogs from Cyprus represent a distinct evolutionary species (P. cypriensis Plötner et al., 2012).

Herein, we present a large mtDNA and nuDNA dataset, collected from eastern Mediterranean region especially Anatolia and neighbouring regions. These data reveal that the markers show discordances in the tree topology, the number of distinct genetic stocks, the levels of genetic differentiation, the times of divergence and the patterns of geographic distribution. Our observations suggest that inconsistency between the two marker sets can be explained by distinct processes, including incomplete lineage sorting and retention of ancestral polymorphism because of large effective population size of nuDNA; and introgression after secondary contact acting with sex-biased or adaptive processes linked to the inheritance pattern of mtDNA.

3.2. Material and Methods

3.2.1. Field Trips, Locality Selection and Sampling

Field trips in Turkey (2010-2011) were planned based on distributions of distinct mitochondrial haplogroups (MHG), broadly known from previous findings. Some regions, especially south-western Anatolia, have been well sampled while eastern Anatolia has been represented by samples from few localities (Akın et al., 2010a; 2010b). Because the study area consists of an extensive geographical region, locality selection was conducted in a way that should adequately reveal distributions of genetic lineages. Our main target for locality selection has been sampling river systems homogeneously throughout Turkey.

A total of 1500 tissue samples from 285 distinct localities at altitudes from 0 m to 2276 m were collected from within Turkey. 111 additional samples from 32 distinct localities from Kazakhstan, Armenia, Ukraine, Iran, Jordan, Poland and Germany were also included. Moreover, a total of 691 samples from previous studies (Plötner et al., 2001; 2008; 2009; 2012; Sumida et al., 2001; Akın et al., 2010a; 2010b) were combined for analysis to reveal the distribution pattern

genetic markers distribution of water frogs in the eastern Mediterranean region (Appendices 3.1 and 3.2).

3.2.2. PCR and Sequencing of Mitochondrial and Nuclear Genes

DNA isolation was carried as described by Akın et al. (2010b). The procedure for amplification of two mitochondrial genes (NADH dehydrogenase subunit 2, ND2, 1038 bp; and NADH dehydrogenase subunit 3, ND3, 340 bp) were given by Plötner et al. (2008). For amplification of the nuclear serum albumin intron 1 (SAI-1), including the non-long terminal repeat (non-LTR) retrotranposon RanaCR1, the primer set Ex1-F5 (forward) and Ex2-R2 (reverse) were used following the method described by Plötner et al. (2009). Two additional internal (Int) primers were designed based on the complete sequences of SAI-1 obtained from homozygous individuals, to read the second part of the gene in heterozygous individuals after positions of G or CA repeats where length polymorphism can start. After G repeats, the 630 bp region of the SAI-1 gene was amplified using primer sets Int1F: 5'- CACTCACTAAAACAAGAAGAAAAGC - 3' or Int1Fv2: 5'- CACTCACGAAAATTAGAAGAAAAGC - 3' and Int1R: 5'- TTCAACAG-CTGGTT-TTCCCAC – 3[°]. The 410 bp region of SAI-1 after CA repeats was also amplified by primer set Int2F: 5'- GGTTGAACTGGATGGACGGA - 3' and Int2R: 5'- CAACAGCTGGTTTTCCCACTG - 3'. Amplification of these partial segments of SAI-1 involved an initial incubation at 96 °C for 3 min, followed by 35 cycles of 94 °C for 1 min, 60 °C for 1 min, 72 °C for 1 min, then a final 5 min extension at 72 °C. Using PCR primers and BigDye terminator chemistry on a 3130XL or 3730XL Genetic Analyzer (Applied Biosystems), PCR products of both the mitochondrial ND2 and ND3 genes and the nuclear SAI-1 gene were directly sequenced by Services in Molecular Biology GmbH (Berlin, Germany) and Mclab (California, United States) respectively. Sequences corresponding to new mitochondrial ND2 and ND3 haplotypes and nuclear SAI-1 alleles have been deposited in the EMBL Nucleotide Sequence Database under the accession numbers listed in Appendices 2.1 and 2.2.

3.2.3. Sequence Alignment and Grouping

All sequence alignments were performed using the CLUSTALW algorithm in MEGA 5.10 (Tamura et al., 2011). Alignments were then checked by eye, and corrected manually. Because of the length polymorphism in the nuclear SAI-1 marker among distinct lineages, several gaps were added to adjust length of sequences. In the *Pelophylax shqipericus* lineage, an additional 353 bp fragment within the retroelement RanaCR1 was recognized. After searching the RepeatMasker database (Institute for Systems Biology; http://www.systemsbiology.org; with the options: search engine=cross match; speed/sensitivity=slow; DNA source=vertebrate), it was determined that this fragment consists of simple repeats (GGTAC) and a member of DNA transposon family (TDR22) (Jurka and Drazkiewicz, 2002). This part of the sequence was therefore removed from all genetic and phylogenetic analysis.

For mitochondrial genes, each sequence was blasted against the GenBank database to establish to which main haplogroup it belongs. For the nuclear marker, the sequences of each individual were grouped according to genotypes as either homozygotes or heterozygotes. Two peaks in the chromatograms at corresponding positions were accepted as heterozygotes. Because there were enough homozygote individuals (nearly 500), we did not clone any nuclear sequences. Instead, nuclear allele phasing or allelic composition of a genotype was estimated on the basis of alleles derived from homozygotes (i.e. known phases) and internal primer pairs that specifically amplified one member of the allele pair within each genotypes becaused by polymorphism at the primer binding site. The resulting alignment of albumin intron alleles was subjected to a test for recombination using the Recombination Detection Program RDP 3.44 (Martin et al., 2010).

3.2.4. Phylogenetic Analysis

Calculation of haplotype and genotype frequency, haplotype diversity (h) and gene diversity (H), expected/observed heterozygosity, and nucleotide diversity (π) (Nei, 1987) were implemented in software ARLEQUIN version 3.5.1.2 (Excoffier and Lischer, 2010). Estimation of model of sequence evolution for both data types was conducted in MEGA 5.10 based on the Bayesian Information Criterion (BIC, Swarz, 1978). The best-fit molecular evolution model for the complete mitochondrial ND2+ND3 (328 sequences) was the general time-reversible model (GTR) with gamma-shaped rate variation (G = 1.046) and a fraction of invariant sites (I=39%) (ln L=-12585.68, BIC=33328.34); the best-fit model for the nuclear SAI-1 + RanaCR1 (145 sequences) was the Tamura-3-parameter (T92) model (Tamura, 1992) with a gamma-distributed shape parameter G = 0.622 (ln L=-6794.10, BIC=17040.46). Because the GTR model is not available in MEGA for genetic distance calculation, the second best-fit model, the Tamura-Nei (TrN) model (Tamura and Nei, 1993) with a gamma-shaped rate variation (G=0.325) and (In L=-12638.43, BIC=33381.97) was used to calculate genetic distance among distinct mitochondrial groups.

Bayesian phylogenetic analysis and divergence time estimation of mitochondrial and nuclear data sets were performed in the program BEAST version 1.7.5 (Drummond et al., 2012). The data sets were prepared using the BEAST assistance program BEAUTI 1.7.5. For the mtDNA data set, Bayesian analysis was initiated from the UPGMA starting tree general time-reversible mutation model (GTR) with gamma-shaped rate variation, using 4 discrete mutation classes, and a percentage of invariant sites (GTR+G+I) (Huelsenbeck and Rannala, 2004). For the nuDNA dataset, the Hasegawa-Kishino-Yano (HKY) mutation model (Hasegawa et al., 1985) with gamma distributed site rate variation, using 4 discrete mutation classes, and a percentage of invariant sites (HKY+G+I), was set to run using a random starting tree. For the nuDNA data set, a relaxed molecular clock (Drummond et al., 2006) was used, whereas for
mtDNA data set, a strict clock was estimated using an uncorrelated lognormal prior and the Yule model as a tree prior. To calculate posterior distribution parameters, the MCMC was run for 100,000,000 steps and sampled every 10,000 steps. The first recorded 1000 trees were discarded as burn-in. The effective sample sizes for posterior probability were evaluated from the logfiles in TRACER 1.5 (Rambaut and Drummond, 2009). Additionally, *P. nigromaculatus* (North Korea), *P. saharicus* (north-western Africa), *P. perezi* (Iberian Peninsula), *P. lessonae* (Central Europe), *P. bergeri* (Apennine Peninsula), *P. shqipericus* (western coast of Balkan Peninsula), and *P. cretensis* (Crete) were included as outgroup comparisons in this study (Appendices 1 and 2).

To reveal evolutionary relationships and current mutational variations and probable ancestral connections among mitochondrial haplotypes and nuclear alleles, median-joining networks (Bandelt et al., 1999) were constructed using the program NETWORK 4.6.1.1. For haplotype phylogeny 1817 ND3 haplotypes and for nuclear phylogeny 1369 SAI-1+RanaCR1 alleles were used. Outgroup sequences were excluded from the network analysis for both mitochondrial and nuclear datasets.

Bayesian tree topologies, numbers of mutational connections and branching patterns among haplotypes and alleles, using TrN+G genetic distance on the basis of joint ND2+ND3 sequences (as suggested by Akın et al., 2010b) and T92 + G genetic distances among SAI-1+RanaCR1 alleles, were accepted as parameters to define the MHGs, the main allele groups (MAGs), subgroups, and allele groups. In addition to these, unique variations such as transitions, transversions, and insertions and deletions (indels) within the SAI-1+*Rana*CR1 marker were used to define allele groups, because when programs calculate genetic distance from a dataset including indels as well as substitutions they generally underestimate genetic distance among lineages.

Divergence times for both mtDNA and nuDNA datasets were calculated by applying two distinct methods. The first was a simple linear regression model as used by Akın et al. (2010b). For mtDNA, pairwise TrN+G genetic distance was divided by two constant evolutionary rates (1.4 and 1.5% per Myr) calculated using as a calibration point two distinct divergence times (5.3 and 5.5 Ma) between Cyprian and Anatolian water frog populations. For nuDNA, pairwise T92 + G genetic distance was divided by two evolutionary rates (0.69 and 0.72% per Myr) calculated using the same two putative divergence times (5.3 and 5.5 Ma) between Cyprian and Anatolian water frog populations. Second, a Bayesian method was applied, using the software BEAST (Drummond and Rambout, 2012).

3.3. Results

3.3.1. Molecular Diversity

In total, 568 ND2 and 1821 ND3 sequences from Turkey, the Middle East, Central Asia, the Levant, Cyprus, Central Europe and Eastern Europe were used in the analysis (Appendix 3.1). The ND2 sequences were represented by 242 distinct haplotypes with h=0.98±0.002 and π =2.76±1.34%, while the ND3 sequences defined 182 distinct haplotypes with h=0.93±0.004 and π =3.03±1.53%. The joint sequences (ND2+ND3; N=328) including outgroups contained 663 variable sites (48.1%) of which 606 (44%) were parsimony-informative. The nuclear marker analysis for SAI-1+*Rana*CR1 yielded sequences for 1369 individuals from the same spread of regions (Appendix 3.2). They were represented by 295 distinct genotypes of which 863 were heterozygous and 506 were homozygous with expected heterozygosity=0.9314 and observed heterozygosity=0.6304. These genotypes were formed by 114 unique alleles with gene diversity H=0.92±0.003 and π =2.05±1.00%. The unique alleles and outgroup sequences (N=145) contained 268 variable sites (23.1%) of which 190 (16.4%)

were parsimony-informative. No evidence for recombination of nuclear serum albumin alleles was found.

3.3.2. Phylogenetic Relationships, Genetic Divergence, and Geographical Pattern of Mitochondrial Haplogroups

The Bayesian tree topology, from joint analysis of ND2+ND3 genes (Fig. 3.2), the median-joining network (Fig. 3.3), the geographical pattern of ND3 haplotypes (Fig 3.1) and pairwise TrN+G among ND2+ND3 sequences (Appendix 3.3) indicate eight well differentiated MHGs as shown by Akın et al., (2010b). The mean TrN+G distances (G=0.325) between these eight MHGs varies from 0.032 (MHG6-MHG7) to 0.102 (MHG1-MHG8) while mean distance values within each MHG was ≤ 0.014 .

MHG1 includes specific haplotypes of European *P. ridibundus*, ranging from France (throughout Europe) to Central Russia, and haplotypes of Balkan lake frogs, *P. kurtmuelleri* (Akın et al., 2010b). MHG2 consists of characteristic haplotypes of *P. bedriagae*, found in Jordan, western Syria and the Nile delta of Egypt. Even though this group has been sampled only to a limited extent, it reveals a high diversity (π =1.0±0.59%), and has been divided into two subgroups. MHG2a was found only in Jordan and As Suwayda in southern Syria whereas MHG2b was recorded in Egypt and elsewhere in Syria. The actual geographic patterns of these subgroups are not known, however, beacuse of limited sampling. MHG3 represents characteristic haplotypes of the recently described species *P. cypriensis* (Plötner et al., 2012) in Cyprus. Sister groups MHG4 (Cilician West) and MHG5 (Cilician East) were found mainly in the Cilician plain sympatrically, but the frequency of MHG5 is higher in the eastern part of the Amanos mountains (Akın et al., 2010b; Fig 3.1. and Appendix 3.1).



Figure 3.1. Distribution of main mitochondrial haplogroups (MHG1-6) and subgroups (MHG6ad) in the eastern Mediterranean region water frog complex (genus *Pelophylax*).

MHG6 (cf. *bedriagae*, Anatolia) is the most diverse group (haplotype diversity, $h=0.89\pm0.01$ and nucleotide diversity, $\pi=1.6\pm0.09$). It consists of four subgroups (a-d), between which the mean genetic distance ranges from 0.017 to 0.027. MHG6a (cf. *caralitanus*) was mainly found in south-western Turkey (the Anatolian Lake District and Konya plain). It coexists with haplotypes of MHG4 and 5 in the eastern border of Konya plain. MHG6b (cf. *cerigensis*) was locally distributed in the region between Antalya and Muğla on the Mediterranean coast of south-western Turkey, also in Rhodos and Karpathos islands. MHG6c (cf. *bedriagae*) is the most widely distributed subgroup, ranging from western

Anatolia, including the Aegean islands of Samos, Lesvos and Chios, to central Russia and to the northern shore of the Caspian Sea (type locality of *P. ridibundus*). MHG6d (Euphrates) was found in the catchments of Euphrates and Tigris rivers (Akın et al., 2010b) in eastern Anatolia. It overlaps with haplotypes of MHG6c in the valleys of the Aras and Kura rivers in northeastern Anatolia and western Iran (Fig.3.1).

MHG7 (Central Asia 1) corresponds to characteristic haplotypes of *P. terentievi*, distributed to the south and east of the Caspian Sea in Iran, Turkmenistan and Uzbekistan. MHG8 (Central Asia 2) represents characteristic haplotypes of the undescribed species, *P. sp. novum*, found in Kyrgyzstan, Kazakhstan and western Tajikistan. It coexists with haplotypes of MHG7 on the river Zeravshan, which is north-east of the river Amu Darya (unpublished data).



Figure 3.2. Maximum clade credibility tree calculated on the basis of Bayesian phylogenetic analysis of the joint ND2+ND3 mtDNA haplotypes (1378 bp) from Western Palearctic and eastern species of eastern Palearctic water frogs (genus *Pelophylax*). Values at nodes are posterior probabilities. The branch labels are haplotype IDs of ND2 and ND3 combinations given in Appendix 3.1.



Figure 3.3. Median joining network of mitochondrial ND3 haplotypes constructed by NETWORK 4.6.1.1 (Bandelt et al., 1999) showing genealogical relationships among main haplogroups (MHG1-8) and subgroups (MHG6a-6d and MHG2a-2b). The sizes of the circles represent haplotype frequency; median vectors (mv) indicate ancestral haplotypes that were not observed in the data set. Numbers on lines show mutational changes from one haplotype to next one in the ND3 alignment.

3.3.3. Phylogenetic Relationships, Genetic Divergence, Geographical Pattern of Nuclear Allele Groups and Their Relationships with Mitochondrial Haplogroups

The Bayesian tree topologies (Fig. 3.4), the median-joining network of the nuclear SAI-1+RanaCR1 alleles (Fig. 3.6), the geographic pattern of nuclear alleles (Fig. 3.5a,b,c), the pairwise T92+G genetic distance (Appendix 3.4) and unique molecular variations among albumin alleles, particularly for subdivision of subgroups and allele groups (Appendix 3.5), reveal five distinct MAGs: MAG1 (Anatolia), MAG2 (Asia), MAG3 (Ridibundus), MAG4 (Levant; BED2) and MAG5 (Cyprus). The mean T92+G distance (G=0.622) between these MAGs ranged from 0.013 (MAG2-MAG3) to 0.050 (MAG1-MAG5); the mean distance values among MAG1, MAG2 and MAG3s were ≤ 0.010 .

The Anatolia MAG1 is the most widespread and well differentiated monophyletic group within the eastern Mediterranean water frog complex, with five allele subgroups (a-e; Fig. 3.4), some widely distributed, others geographically restricted. The mean distance values between the five subgroups are 0.005-0.018. Each of these subgroups in turn consists of one or more allele groups. MAG1 has variations in G repeats (3-5) at positions between 514 and 518 within the *Rana*CR1 element and in T repeats at positions 1135-1140 at the end of the intron.



Figure 3.4. Maximum clade credibility tree constructed using Bayesian phylogenetic analysis of nuclear SAI-1 + RanaCR1 alleles from western Palearctic and eastern taxa of Eastern Palearctic water frogs (genus *Pelophylax*). Values at nodes indicates posterior probabilities. The branch labels are names of alleles given in Appendix 3.2.

MAG1a has three allele groups: ANT1, ANT2 and ANT3. Alleles of ANT1 (ant1-1 - ant1-6) were found only in the closed Konya basin (most frequently, including the type locality of *P. cf caralitanus*), in Kayseri-Sultansazlığı and Akşehir-Eber lakes (Isparta province), and in Işıklı lake (Kütahya and Uşak provinces). Except for Kütahya and Uşak, haplotypes characteristic of MHG6a (cf. *caraliranus*) were also found at all of these localities. The ANT2 allele (ant2-1) was most frequently recorded in the Anatolian Lake District region except in a few localities (Afyon and İzmir Çandarlı) where individuals had mostly cf. *caralitanus* specific haplotypes (MHG6a). In contrast, alleles of ANT3 (ant3-1 - ant3-6) were distributed in most parts of Turkey except western Anatolia. Its distribution, therefore, covers not only the cf. *caralitanus* subgroup (MHG6a) but also other mitochondrial haplogroups (Fig. 3.1).

MAG1b has two geographically restricted allele groups: ANT4 and ANT5. Alleles of ANT4 (ant4-1 - ant4-4) were frequently found in the Mediterranean coastal region of SW Anatolia between Antalya and Muğla where haplotypes specific for cf. *cerigensis* (MHG6b) occurred. They were also found in the Lake District region where cf. *caralitanus* specific haplotypes (MHG6a) were found, and less frequently in the catchment of the Büyük Menderes River where haplotypes specific of cf. *bedriagae* (MHG6c) were present. In contrast, alleles of ANT5 (ant5-1 - ant5-7) were predominantly recorded farther east along the Mediterranean coast of Anatolia between Antalya and Mersin where haplotypes specific for cf. *caralitanus* (MHG6a) occurred. The MAG1c (cer1-1) allele was only found on the islands of Rhodos and Karpathos, where only *cf. cerigensis* specific haplotypes (MHG6b) have been reported.



Figure 3.5a. Distribution of allele groups of the Anatolia main allele group (MAG1a-e), Asia (MAG2: CIL1, TER, CIL3, BED1), Cyprus (MAG5) and Levant (MAG4: BED2). Distribution of RID (MAG3) is shown on this map without giving group divisions.

MAG1d has three allele groups: ANT6, ANT7 and ANT8. Alleles of ANT6 (ant6-1 - ant6-2) were found between north-eastern Anatolia, Ukraine and Kazakhstan, including throughout the Caucasus. Alleles of ANT7 (ant7-1 - ant7-3) had the widest distribution, including western and central Anatolia, western and central parts of the Black Sea region (but not the eastern Black Sea region), and Thracia, Ukraine and Kazakhstan. The absence of this allele group from the eastern Black Sea region indicates that it probably spread over Thracia and the western Black Sea region from Anatolia to eastern Europe. ANT8 includes several rare alleles

(ant8-1, 8-4 and 8-5), found in a few localities, as well as common alleles ant8-3 (in western Anatolia) and ant8-2 (several localities in Ukraine, Kazakhstan but only one locality in Turkey, Akşehir-Eber lake), which suggests that they occurred frequently in Anatolia in the past, but later they were lost or became rare. The distribution of this MAGd shows a similar pattern to haplotype MHG6c (cf. *bedriagae*; Fig. 3.1).

MAG1e has four allele groups, ANT9, ANT10, ANT11 and ANT12. Alleles of ANT9 (ant9-1 - ant9-3) were only found in south-western Anatolia. Alleles of ANT10 (ant10-1 - ant10-5) were distributed in south-western, western, and central Anatolia, the western Black Sea region and Thracia. Alleles of ANT11 (ant11-1 - ant11-10) were found throughout Turkey except for southern Mediterranean coastal localities.

The Asia MAG2 is probably the most interesting group, consisting of geographically overlapping or adjacent western Asian allele groups (CIL1, CIL3, and BED1) and adjacent Central Asian allele groups (TER and SP NOV); the mean distance values between these being 0.004-0.011. Within MAG2, there are variations in the number of CA microsatellite repeats (4-6) at the 3' UTR region of the *Rana*CR1 retrotransposon, and all share a 25 bp deletion at position 1051-1075 (Plötner et al., 2012).

Alleles of CIL1 (cil1-1 - cil1-8) occur in the Cilician and Narlı plains on both the western and eastern sides of the Amanos Mountains. At the boundaries of these plains, several heterozygous individuals, carrying alleles of the CIL-1 subgroup and either MAG1 or MAG3 groups were found.

Alleles of CIL3 (cil3-1 - cil3-3) were interestingly found in the eastern Amanos mountains in a few localities (Kırıkhan in Hatay province and Lake Kumaşır in Kahramanmaraş province) and in Jordan (Al Kerak and near Jesus where individuals were heterozygous with an allele of BED1). Alleles of BED1 (bed1-3,

bed1-3 - bed1-5) were also recorded in Jordan, Syria and one locality in Turkey (Lake Kumaşır, where one individual was heterozygous carrying alleles of CIL1 and BED1 groups). There may, however, well be individuals carrying alleles belonging to CIL3 and BED1 in the region between Hatay in Turkey and Jordan particularly throughout the catchment of the River Orontes. No field studies have so far been carried out in this region; sampling there (which will only be possible when the political situation in Syria and Lebanon improves) could make clear both the phylogenetic positions of these groups and their relationships to the CIL1 group, which are now only supported by a low posterior probability value, 0.3 (Fig. 3.5). In respect of mtDNA, central southern Turkey includes the Cilician West (MHG4) haplotype (west of the Amanos Mountains) and Cilician East (MHG5) haplotype (east of the Amanos Mountains), with haplotypes of *P. bedriagae* (MHG2) distributed in Syria and Jordan (Fig. 3.1). Just as for the nuDNA data, the haplotype pattern is not known in the catchment of the River Orontes between Hatay and Syria.

The TER group (alleles cil2-1, cil2-2, ter1-ter6), one of the Central Asian groups of MAG2, was mostly found to the south and east of the Caspian Sea, in Iran, Turkmenistan and Uzbekistan. The distribution of these alleles overlaps with that of haplotypes specific for *P. terentievi* (unpublished data). It is difficult to explain why two alleles, cil2-1 and cil2-2, were only found in central-southern Turkey, forming a clade within the TER group, not the CIL1 group. Individuals carrying these two alleles, have haplotypes from distinct groups (MHG4 and 5 and MHG6c and d), depending on the locality. Alleles of the other Central Asian group, SP NOV, were found in Kyrgyzstan, Kazakhstan and western Tajikistan, coincident with haplotypes specific for *P. sp nov*. (unpublished data).



Figure 3.5b. Distribution of allele groups in central-southern Turkey (MAG1a, MAG1b, MAG1d, MAG1e, RID MAG3, Asia MAG4 (CIL1, CIL2=TER, CIL3, BED1) and the Levant (Levant MAG-BED2, BED1 and CIL3).

The Ridibundus MAG is widely distributed. It shows variations in CA microsatellite repeats (5-10) at the 3' UTR region of the *Rana*CR1 retrotransposon within SAI-1 and T repeats (5-6) at the end of the intron (positions 1135-1140). It consists of four allele groups (RID1-RID3 and KUR); mean distance values between these are 0.005-0.013. Alleles of RID1 are mainly distributed in the European part of Turkey, the western Black Sea region, Kazakhstan (including the allele rid1-5, found at the type locality of *R. ridibundus*) and Ukraine (Fig 3.4b), where some individuals were heterozygous for one allele from the RID1 group, the other allele from the Anatolia MAG1d or the MAG1e subgroups. This sympatric distribution of RID1 and ANT allele groups is also supported by the overlapping distribution of mtDNA haplotypes belonging to MHGc (cf. *bedriagae*) and MHG1 (*P. ridibundus*) in these regions, except in the western Black Sea where haplotypes of cf. *bedriagae* type were only recorded but no *ridibundus* specific haplotypes.

RID2 includes Central European specific *ridibundus* alleles (rid2-3 and 2-4). RID2 also includes one of the most common *ridibundus* alleles (rid2-1) in Central Anatolia and the Black Sea region, as well as rid2-6 and 2-7 (unique to the European part of Turkey) and rid2-5 (a common allele in north-eastern Anatolia). Relationships between these last three alleles and the others are, however, not clear. Alleles of RID2 in Turkey were carried as homozygotes or heterozygotes in individuals with alleles from other ANT or RID groups. Except in Central Europe and the European part of Turkey, the haplotype pattern of Central Anatolia, the Black Sea Region and north-eastern Anatolia (where the haplotype specific for *cf. bedriagae* was present), was not concordant with the distribution of nuclear alleles of RID2 group, since no *ridibundus* haplotypes were found in these regions.

RID3, the most frequent rid allele group in Turkey, includes alleles specifically distributed in the eastern Anatolia (rid3-7, 3-8 and 3-9), the rid3-5 allele found in some localities near the Tuz Gölü salt lake in central Anatolia, in the European part of Turkey and in south-eastern Anatolia, and the rid3-6 allele recorded only at a few localities near the Tuz Gölü. Individuals were either homozygote, especially in eastern and central Anatolia, or heterozygote with alleles from the ANT, CIL and RID groups. The RID3 group probably reveals the most discordant haplotype pattern; the region where alleles of this group are found is dominated by haplotypes of the cf. *caralitanus*, cf. *bedriagae* or Euphrates haplogroups (Akın et al., 2010b). Except in the European part of Turkey, none of haplotypes of the *ridibundus* MHG were found in these regions.



Figure 3.5c. Distribution of allele groups of RID MAG3 (RID1, RID2, RID3 and KUR). Distributions of other MAGs, including Anatolia, Asia, Cyprus and Levant are shown only without group divisions.

Group KUR contains alleles characteristic of Balkan lake frogs, *P. kurtmuelleri*. It is well resolved from the other RID allele groups by the high posterior probability value of 0.93, the number of mutational connections in the median-joining network and several unique mutations within the SAI-1+RanaCR1 marker, including a 124 bp deletion starting after the CA repeats at positions 758-890 (Plötner et al., 2009). The distribution of alleles of this group is concordant with the haplotype pattern specific for *P. kurtmuelleri*. Its complete distribution and its relationships with other European water frogs are not known.

The Levant MAG4 (BED2) is the other group that carries *P. bedriagae* specific haplotypes. In contrast to the BED1 group, which is closely related to Cilician and other Central Asian groups, BED2 forms a clade among outgroup species. It seems to have sister relationships with *P. shqipericus*, *P. lessonae and P. bergeri*, although, this conclusion is supported by a low posterior probability value (Fig. 3.5). Alleles of this group share with the Asia MAG2 a 25 bp deletion at positions 1051-1075, but also has a 7 bp deletion at positions 758-764 and several unique transitions and transversions. Alleles of this group were found in Jordan, but its actual distribution is not known because of sampling deficiency. Finally, the Cyprus MAG5 has alleles characteristic of the recently described Cyprus water frog species, *P. cypriensis* (Plötner et al., 2012) and it carries haplotypes specific of this species.



Figure 3.6. Median joining network of nuclear SAI-1+RanaCR1 alleles constructed by NETWORK 4.6.1.1 (Bandelt et al., 1999) demonstrating phylogenetic relationships among main allele groups (MAG1 and subgroups MAG1a-e, Anatolia; MAG3, Ridibundus; and MAG2, Asia). The sizes of the circles indicate allele frequency; median vectors (mv) represent ancestral alleles that were not found in the dataset. Numbers on lines show mutational changes (substitutions and indels) from one allele to next in the SAI-1+RanaCR1 alignment.

3.3.4. Comparison of mtDNA and nuDNA Tree Topologies and Estimation of Times of Divergence

Our Bayesian phylogenetic results reveal clear discrepancies between the mitochondrial and nuclear gene trees (Fig. 3.7). The topology obtained from mitochondrial genes consists of several reciprocally monophyletic groups, supported by high posterior probabilities. In contrast, the topology based on the nuclear marker contains both weakly supported groups, some of which showed paraphyletic relationships; and some well supported clades. Thus, for example, in the mtDNA tree, P. bedriagae (MHG2) represents an ingroup species, clustering with P. ridibundus (MHG1) as a sister group, whereas in the nuDNA tree it (Levant MAG5) forms a clade within the outgroup species P. shqipericus/P. lessonae/ P. bergeri. P. epeiroticus split off before P. cretensis/P. cypriensis in the mtDNA tree, but in the nuDNA tree seems more closely related with the Anatolia and Asia/Rid MAGs. P. cretensis diverges earlier than P. cypriensis in the mtDNA tree, but these two form sister species in the nuDNA tree (Plötner et al., 2012). P. ridibundus in the mtDNA tree forms a sister group with P. bedriagae from the Levant whereas in the nuDNA tree it clusters with the Asia MAG2, including groups from Central Asia, central-southern Turkey and the Levant (BED1). In contrast to the well characterized Cilician (MHG4 and 5) and Central Asia (MHG7 and 8) monophyletic groups in the mtDNA tree, their relationships within the Asian MAG2 are not fully resolved; some clades within this group were indeed only weakly supported.



Figure 3.7. Bayesian phylogenetic tree topologies of mtDNA (ND2 + ND3 genes) on the left and nuDNA (SAI-1 + RanaCR1) on the right. Tree topologies include only main groups (MHG or MAG) and subgroups without showing haplotype and allele names.

The two methods for estimating divergence times, a simple linear regression model and analysis using BEAST, provided similar results within a data type (Table 3.1). As already noted, for both mtDNA and nuDNA, divergence times between the Cyprian and Anatolian water frog populations (5.3 and 5.5 Ma) were used as a calibration points, based on the idea that these populations became separated around the end of the MSC when the Mediterranean Sea reverted to being a marine basin (Akın et al., 2010b). Using these calibration points, however, there are discrepancies between mtDNA and nuDNA for some groups or species. Thus, for example, the estimated rate for mtDNA evolution indicates that P. cretensis diverged before 9 Ma, whereas the estimated rate for nuDNA evolution indicates that its divergence occurred around the same time as that of P. *cypriensis*, about 6-5 Ma. mtDNA indicates a deep divergence (about 3.5-4.5 Ma) between Cilician groups (MHG4 and 5) and Central Asian groups (MHG7 and 8) whereas nuDNA results indicate a very shallow divergence among these groups (CIL1 and 3-TER and SP NOV), diversifying about 1.5-0.5 Ma. Similarly, in contrast to the mtDNA results, divergence between the RID MAG3 and the Asian MAG2 seems very recent in the nuDNA results (~2.0-1.7 Ma).

Table 3.1. Estimates of divergence times for mtDNA and nuDNA markers. Divergence times calculated between mitochondrial groups (MHGs) and subgroups on the basis of TrN+G distances and between nuclear groups (MAGs) and subgroups on the basis of T92+G distances. An evolutionary rate for mtDNA was estimated by dividing genetic distance between MHG3 and MHGs 4, 5, and 6 (0.077) by the two divergence times (5.3 and 5.5 Ma), giving rates of 1.45% Myr⁻¹ and 1.4% Myr⁻¹ respectively. For nuDNA, the T92+G distance between the Cyprus MAG5 and subgroups CIL1, CIL3, and BED1 of Anatolian MAG1 (0.038) was divided by the same two time estimates, yielding rates of 0.72% Myr⁻¹ and 0.69 Myr⁻¹ respectively. Using BEAST, divergence times were estimated using the calibration point *P. cypriensis* - sister clade (Anatolian populations) at 5.3 ± 1.0 Ma. The tree prior was a birth-death process.

		Divergence Time Estimates (Ma)		
			BEAST	
Group Comparisons	TrN + G	Regression Model	Mean	95% credibility
MHG1/2-MHG3-8	0.082	5.47		
MHG1-MHG2	0.079	5.27	5.1	2.38-6.81
MHG3-MHG4/5/6	0.077	5.3***		
MHG8-MHG6/7	0.052	3.47		
MHG4/5-MHG6/7/8	0.047	3.13		
MHG4-MHG5	0.041	2.73		
MHG6-MHG7	0.032	2.13	4.27	2.12-5.49
MHG6a-MHG6b/c/d	0.021	1.40	3.87	1.9-4.96
MHG6d-MHG6b/c	0.018	1.20	3.42	1.6-4.31
MHG6b-MHG6c	0.017	1.13	3.12	1.36-3.88
P. cretensis-MHG1-8	0.176	11.73	7.52	4.02-10.42
Anatolia MAG - Ridibundus/Asia MAGs	0.040	5.56		
Cyprus - Anatolia MAG/CIL1/CIL3/BED1	0.038	5.3***	4.76	2.85-6.59
MAGa-MAGb/c/d/e	0.015	2.08	3.3	1.57-5.21
MAGe-MAGb/c/d	0.010	1.39	2.2	1.1-3.56
MAGb-MAGc/d	0.007	0.97	1.89	
MAGc-MAGd	0.005	0.69	1.66	
Ridibundus MAG-Asia MAG	0.013	1.81	3.4	1.69-5.48
RID1-KUR/RID2/RID3	0.008	1.11	2.24	1.04-3.74
KUR-RID2/RID3	0.010	1.39	1.55	0.72-2.67
RID2-RID3	0.005	0.69	0.92	0.32-1.69
BED1-SP NOV/CIL3/TER/CIL1	0.008	1.11	1.68	
SP NOV-TER/CIL1	0.010	1.39	1.28	
TER-CIL1	0.005	0.69	0.96	0.42-1.67
BED2- Anatolia/Asia/Ridibundus	0.048	6.67	7.94	4.07-12.36
P.cretensis- Anatolia/Ridibundus/Asia MAG	0.035	4.86		

3.4. Discussion

Our analysis of both mtDNA and nuDNA markers in the eastern Mediterranean water frog complex has uncovered several striking discordances. The evolutionary rate estimated for two genomes is different, with the mitochondrial rate ($\sim 0.7\%$ Myr⁻¹) nearly twice the nuclear rate (1.4% Myr⁻¹). Six major discordant groups and their probable causes are discussed below.

3.4.1. Potential Mechanisms Causing Phylogeographic Discordance among mtDNA and nuDNA markers

3.4.1.1. P. cypriensis versus P. cretensis

3.4.1.1.1. Isolation of Cyprus

We have used as a calibration point the isolation of Cyprus from Anatolia at ~5.5-5.3 Ma based on the palaeogeography of the region at the time of the MSC (Plötner et al., 2010; Akın et al., 2010b). Phylogenetic results for mtDNA, constrained such that Cyprian water frogs became isolated from the mainland after the MSC, indicated that populations on Crete diverged ~10-9 Ma, before the MSC (Plötner et al., 2010; Akın et al., 2010b). In contrast, our new results for the nuDNA marker (Fig. 3.7) indicate that both the Cyprian and Cretan populations diverged from the mainland at the end of the MSC and are a sister species.

Plötner et al. (2010) and Akın et al. (2010b) envisaged that during the latter stages of the MSC, when the Mediterranean basin had partially re-flooded, a land bridge linked Cyprus to Anatolia, allowing exchange of frog populations. This land bridge became flooded and the frog populations became isolated when the Mediterranean Sea rose to its present level at the end of the MSC and the start of the Pliocene. A more recent synthesis of the palaeogeography and palaeoenvironments in the Mediterranean region during the MSC (Bache et al., 2012) agrees that the MSC began about ~6 Ma when uplift of Spain and Morocco restricted the connections between the Mediterranean Sea and the Atlantic Ocean (cf Krijgsman et al., 1999a;b), resulting in the desiccation of the Mediterranean basin; and also agrees that the ending of the MSC involved partial re-flooding of the basin, at which point the sea surface was ~650 m below its present level. Subsequent re-flooding to the present sea-level occurred at an orbitally-tuned age of 5.46 Ma. This is the best current estimate for the timing of the end of the MSC. At present, Cyprus and Anatolia are separated by a seaway that in places is ~ 750 m deep. The present sea floor does not mark the sea floor at the end of the MSC, however, because it is underlain by Pliocene and Pleistocene sediment; between Cyprus and NW Syria, up to 2500 m of post-Messinian sediments have been deposited (Hall et al., 2005). Loading by this sediment can be expected to have depressed the adjoining crust, such that prior to the end of the MSC the land surface may well have been above the contemporaneous (low) sea level. The seafloor in this region has experienced significant crustal deformation since the Messinian (Hall et al. 2005; Sevrek et al., 2014); it forms part of a major active fault zone, the Misis-Kyrenia Fault Zone. The palaeogeography at the time of partial re-flooding near the end of the MSC, depicted by Bache et al. (2012: Fig. 16), shows a ~20 km wide palaeo-strait separating Cyprus and Anatolia. This illustration has, however, not been corrected for the effects of post-Messinian sediment loading and other crustal deformation. We therefore continue to regard as reasonable the assumption that the frog populations in Cyprus and Anatolia became isolated at the start of the Pliocene; hence its continued use in the present study. No land bridge between Cyprus and Anatolia is plausible since then (cf. Hadjisterkotis et al. 2000).

3.4.1.1.2 Isolation of Crete

The palaeogeographic history of Crete has been rather different, being related primarily to the development of the Hellenic subduction zone, along which the leading edge of the African plate has plunged beneath Eurasia during the Cenozoic. The dynamics of the subduction process have affected the stress field, causing extension within the overriding plate, which has resulted in significant change in its shape and, consequently, significant relative movements between Crete, the Peloponnese to the west, and western Anatolia to the east. The palaeoenvironmental changes associated with the MSC have been superimposed onto the effects of these processes. It should be noted, however, that the palaeogeographic maps published by Bache et al. (2012) illustrating conditions during the MSC do not take account of subsequent changes in the shape of the region (i.e., are non-palinspastic) and should be viewed in the context of dynamic reconstructions. Recent syntheses of the evolution of this subduction-related system include works by Ring et al. (2010) and Royden and Papanikolaou (2011). These reconstructions are, however, subject to some uncertainty, because the magnitude of the subduction-related extension has been inferred largely from cooling histories, but these histories will in turn be affected by changes to the geometry of the subduction (Westaway, 2006).

Royden and Papanikolaou (2011) estimated that since the subduction began at ~40 Ma this part of the African plate has moved northward relative to Eurasia by ~1000 km, but the Eurasian crust has extended by ~100 km north of the Peloponnesos and by ~250 km north of Crete; Crete has thus moved ~150 km southward relative to the Peloponnesos. Furthermore, the southern margin of Eurasia was initially almost straight and oriented west-east, but has evolved (as an example of 'oroclinal bending') to its present strongly curved shape as a result of forces associated with the subduction. Crete thus has also experienced components of east-west separation relative to both the Peloponnesos and ~230 km from SW Anatolia but reconstructions by Royden and Papanikolaou (2011) show that these distances were smaller at earlier periods (~80 km and ~150 km at 4 Ma and <20 km and ~80 km at 15 Ma).

During the Middle Miocene the entire modern Aegean region is thought to have been subaerial (e.g., Rögl, 1999), so no isolation of frog populations would be expected. During the Tortonian stage of the Late Miocene (~11-7 Ma), a marine transgression entered the Aegean region from the south (e.g., Rögl, 1999), forming an elongated north-south marine embayment that may have been no more than ~100 km wide (e.g., Popov et al., 2006). This was evidently a time of complex changes to marine and terrestrial environments, as discussed for example by Köhler et al. (2010), Krijgsman et al. (2010) and Vasiliev et al. (2011); it is unclear whether the marine connection at this time between the Mediterranean and the Aegean was located to the west or to the east of Crete.

During the peak of the MSC (~5.6 Ma) lake basins developed in parts of the central Aegean Sea (e.g., Anastasakis et al., 2006). These received sediment input from adjoining areas of land (e.g., Hejl et al., 2008). There is no evidence, however, that these lacustrine environments reached as far south as Crete. At this time, what is now Crete can be envisaged as an upland rising kilometres above the playas that occupied the desiccated Mediterranean and southern Aegean basins. The heights of the lowest cols separating Crete from Anatolia or the Peloponnesos are unclear, however, making it difficult to form any conclusion from the geological evidence as to whether the frog populations on Crete were isolated or not. The partial marine flooding that occurred late in the MSC is, however, envisaged (Bache et al., 2012) as isolating Crete from any adjoining land areas. Nonetheless, much of present area of the central and northern Aegean Sea remained land for much of the Pliocene (e.g., Anastasakis et al., 2006; Hejl et al., 2008); the progressive subsidence that has widened this sea to its present dimensions has presumably involved outward flow of mobile lower-crustal material and is coupled to the uplift of the adjoining land areas (cf. Westaway et al., 2009).

The complexity of the above-mentioned geological history makes clear that it is unwise to use any inferred timing of the 'separation' of Crete to calibrate our records; basing the calibration on Cyprus is evidently preferable.

3.4.1.1.3 Possible Explanations for the Discrepancy

In principle, the inconsistency between our mtDNA and nuDNA markers for the divergence time of *P. cretensis* could be explained in two ways. First, the divergence time from the nuDNA is correct, in which case the freshwater environments that existed in the central Aegean Sea during the MSC somehow enabled linkage between the Cretan and Anatolian frog populations. It would, however, be difficult to test this hypothesis. Second, the divergence time from the mtDNA is correct, in which case Crete became isolated from Anatolia at ~9 Ma, as a result of the partial marine inundation of the Aegean basin early in the Late Miocene. This seems more plausible since distinct selective constrains on both mitochondrial and nuclear genes and lineages could result in distinct evolutionary rates. An important precaution in the analysing biogeographic data is that a lineage divergence and the formation of a geographic barrier may not be consistent with each other. It is also possible that large differences among mtDNA genes and rate heterogeneties between lineages and within a lineages over time are too large for molecular clock calibrations (Heads, 2005).

3.4.1.2. The Levant (*P. bedriagae*)

Another striking discordance between results for mtDNA and nuDNA arises for the Levant group of frog populations, including *P. bedriagae*. mtDNA markers indicate an ingroup relationship of this group with other eastern Mediterranean water frog groups, which form a sister group with *P. ridibundus* from Europe. On the other hand, the nuDNA marker places it as an outgroup to all other members of the *P. ridibundus* group on the phylogenetic tree, indicating a time of divergence rather older than MSC. Moreover, the geographic distribution of the two markers (Figs. 3.4 and 3.7) reveals a complex pattern that is difficult to interpret. According to mtDNA, the Levant, including the Nile delta, the Jordan valley and upper reaches of the River Orontes, are occupied exclusively by *P. bedriagae* specific haplotypes. In contrast, the nuDNA reveals the existence of two divergent groups, Levant (BED2) and Asia (CIL3 and BED1), in the Levant region (Fig 3.5).

The present configuration of the Levant began to develop circa 15 Ma, when uplift of the northern Arabian Platform closed the former east-west oriented Tethys Seaway, forming a narrow land bridge between Africa and Eurasia in the vicinity of Gaziantep and Aleppo (e.g., Steininger and Rögl, 1984; Rögl, 1999). This region has since become transected by the Dead Sea Fault Zone (DSFZ), the boundary between the African and Arabian plates, which has had major effects on both topography and drainage (e.g., Westaway, 2004; Seyrek et al., 2014). The southern DSFZ, which has existed since the Middle Miocene (e.g., Garfunkel, 1981), is drained by the southward-flowing River Jordan, which has headwaters in southern Lebanon. The northern DSFZ, which may have developed as recently as the latter part of the Late Miocene (~6 Ma; e.g., Seyrek et al., 2014), is drained primarily by the River Orontes, which has its headwaters in northern Lebanon and flows northward across NW Syria to the Mediterranean coast in central-southern Turkey. The headwaters of the Jordan and Orontes rivers are ~80 km apart, the region in between being drained by the River Litani, which flows southward along the DSFZ then westward to the Mediterranean coast in southern Lebanon. Various hypotheses have been proposed for significant drainage changes in this region; for example, Horowitz (2001) proposed that the Orontes headwaters formerly drained southward into the Jordan, perhaps as recently as the Early Pliocene, whereas Butler et al. (1998) envisaged a succession of diversions of the Litani as active faulting has progressed. In contrast, Bridgland et al. (2012) envisaged that the Orontes catchment has remained constant since ~6-5 Ma, with no evidence of capture of the former headwaters of the Jordan. On the other hand, Bridgland et al. (2012) noted two major gorges along the Orontes, upstream and downstream of the Ghab Basin in NW Syria. Either or both might act to isolate frog populations, as has evidently occurred as a result of the development of the Berke Gorge on the River Ceyhan through the Amanos Mountains in central-southern Turkey, isolating the MHG4 and MHG5 frog populations (Fig. 3.4). At this stage, however, the lack of sampling in Syria and Lebanon makes it impossible to say what effect these gorges and/or putative drainage diversions might have had on isolation of frog populations.

Our mtDNA results indicate that the separation of P. bedriagae occurred during the MSC (~6-5 Ma), but nuDNA indicates that it was earlier. This discordance between mtDNA and nuDNA markers could be explained by recent introgression, male-biased dispersal during Pleistocene, or more recently at ~70 ka which was the last wet phase, by a large salty lake that covered the central Jordan Valley (Horowitz, 2001). Thus, although no samples are available from the Nile delta, Israel, Lebanon or the Upper and Middle reaches of the River Orontes, the pattern suggests male mediated gene flow, facilitating connectivity between these two lineages. Male-biased dispersal was shown in the closely related species Pelophylax ridibundus (Holenweg Peter, 2001). The distribution of mtDNA haplogroups reveals that females seem to show a strong philopatry. It is hypothesized that males from the Asia group (CIL3, BED1) may have dispersed into the Jordan valley from the Orontes catchment and mated with females of the BED2 group, acquiring their mtDNA. Another possible explanation is that the dispersal has not been sex-biased, but results from coalescence time differences between the different markers caused by effective population size differences between them; mtDNA rapidly coalescences and sorts among lineages before historical signs of introgression disappear in nuDNA.

3.4.1.3. Cilician versus Central Asia Group

The relationships between Cilician and Central Asia groups are probably the most complicated. Although both markers support close relationships among these groups, the mtDNA topology indicates clear monophyletic groups (MHG4/5 and MHG7/8), whereas topological relationships within the Asia MAG are not resolved by nuDNA. In particular, two alleles found exclusively in the Cilician and Narlı plains form a clade within the TER group, one of the Central Asia groups. It is not clear how such geographically distant groups that have developed in allopatry with no geographical connection, share these alleles. A second issue relating to these groups concerns how their genetic distance could have remained so shallow in nuDNA (Asia MAG), whereas the genetic distance from the mtDNA (MHG4/5 and MHG7/8) increases with geographic distance, indicating the expected deep divergence. Possibly these aspects result from incomplete lineage sorting and retention of ancestral alleles in the populations and pattern of inheritance differences between nuDNA and mtDNA markers (Funk and Omland, 2003; Ballard and Whitlock, 2004; Maddison and Knowles, 2006). Thus, these two groups were probably connected in the distant past, but were later completely isolated from each other. Some alleles, such as TER specific alleles, may have remained in the Cilician population as a relict. In contrast, lineage sorting of nuclear genes takes longer than for mtDNA because of the large effective population size and recombination. When nuDNA still harbours imprints of past polymorphism, mtDNA quickly evolves and differentiates into distinct lineages, erasing the ancestral history especially in the case of strong female philopatry.

3.4.1.4. Discordance within the Cilician Groups

The other prominent discordance within the Cilician groups is that mtDNA reveals two deeply divergent monophyletic groups (MHG4 in both Cilician and Narlı plains and MHG5 in mostly Narlı plain about 3.8 Ma) but nuDNA does not

support such a divergence and the pattern of geographic distribution. The main barrier between the Cilician and Narlı plain, probably separating MHG4 and MHG5, is the Amanos mountain range. It has developed recently (about 3.7-3.6 Ma) as a result of the modern pattern of plate motions (Seyrek et al., 2007; 2008). This mountain range possibly become a significant geographic barrier to gene flow within a few hundred thousand years; the development of two distinct mitochondrial DNA haplogroups due to a high mutation rate and small effective population size. Unlike mtDNA, nuDNA has a relatively slow evolutionary rate and a large effective population size, so that this mountain range has not been an effective barrier enough to prevent gene flow. Particulary, in the northern part of the range, upper reaches of rivers in the Cilician plain and Narli plains are present in the close proximity, which could make gene flow among water frog populations a possible. Moreover, recent dispersal of water frogs by human is possible because water frog collecting is very widespread in the Cilician and Narli plains (Akın & Bilgin, 2010). This might increase genetic mixture among mitochondrial haplogroups and nuclear alleles.

3.4.1.5. *Ridibundus* versus Anatolia groups

Discordance between the Ridibundus group and the Anatolia groups is evident from the geographic pattern of mtDNA and nuDNA markers in central and eastern Anatolia. mtDNA data reveal only haplotypes specific to MHG6 (cf. *bedriagae*; Anatolia group) in these regions, whereas nuDNA data also document the existence of allele groups (RID2 and RID3) from the Ridibundus MAG. Very deep divergence between these two groups is supported by both markers, their estimated divergence time corresponding to the MSC (~6-5 Ma).

As near the Mediterranean Sea, dramatic environmental changes have also occurred farther north and east. The region extending west-east from the Vienna area of Austria to Kazakhstan and Turkmenistan and north-south from central Ukraine to the modern Black Sea coast of Turkey was formerly covered by a vast water body known as the Paratethys Sea (see, e.g., Rögl, 1999, or Popov et al., 2006, for detailed documentation and palaeogeographic maps). Around 12 Ma this become isolated from the Mediterranean Sea as a result of uplift of intervening barriers, and subsequently developed into a reduced salinity environment. The Paratethys included the modern Black and Caspian Sea basins and other lowlands farther west such as the Hungarian Plain and the Vienna Basin. Subsequent vertical crustal motions, both uplift and subsidence, have subdivided the Paratethys region into these distinct basins that are now separated by uplands or by mountain ranges such as Greater and Lesser Caucasus ranges (Popov et al., 2006). During the peak of the MSC the Black Sea basin, then apparently isolated from both the Mediterranean Sea and the inflow from the River Danube, experienced a significant fall in water level, as is illuatrated in the sequence of palaeogeographic maps by Bache et al. (2012); this event is known as the Pontian desiccation. Fluctuations in the level of the Black and Caspian seas have continued during the Pliocenne and Quaternary, a notable instance being the transition in the Caspian basin from the Kimmerian lowstand to the Akchagyl highstand in the Middle Pliocene, although there is uncertainty in the timing of this event (e.g., ~4.2 Ma, Steininger et al., 1996; ~3.4 Ma; Hall et al., 2009; ~3.2 Ma, Van Baak et al., 2013). Following the mid-Pliocene climatic optimum, when conditions in the study region were warmer and wetter than at present (e.g., Haywood et al., 2000), the climate has become cooler and more arid, and has fluctuated as a result of Milankovitch forcing in response to variations in the Earth's orbit (e.g., Rohling and Hilgen, 1991; Ehlers and Gibbard, 2007). The transition from the Early Pleistocene to the Middle Pleistocene, circa 800 ka, involved a switch from dominance of the climate by shorter-period fluctuations to a predominant 100 kyr periodicity, known as the Mid-Pleistocene Revolution (MPR), which has been associated with even colder and more arid conditions during the cold stages (e.g., Maslin and Ridgewell, 2005; Head et al., 2008). Associated feedbacks, whereby reduced vegetation cover at these times of harsh climate results in faster erosion, have resulted in increased rates of vertical crustal

motion at this time, contributing to the development of topographic relief (e.g., Westaway et al., 2009). During the Late Pleistocene to Holocene transition, the Black Sea was isolated from the global marine environment and was a fresh to brackish water lake, in part because of the melt water entering its via the River Danube, from the Alpine ice sheet, and from Scandinavian ice sheet, via the River Dnieper (e.g., Chepalyga, 2007). This period ended around 9.4 ka with the incursion of Mediterranean water over the Bosphorus sill, creating the modern marine-connected Black Sea (e.g., Ryan et al., 2003; Hiscott et al., 2007).

We infer that this combination of effects of climate change and crustal deformation affecting the Black Sea and its surroundings around the time of the MSC resulted in the primary divergence between the Ridibundus group and Anatolia groups. However, the geographical pattern of both mtDNA and nuDNA data suggests that these two groups may have come into secondary contact as recently as the latest Pleistocene or earliest Holocene, possibly during the period of lacustrine isolation noted above. Thus, introgression might have occurred between these groups (both in northern Turkey and in eastern Europe), until the more saline conditions created by the marine reconnection during the Holocene reduced the distribution of both groups (which both occur in Ukraine according to both mtDNA and nuDNA data). In contrast, evidence of introgression is only supported by nuDNA, not mtDNA, in Central to Eastern Anatolia. This could be explained by male-biased dispersal of the Ridibundus group into Anatolia during low-salinity conditions of Black Sea (Holenweg Peter, 2001). In the course of time, gene flow among the high density of local Anatolia females and the less frequent invading ridibundus males would result in individuals carrying the haplotype specific for Anatolia but their genotypes would be heterozygous alleles from both Anatolia and Ridibundus groups. It follows that repeated backcrossing throughout several generations could produce several combinations of heterozygous and homozygous individuals in each of the groups. The geographic patterns of mtDNA and nuDNA in central and eastern Anatolia might thus be explained.

3.4.1.6. Discordance Patterns within the Anatolia MAG

As in the groups already discussed, although mtDNA results support well separated monophyletic groups, nuDNA reveals unresolved relationships among some clades (MAGc, d and e) within the Anatolia MAG. Thus, mtDNA subgroups indicate divergence times of 1.5-1.0 Ma, reflecting the MPR, whereas time estimates of nuDNA subgroups indicate 2.1-0.8 Ma (as a result of divergence of MAGa earlier than the rest), covering both the Pliocene-Pleistocene boundary and the MPR, thus indicating spans of time when significant global coolings have occurred (e.g., Ehlers and Gibbard, 2007).

The first discordant pattern is caused by the caralitanus-related group (MAGa in nuDNA, MAGa (cf. caralitanus) in mtDNA): cf. caralitanus, recognized from mtDNA, only occurs in the south-western Anatolia, whereas the MAGa allele group, recognized from nuDNA, occurs throughout the Black Sea region and eastern and south-eastern Anatolia, as well as south-western Anatolia. Both the mtDNA and nuDNA markers indicate that these *caralitanus* specific groups may well be the oldest group within the Anatolia MAG. Decades ago, Kosswig (1955) proposed that much of the biogeography of Anatolia is explicable in terms of the existence of the 'Central Anatolian Lake System', a vast, interconnected wetland environment that extended across much of Anatolia, which he believed existed in the Pliocene. One of course has to be extremely careful with old concepts such as this, because the chronologies of the 'continental' and 'marine' Pliocene were not standardized and much of the stratigraphic record that was then thought to be 'continental Pliocene' subsequently turned out to be Late Miocene (e.g., Berggren and Van Couvering, 1974, pp. 37-56). Nonetheless, this idea is consistent with the wet climate now evident in Anatolia during the Pliocene (e.g., Haywood et al., 2000), and with the much lower relief that existed then compared with at present, prior to phases of post-Pliocene uplift that are estimated as several hundred metres in western Anatolia, rising eastward significantly towards eastern Anatolia, and the many sites where large lakes can be reconstructed and dated to the Pliocene (e.g., Westaway et al., 2006; Seyrek et al., 2008; Demir et al., 2009). We therefore suggest that a widespread Pliocene lacustrine environment, as envisaged by Kosswig (1955), made possible the dispersal across most of Anatolia of the caralitanus-related allele group MAGa that is now evident from the nuDNA records as the retention of ancestral polymorphism in populations.

The second inconsistency between the mtDNA and nuDNA markers occurs in the cf. *cerigensis* group, which has been described as a new species, *P. cerigensis*, in Rhodos and Karpathos islands (Beerli et al., 1994). mtDNA data indicate that both islands and the adjacent mainland (Anatolia) have identical mtDNA haplotypes, whereas nuDNA indicates that alleles of islands (MAGc) was distinct from the mainland. This inconsistency could be as a result of migration of few individuals versus genetic drift.

3.4.2. Systematic Implications

For a long time, Anatolian water frogs have been designated as *P. cf. bedriagae*. Notwithstanding the extensive overlapping and mixing distribution of distinct genetic lineages evident from both nuDNA and mtDNA markers, which occur not only in transition zones but also across wider geographic regions, Anatolian water frogs clearly consist of two distinct genetic lineages. The first and larger group is the Anatolia group (Anatolia MAG, nuDNA; MHG6, mtDNA), consisting of several subgroups as previously discussed (Figs 3.4 and 3.7). The second and geographically more restricted group is the Cilician group (CIL1, CIL3, and BED1, nuDNA; MHG4 and MHG5, mtDNA), likewise including several subgroups (Figs 3.4 and 3.7). The extent of its distribution southward and its relationship with the Levant group are, however, unclear because of the lack of samples from Syria and Lebanon. These results indicate that Anatolian water frogs are conspecific neither with *P. ridibundus* (Europe) nor with *P. bedriagae* (the Levant); they therefore warrant species status and the names of these two new

species should be clarified. This study revealed the possibility of an extensive gene flow among several genetically distinct water frog lineages despite biological species concept. Several recent studies, however, showed that speciation can be possible with ongoing gene flow (Nosil, 2008; Feder et al., 2012).
3.5. References

Akın, Ç., Bilgin, M., Bilgin, C.C. (2010a) Discordance between ventral colour and mtDNA haplotype in the water frog Rana (ridibunda) caralitana, 1988 Arıkan. *Amphibia-Reptilia* 31: 9–20.

Akın, Ç., Bilgin, C.C., Beerli, P., Westaway, R., Ohst, T., Litvinchuk, S.N., et al. (2010b) Phylogeographic patterns of genetic diversity in eastern Mediterranean water frogs were determined by geological processes and climate change in the Late Cenozoic. *J Biogeogr* 37: 211–2124.

Akın, Ç. and Bilgin, C.C. (2010) Türkiye'de Su Kurbağalarının Toplanması, İşlenmesi ve İhracatına İlişkin Ön Rapor (KKGM'ye sunulmuştur). ODTÜ, Ankara, 10 pages.

Anastasakis, G., Piper, D.J.W., Dermitzakis, M.D., Karakitsios, V., (2006) Upper Cenozoic stratigraphy and paleogeographic evolution of Myrtoon and adjacent basins, Aegean Sea, Greece. *Mar Petrol Geol* 23: 353-369.

Avise, J.C., (1994) Molecular Markers, Natural History, and Evolution. Chapman & Hall, New York.

Bache, F., Popescu, S.-M., Rabineau, M. et al. (2012) A two-step process for the reflooding of the Mediterranean after the Messinian Salinity Crisis. *Basin Res* 24: 125-153.

Ballard, J., Whitlock, M. (2004) The incomplete natural history of mitochondria. *Molecular Ecol* 13: 729-744.

Bandelt, H. J., Forster, P., Röhl, A. (1999) Median-joining networks for inferring intraspecific phylogenies. *Mol Biol Evol* 16: 37-48.

Beerli, P., Hotz, H., Tunner, H., Heppich, S., Uzzell, T. (1994) Two new water frog species from the Aegean islands Crete and Karpathos (Amphibia, Salientia, Ranidae). *Notulae Naturae, Academy of Natural Sciences of Philadelphia* 470:1-9.

Beerli, P., Hotz, H., Uzzell, T. (1996) Geologically dated sea barriers calibrate a protein clock for Aegean water frogs. *Evolution* 50: 1676–1687.

Berggren, W.A., Van Couvering, J.A., (1974) The Late Neogene: Biostratigraphy, geochronology and paleoclimatology of the last 15 million years in marine and continental sequences. Developments in Palaeontology and Stratigraphy, 2. Elsevier, Amsterdam, 215 pp.

Bridgland, D.R., Westaway, R., Romieh, M.A., Candy, I. et al. (2012) The River Orontes in Syria and Turkey: Downstream variation of fluvial archives in different crustal blocks. *Geomorphology* 165-66: 25-49.

Butler, R.W.H., Spencer, S., Griffiths, H.M., (1998) The structural response to evolving plate kinematics during transpression: evolution of the Lebanese restraining bend of the Dead Sea Transform. In: Dewey, J.F., Holdsworth, R.E., Strachan, R.A. (eds.), Transpression and transtension zones. Geological Society, London, Special Publications, 135, 81-106.

Chepalyga, A.L. (2007) The late glacial great flood in the Ponto-Caspian basin. In: Yanko-Hombach, V.V., Gilbert, A.S., Panin, N., Dolukhanov, P.M. (Eds.), The Black Sea Flood Question: Changes in Coastline, Climate and Human Settlement. Springer, New York.

Cosentino, D., Gliozzi, E., Pipponzi, G. (2007) The late Messinian lago-mare episode in the Mediterranean basin: preliminary report on the occurrence of Paratethyan ostracod fauna from central Crete (Greece). *Geobios* 40: 339-349.

Coyne, J.A., Orr, H.A. (2004) Speciation. Sinauer Associates, Sunderland, MA. 545 pp.

Currat, M., Ruedi, M., Petit, R.J., Excoffier, L., (2008) The hidden side of invasions: massive introgression by local genes. *Evolution* 62: 1908–1920.

Degnan, J.H., Rosenberg, N.A. (2009) Gene tree discordance, phylogenetic inference and the multispecies coalescent. *Trends Ecol Evol* 24: 332-340.

Demir, T., Seyrek, A., Guillou, H., Scaillet, S., Westaway, R., Bridgland, D., (2009) Preservation by basalt of a staircase of latest Pliocene terraces of the River Murat in eastern Turkey: evidence for rapid uplift of the eastern Anatolian Plateau. *Global Planet Change* 68: 254-269.

Dermitzakis, M. (1990) Paleogeography, geodynamics processes and event stratigraphy during the Late Cenozoic of the Aegean area. International symposium on: biogeographical aspect of insularity, Roma 1987. *Accad Naz Lincei* 85: 263-288.

Dermitzakis, M., Papanikolaou, D.J. (1981) Paleogeography and geodynamics of the Aegean region during the neogene. *Ann Geol Pays Hellen* 20: 245-289.

Drummond, A. J., Ho, S. Y. W., Phillips, M. J., Rambaut, A. (2006) Relaxed phylogenetics and dating with confidence. *PLoS Biol* 4: E88.

Drummond, A. J., Suchard, M. A., Xie, D., Rambaut, A. (2012) Bayesian phylogenetics with BEAUti and the BEAST 1.7. *Mol Biol Evol* DOI: 10.1093/molbev/mss075.

Ehlers, J., Gibbard, P.L. (2007) The extent and chronology of Cenozoic global glaciation. *Quatern Int* 164–165: 6–20.

Edwards, S.V. (2009) Is a new and general theory of molecular systematics emerging? *Evolution* 63: 1-19.

Feder, J., Egan, S., Nosil, P. (2012) The genomics of speciation-with-gene flow. *Trends Genet* 28:342–350.

Freeand, J.R., Boag, P.T (1999) The mitochondrial and nuclear genetic homogeneity of the phenotypically diverse Darwin's ground finches. *Evolution* 53: 1553-1563.

Funk, D.J., Omland, K.E. (2003) Species-level paraphyly and polyphyly: frequency, causes and consequences with insights from animal mitochondrial DNA. *Ann Rev Ecol Evol Syst* 34: 397-423.

Excoffier, L., Lischer H.E.L. (2010) Arlequin suite ver 3.5: A new series of programs to perform population genetics analyses under Linux and Windows. *Mol Ecol Resour* 10: 564-567.

Garfunkel, Z., (1981) Internal structure of the Dead Sea leaky transform (rift) in relation to plate kinematics: *Tectonophysics* 80: 81-108.

Garfunkel Z. (1988) The pre-Quaternary geology of Israel. In: Yom-Tov Y, Tchernov E, eds. *The zoogeography of Israel*. Dordrecht: Dr Junk, 7–34.

Götz, L.-G. (1996) Beschreibung und Vergleich der Tektonik pazifischer und mediterraner Backarc-Becken hergeleitet aus echographischen und bathymetrischen Vermessungen. Ber. Zentrum Meeres- Klimaforschung, Reihe C: *Geophysik* 9: 1–165.

Hadjisterkotis, E., Masala, B., Reese. D.S. (2000) The origin and extinction of the large endemic Pleistocene mammals of Cyprus. *Biogeographia* 21: 593-606.

Hall, J., Aksu, A.E., Calon, T.J., Yaşar, D. (2005) Varying tectonic control on basin development at an active microplate margin: Latakia Basin, Eastern Mediterranean. *Mar Geol* 221: 15–60.

Hasegawa, M., Kishino, H., Yano, T.A. (1985) Dating the human-ape splitting by a molecular clock of mitochondrial DNA. *J Mol Evol* 22: 160-174.

Haywood, A.M., Sellwood, B.W., Valdes, P.J. (2000) Regional warming: Pliocene (3 Ma) paleoclimate of Europe and the Mediterranean. *Geology* 28: 1063-1066.

Heads, M. (2005) Dating nodes on molecular phylogenies: a critique of molecular biogeography. *Cladistics* 21: 62–78.

Head, M.J., Pillans, B., Farquhar, S.A. (2008) The Early–Middle Pleistocene Transition: characterization and proposed guide for the defining boundary. *Episodes* 31: 255-259.

Hejl, E., Grave, J.D., Riedl, H., Weingartner, H., Haute, P.V.D. (2008) Fission-track thermochronology of the Middle Aegean Island Bridge implications for Neogene geomorphology and palaeogeography. *Zeitschrift der Deutschen Gesellschaft für Geowissenschaften* 159: 495-512.

Hiscott, R.N., Aksu, A.E., Muddie, P.J., Marret, F., Abrajano, T., Kaminski, MA., Evans, J., Çakıroğlu, A.I., Yaşar, D. (2007) A gradual drowning of the southwestern Black Sea shelf: evidence for a progressive rather than abrupt Holocene reconnection with the eastern Mediterranean Sea through the Marmara Sea Gateway. *Quatern Int* 167-168: 9-34.

Holenweg Peter, A.-K. (2001) Dispersal rates and distances in adult water frogs, *Rana lessonae*, *R. ridibunda*, and their hybridogenetic associate *R. esculenta*. *Herpetologica* 57 (4): 449-460.

Horowitz, A. (2001) The Jordan Rift Valley. Lisse, Netherlands: A.A. Balkema.

Huelsenbeck, J.P., Rannala, B. (2004) Frequentist properties of Bayesian posterior probabilities of phylogenetic trees under simple and complex substitution models. *Systematic Biol* 53: 904–913.

Jurka, J., Drazkiewicz, A. (2002) TDR22: a putative non-autonomous DNA transposon from Danio rerio. *Repbase Reports* 2(3): 15-15.

Kissel, C., C. Laj. (1988) The Tertiary geodynamical evolution of the Aegean arc: a paleomagnetic reconstruction. *Tectonophysics* 146: 183–201.

Kosswig, C. (1955) Zoogeography of the Near East. Syst Zool 4: 49-73.

Köhler, C.M., Heslop, D., Krijgsman, W., Dekkers, M.J. (2010) Late Miocene paleoenvironmental changes in North Africa and the Mediterranean recorded by geochemical proxies (Monte Gibliscemi section, Sicily) *Palaeogeogr Palaeocl* 285: 66-73.

Krijgsman, W. (2002) The Mediterranean: mare nostrum of earth sciences. *Earth Planet Sci Lett* 205: 1-12.

Krijgsman, W., Hilgen, F.J., Raffi, I., Sierro, F.J. Wilson, D.S. (1999a) Chronology, causes and progression of the Messinian salinity crisis. *Nature* 400: 652–655. Krijgsman, W., Langereis, C.G., Zachariasse, W.J., Boccaletti, M., Moratti, G., Gelati, R., Iaccarino, S., Papani, G. Villa, G. (1999b) Late Neogene evolution of the Taza- Guercif Basin (Rifian Corridor, Morocco) and implications for the Messinian salinity crisis. *Mar Geol* 153: 147–160.

Krijgsman, W., Stoica, M., Vasiliev, I., Popov, V.V. (2010) Rise and fall of the Paratethys Sea during the Messinian Salinity Crisis. *Earth Planet Sc Lett* 290: 183-191.

Leaché, A.D. (2009) Species tree discordance traces to phylogeographic clade boundaries in North American Fence lizards (Sceloporus). *Syst Biol* 58: 547–559.

Liu, K., Wang, F., Chen, W., et al. (2010) Rampant historical mitochondrial genome introgression between two species of green pond frogs, *Pelophylax nigromaculatus* and *P. plancyi. BMC Evol Biol* 10: 1-14.

Maddison, W.P., Knowles, L.L. (2006) Inferring phylogeny despite incomplete linaege sorting. *Syst Biol* 55(1): 21-30.

Maroja, L.S., Andres, J.A., Walters, J.R., Harrison, R.G. (2009) Multiple barriers to gene exchange in a field cricket hybridzone. *Biol J Linn Soc* 97: 390–402.

Martin, D.P., Lemey, P., Lott, M., Moulton, V., Posada, D., Lefeuvre, P. (2010) RDP3: a flexible and fast computer program for analyzing recombination. *Bioinformatics* 26: 2462-2463.

Maslin, M.A., Ridgwell, A.J. (2005) Mid-Pleistocene revolution and the 'eccentricity myth'. In: M.J. Head and P.L. Gibbard (Editors), Early–Middle Pleistocene transitions: the land–ocean evidence: *Geological Society of London Special Publication* 247: pp. 19–34.

Mavarez, J., Salazar, C., Bermingham, E., Salcedo, C., Jiggins, C.D., Linares, M. (2006) Speciation by hybridization in Heliconius butterflies. *Nature* 441:868–871

Nei, M. (1987) Molecular evolutionary genetics. Columbia University Press, New York.

Nosil, P. (2008) Speciation with gene flow could be common. Mol Ecol 17(9): 2103-2106.

Petit, R.J., Excoffier, L. (2009) Gene flow and species delimitation. *Trends Ecol Evol* 24: 386–393.

Plötner, J., Ohst, T., Böhme, W., Schreiber, R. (2001) Divergence in mitochondrial DNA of Near Eastern water frogs with special reference to the systematic status of Cypriote and Anatolian populations (Anura, Ranidae). *Amphibia-Reptilia* 22: 397–412.

Plötner, J., Uzzell, T., Beerli, P., Spolsky, C., Ohst, T., Litvinchuk, S. N., Guex, G.-D., Reyer, H.-U., Hotz, H. (2008) Widespread unidirectional transfer of mitochondrial DNA: a case in western Palearctic water frogs. *J. Evol Biol* 21: 668–681.

Plötner, J., Köhler, F., Uzzell, T., Beerli, P., Schreiber, R., Guex, G.-D. & Hotz, H. (2009) Evolution of serum albumin intron-1 is shaped by a 5' truncated non-long terminal repeat retrotransposon in western Palearctic water frogs (Neobatrachia). *Mol Phylogenet Evol* 53: 784–791.

Plötner, J., Uzzell, T., Beerli, P., Akın, Ç., Bilgin, C.C., Haefeli, C., Ohst, T., Köhler, F., Schreiber, R., Guex, G.-D., Litvinchuk, A. N., Westaway, R., Reyer, H.-U. & Hotz, H. (2010) Genetic divergence and evolution of reproductive isolation in eastern Mediterranean water frogs. Evolution in action. Case studies in adaptive radiation and the origin of biodiversity. Special volume from the SPP 1127 'Radiations – Genesis of Biological diversity' of the DFG (ed. by M. Glaubrecht), pp. 373–403. Springer, Heidelberg, Berlin.

Plötner, J., Baier, F., Akın, Ç., Mazepa, G., Schreiber, R., Beerli, P., Litvinchuk, S.N., Bilgin, C.C., Borkin, L., Uzzell, T. (2012) Genetic data reveal that water frogs of Cyprus (genus Pelophylax) are an endemic species of Messinian origin. *Zoosyst Evol* 88: 261-283.

Popov, S.V., Shcherba, I.G., Ilyina, L.B., Nevesskaya, L.A., Paramonova, N.P., Khondkarian, S.O., Maygar, I. (2006) Late Miocene to Pliocene palaeogeography of the Paratethys and its relation to the Mediterranean. *Palaeogeog Palaeocl* 238: 91-106.

Rambaut, A., Drummond, A.J. (2009) *Tracer v1.5.* Available at http://beast.bio.ed.ac.uk/Tracer (last accessed 04 March 2012).

Ring, U., Glodny, J., Will, T.M., Thompson, S.N. (2010) The retreating Hellenic subduction system: high-pressure metamorphism, exhumation, normal faulting and large-scale extension. Annu Rev Earth Pl Sc 38: 45-76.

Rohling E.F, Hilgen, F.J. (1991) The eastern Mediterranean climate at times of sapropel formation: a review. *Geol Mijnbouw* 70: 253-264.

Royden, L.H., D.J. Papanikolaou (2011) Slab segmentation and Late Cenozoic disruption of the Hellenic Arc. *Geochem Geophy Geosy* 12: Q03010, 24 pp., doi: 10.1029/2010GC003280.

Rögl, F. (1999) Mediterranean and Paratethys. Facts and hypotheses of an Oligocene to Miocene paleogeography (short overview). *Geol Carpath* 50: 339-349.

Ryan, W.B.F., Major, C.O., Lericolais, G., Goldstein, S.L. (2003) Catastrophic flooding of the Black Sea. *Annu Rev Earth Pl Sc* 31: 525-554.

Salzburger, W., Baric, S., Sturmbauer, C. (2002) Speciation via introgressive hybridization in east African cichlids? *Mol Ecol* 11: 619-625.

Schwarz, G. (1978). Estimating the dimension of a model. Ann Stat 6 (2): 461-464.

Seehausen, O. (2004) Hybridization and adaptive radiation. *TRENDS Ecol Evol* 4(19): 198-207.

Seyrek, A., Demir, T., Pringle, M.S., Yurtmen, S., Westaway, R., Beck, A., Rowbotham, G., (2007) Kinematics of the Amanos Fault, southern Turkey, from Ar/Ar dating of offset Pleistocene basalt flows: Transpression between the African and Arabian plates. *Geological Society, Special Publications,* 290: London, pp. 255–284.

Seyrek, A., Westaway, R., Pringle, M., Yurtmen, S., Demir, T., Rowbotham, G., (2008) Timing of the Quaternary Elazığ volcanism, eastern Turkey, and its significance for constraining landscape evolution and surface uplift. *Turk J Earth Sci* 17: 497-541.

Seyrek, A., Demir, T., Westaway, R., Guillou, H., Scaillet, S., White, T.S., Bridgland, D.R. (2014) The kinematics of central-southern Turkey and northwest Syria revisited. *Tectonophysics* in press (online preprint available).

Steininger, F.F., Rögl, F. (1984) Paleogeography and palinspastic reconstruction of the Neogene of the Mediterranean and Paratethys. In: Dixon JE, Robertson AHF (eds), The Geological Evolution of the Eastern Mediterranean. *Geological Society London SpecialPublications* 17: 659-668 (reprinted 1996).

Steininger, F.F., Berggren, W.A., Kent, D.V., Bernor, R.L., Sen, S., Agustí, J., (1996) Circum-Mediterranean Neogene (Miocene and Pliocene) marinecontinental chronologic correlations of European mammal units. In: Bernor, R.L., Fahlbusch, V., Mittmann, H.-W. (Eds), The Evolution of Western Eurasian Neogene Mammal Faunas. Columbia Univ Press, New York, pp. 7-46.

Sumida, M., Kanamori, Y., Kaneda, H., Kato, Y., Nishioka, M., Hasegawa,
M., Yonekawa, H. (2001) Complete nucleotide sequence and gene rearrangement of the mitochondrial genome of the Japanese pond frog *Rana nigromaculata*. *Genes Genet Syst* 76: 311–325.

Tamura, K. (1992) Estimation of the number of nucleotide substitutions when there are strong transition-transversion and G + C-content biases. *Mol Biol Evol* 9: 678-687.

Tamura, K., Nei, M. (1993) Estimation of the number of nucleotide substitutions in the control region of mitochondrial DNA in humans and chimpanzees. Molecular Biology and Evolution, 10: 512–526.

Tamura, K., Peterson, D., Peterson, N., Stecher, G., Nei, M., Kumar, S. (2011) MEGA5: Molecular Evolutionary Genetics Analysis using Maximum Likelihood, Evolutionary Distance, and Maximum Parsimony Methods. *Mol Biol Evol* 28: 2731-2739.

Toews D.P.L., Brelsford, A. (2012) The biogeography of mitochondrial and nuclear discordance in animals. *Mol Ecol* 21: 3907-3930.

Van Baak, C.G.C., Vasiliev, I., Stoica, M., Kuiper, K.F., Forte, A.M., Aliyeva, E., Krijgsman, W. (2013) Paleomagnetic A magnetostratigraphic time frame for Plio-Pleistocene transgressions in the South Caspian Basin, Azerbaijan. *Global Planet Change* 103: 119–134.

Vasiliev, I., Iosifidi, A.G., Khramov, A.N., Krijgsman, W., Kuiper, K., Langereis, C.G., Popov, V.V., Stoica, M., Tomsha, V.A., Yudin, S.V. (2011) Magnetostratigraphy and radio-isotope dating of upper Miocene-lower Pliocene sedimentary successions of the Black Sea (Taman Peninsula, Russia). *Palaeogeography, Palaeoclimatology, Palaeoecology* 310(3-4): 163-175.

Westaway, R. (2004) Kinematic consistency between the Dead Sea Fault Zone and the Neogene and Quaternary left-lateral faulting in SE Turkey. *Tectonophysics* **391**: 203–237.

Westaway, R. (2006) Cenozoic cooling histories in the Menderes Massif, western Turkey, may be caused by erosion and flat subduction, not low-angle normal faulting. *Tectonophysics* 412: 1-25.

Westaway, R., Guillou, H., Yurtmen, S., Beck, A., Bridgland, D., Demir, T., Scaillet, S., Rowbotham, G., (2006) Late Cenozoic uplift of western Turkey: Improved dating of the Kula Quaternary volcanic field and numerical modelling of the Gediz river terrace staircase. *Global Planet Change* 51: 131-171.

Westaway, R., Bridgland, D. R., Sinha, R., Demir, T. (2009) Fluvial sequences as evidence for landscape and climatic evolution in the Late Cenozoic: a synthesis of data from IGCP 518. *Global Planet Change* 68: 237–253.

3.6. Appendices

Table 3.2. Locality information for mitochondrial ND2 and ND3 haplotypes. This table gives details of localities, collection or laboratory IDs, and old and new haplotype IDs for the investigated individuals. Since several additional new haplotypes have been included for Anatolian groups (MHG4, 5 and 6), a new haplotype abbrevation system is used to enable the distribution of haplotypes to be easily followed. Samples used for the first time in this study are marked with a grey background, while samples taken from previous studies are listed with white backgrounds. Accession numbers starting with AB, AJ, AM, GQ and GU were taken from the EMBL database. They correspond to the publications of Sumida et al., (2001), Plötner et al., (2001, 2008, 2009), and Akın et al., (2010a,b). Abbreviations denote: MHG, main haplogroup; ND2, mitochondrial gene for NADH dehydrogenase subunit 2; ND3, mitochondrial gene for NADH dehydrogenase subunit 3; ZFMK, Zoologisches Forschungsmuseum und Alexander Koenig, Bonn; ZISP, Zoological Institute St. Petersburg; and ZMB, Zoologisches Museum Berlin.

Country	Locality	Lat	Long		Haplo (c	Haplotype ID (old)		otype ID new)	Accession Number		MHG/ Species
				Collection/ Laboratory ID	ND2	ND3	ND2	ND3	ND2	ND3	species
Albania	Durräc	41 3247	10 4268	-		RE4				AJ310330	MHG1
Albania	Duries	41.3247	19.4208	-	R11	RE4			AM749704	AJ310330	MIIOT
Armenia	Megri	38.9135	46.2340	ZISPSp.308		AN41		ANT16		GU812198	MHG6c
	Alepu swamps near Sozopol	42.3995	27.6930	ZMB47094	R1	RE1			AM900661	AJ310338	MHG1
	Kazanlak	42.4006	25.3010	Hotz17994		RE1				AJ310338	
Dulgorio				Hotz17995		RE1				AJ310338	MHG1
Bulgaria				Hotz17996		RE1				AJ310338	
				Hotz17997		RE1				AJ310338	
	Nessebar	42.6054	27.6062	ZMB51288	R2	RE1			AM749712	AJ310338	MHG1
				Hotz19410	A1	AN12	CA1	CAR12	GU812088	GQ902088	MHG6a
				Hotz19411	CP6	CY4			HE820104	HE861955	
Cyprus	Northern Cyprus	35.1855	33.5674	Hotz19412	CP6	CY4			HE820104	HE861955	MHG3
			-	Hotz19413	CP6	CY4			HE820104	HE861955	
				Hotz19416	A1	AN12	CA1	CAR12	GU812088	GQ902088	MHG6a

Table 3.2. (continued).	
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	Argaka reservoir	35.047	32.506	ZFMK48755		CY7			HE861948	MHG3
	Potamos tis Ezousas near Episkopi	34.671	32.916	ZFMK48757		CY1			HE861949	MHG3
				P21	CP8	CY3		HE820092	GU812161	
				P22	CP8	CY3		HE820093	GU812161	
	Cape Greco (water reservoir in the west)	34.9794	34.0548	P23	CP8	CY3		HE820094	GU812161	MHG3
				P24	CP8	CY3		HE820095	GU812161	
				P25	CP5	CY3		HE820096	GU812161	
	Stream close to Gialia	35 0769	32 5690	P32	CP4	CY6		HE820097	HE861950	MHG3
	Sitean close to Glana	33.0707	32.3070	P33	CP4	CY6		HE820098	HE861951	WIIG5
	Gönyeli dam	35.2332	33.2937	Р3	CP8	CY3		HE820099	GU812161	MHG3
	Lefkosa (stream system of the	35.2017	33.3522							
Cuprus	Pediaios)			P2	CP8	CY3		HE820100	GU812161	MHG3
	Maroullenas-1	35.0084	33.1477	P5	CP8	CY4		HE820101	HE861952	MHG3
Cyprus	Maroullenas-2	34.9996	33.1423	P6	CP8	CY4		HE820102	HE861953	MHG3
	Maroullenas-3	35.0054	33 1453	P7	CP8	CY4		HE820103	HE861954	MHG3
			55.1455	P8	CP6	CY4		HE820104	HE861955	MIIOS
				-	CP1	CY1		GU812078	AJ310334	MHG3
	Nicosia-1	35.1682	33.5273	-	CP1	CY1		GU812078	AJ310334	
				ZFMK48760		CY1			AJ310334	
				DB238		CY2			GU812160	
	Nicosia-2	35.1951	33.3189	DB239		CY3			GU812161	MHG3
				DB240		CY3			GU812161	
1				P15	CP8	CY3		HE820105	GU812161	MHG3
	Nicosia (channel in the forest part)	35 1707	33.3567	P16	CP8	CY3		HE820106	GU812161	
	ricessa (chainer in the forest part)	55.1707		P17	CP8	CY3		HE820107	GU812161	
				P18	CP7	C74		HE820108	HE861956	

Table 3.2. (continued)	•
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	Nicosia (channel in the forest part)	35.1707	33.3567	P19	CP8	CY3		HE820109	GU812161	MHG3
	2 km southwest of Palaichhori	34.9140	33.0681	P1	CP1	CY1		GU812078	AJ310334	MHG3
	Panagia (forest station at stream Peristerona)	35.0156	33.0833	P4	CP1	CY1		GU812078	AJ310334	MHG3
	Lake Paralimni	35.0358	33.9753	P20	CP8	CY3		HE820110	GU812161	MHG3
	Paristarana 1	34 9770	22.0012	P34	CP1	CY1		GU812078	AJ310334	MUC2
	T CHSteronia-1	34.9770	55.0912	P35	CP1	CY1		GU812078	AJ310334	WHO5
				P26	CP1	CY1		GU812078	AJ310334	
				P27	CP1	CY1		GU812078	AJ310334	
	Peristerona-2 (headwater)	3/ 9953	33.0926	P28	CP1	CY1		GU812078	AJ310334	MHG3
		54.7755		P29	CP1	CY1		GU812078	AJ310334	WII05
Cyprus				P30	CP1	CY1		GU812078	AJ310334	
				P31	CP1	CY1		GU812078	AJ310334	
	Potamos tou Limniti, north of Stavros	35.161	32.734	ZFMK48758		CY7			HE820111	MHG3
				Р9	CP4	CY6		HE820112	HE861957	
	Pyrgos Dam	35.1495	32.6549	P12	CP3	CY7		HE820113	HE861958	MHG3
				P13	CP4	CY6		HE820114	HE861959	
	near Stavros	35.0673	32.6310	P14	CP4	CY6		HE820115	HE861960	MHG3
				ZFMK59099	CP2	CY1		GU812079	AJ310334	
	Troodos-1	34.7269	32.9095	ZFMK590100		CY1			AJ310334	MHG3
				ZFMK590102		CY1			AJ310334	
	Troodos-2	34 9161	32.9003 -	ZMB77443	CP8	CY8		HE820116	HE861961	MHG3
		34.9161		ZISP10531	CP8	CY8		HE820117	HE861962	MIIGS

Cyprus	Lake Xyliatos	35.0375	33.0381		CP1 CP1	CY1 CY1					MHG3
				Hotz16375	B1	BN1			GU812072	AJ310322	
Egypt	Cairo	29.9638	31.2289	Hotz16376	B1	BN1			GU812072	AJ310322	MHG2
				Hotz16377		BN1				AJ310322	
France	Aramon	43.8967	46.827	-	R4	RE6			AM900652	AM900653	MHG1
Trance	St. Étienne du Gres	43.7786	4.6533	-	R1	RE1			AM900661	AJ310338	MHG1
	Batumi	41 6420	41 6800	ZMB47390	A25	AN26	AT30	ANT1	GU812112	AJ310337	MHG6c
Georgia	Batum	41.0420	41.0000	ZMB47490	A33	AN42	AT75	ANT17	GU812119	GU812199	WINGOU
Georgia	Sukhumi	42 9970	40.9825	ZMB44562	A34	AN42	AT73	ANT17	GU812120	GU812199	MHG6c
	Sukiluini	42.9970	40.9025	ZMB44563	A35	AN42	AT74	ANT17	GU812121	GU812199	WINGOU
			23.0841	Hotz17357	R5	RE5			AM749705	AM749705	
	Aliartos	38.3688		Hotz17359	R8	RE12			AM749709	AM749709	MHG1
Crassa				Hotz17360	R9	RE12			AM749710	AM749709	
Gieece	Apolakkia/Rhodos	36.0592	27.7876	Hotz18167	A8	AN24	CE2	CER4	GU812095	GU812183	MHG6b
	Apollona/Rhodos	36.2543	27.9779	-		AN24		CER4		GU812183	MHG6b
	Archipolis/Rhodos	36.3117	28.1360	Hotz18164		AN24		CER4		GU812183	MHG6b
	Archinglis/Phodos	26 2117	28 1260	Hotz17218	A8	AN24	CE2	CER4	GU812095	GU812183	MHC6b
	Archipons/Kilodos	50.5117	28.1300	-		AN24		CER4		GU812183	MHG00
				Ploetner137.03	A15	AN54	AT99	ANT29	GU812102	GU812209	
	Chios	38.3568	26.1433	Ploetner138.03	A15	AN54	AT99	ANT29	GU812102	GU812209	MHG6c
Greece				Ploetner143.03	A15	AN54	AT99	ANT29	GU812102	GU812209	
	Dadia	41 1410	26.2846	Hotz22714		RE1				AJ310338	MHG1
	Daula	41.1419	26.2846	Hotz22717		RE1				AJ310338	WIIGI
	Ikaria	37 6075	5 26.1521 -	Hotz17325	A3	AN19	CA10	CAR19	GU812090	GU812180	0 MHG6a
	Ikaria	37.6075		Hotz17327	A3	AN19	CA10	CAR19	GU812090	GU812180	

	Ikaria	37.6075	26.1521	Hotz17329	A3	AN19	CA10	CAR19	GU812090	GU812180	MHG6a
	Iviro	40,0004	22 6005	Hotz17979		RE7				GU812153	MHC1
	Ivita	40.9004	23.0005	Hotz17984		RE6				AM900653	MHOI
	Kalanistra	38.0633	21.8485	ZMB49804	R17	RE11			AM900644	AJ310329	MHG1
	Kaminia	39.7653	20.8678	ZMB49822	R14	RE11			AM749699	AJ310329	MHG1
	Kayasilas	37 8762	21 2854	Hotz19510	R12	RE13			AM749702	AM749702	MHG1
	Kavasiias	37.8702	21.2034	Hotz19528	R17	RE14			AM900644	AM749697	MITGI
	Karamoti	40.0351	24 4038	Hotz17412		RE1				AJ310338	MHG1
	Keramou	40.9331	24.4038	Hotz17413		RE2				GU812151	MHOI
	Kymina	40 5565	22 7005	Hotz17986		RE6				AM900653	MHG1
	Kyiiina	40.5505	22.1000	Hotz17988		RE6				AM900653	WIIGI
	Kythira	36.1530	22.9871	Hotz17408	R15	RE11			AM900646	AJ310329	
Graaaa				Hotz17409	R15	RE11			AM900646	AJ310329	MHG1
Gleece				Hotz17424	R15	RE11			AM900646	AJ310329	
	Lefkes	40.9015	25.8001	Hotz17430		RE1				AJ310338	MHG1
				Ploetner145.03	A13	AN54	AT92	ANT29	GU812100	GU812209	- MHG6c
	Lasvos	30 2103	26 1787	Ploetner146.03	A12	AN54	AT93	ANT29	GU812099	GU812209	
	Lesvos	39.2103	20.1787	Ploetner182.03	A14	AN54	AT91	ANT29	GU812101	GU812209	
				ZMB56920	A12	AN54	AT93	ANT29	GU812099	GU812209	
				Hotz22721		RE1				AJ310338	
	Mangana	10 9289	24 8456	Hotz22723		RE6				AM900653	MHG1
	ivialigalia	40.7207	24.0450	Hotz22724		RE6				AM900653	WIIGI
_				Hotz22725		RE1				AJ310338	
				Hotz17341	R7	RE6			AM900653	AM900653	MHG1
	Mistros 3	38.5228	23.8325	Hotz17342	R6	RE4			AM749706	AJ310330	
				Hotz17343	R7	RE6			AM900653	AM900653	

				Hotz17365		RE1				AJ310338	MHG1
	Monostiraki	40.8515	26 1026	Hotz17366		RE1				AJ310338	MHOI
	Wonastnaki	40.6515	20.1020	Hotz17367		AN38		ANT13		GU812195	MHG6c
				Hotz17368		RE1				AJ310338	MHG1
				Hotz19532	R17	RE11			AM900644	AJ310329	
	Nea Manolada	38.0316	21.3611	Hotz19533		RE11				AJ310329	MHG1
				Hotz19548	R13	RE13			AM749703	AM749702	
			27.1685	Hotz17207	A6	AN21	CE1	CER1	GU812093	GU812181	
	Olympos/Karpathos	35.7427		Hotz17208	A6	AN21	CE1	CER1	GU812093	GU812181	MHG6b
				Hotz17420	A6	AN21	CE1	CER1	GU812093	GU812181	
	Pagouria	40.9164	25.3780	Hotz17446		RE1				AJ310338	MHG1
			23.2110	Hotz12923	R3	RE1			AM749713	AJ310338	MHG1
Greece				Hotz12926		RE1				AJ310338	
Greece		39.2993		Hotz12927		RE6				AM900653	
				Hotz12928		RE1				AJ310338	
	Paradisos			Hotz12931		RE2				GU812151	
				Hotz12932		RE1				AJ310338	
				Hotz12933		RE6				AM900653	
				Hotz12934		RE8				GU812154	
				Hotz12935		RE1				AJ310338	
_	Samos	37 7641	26.9750	Hotz17322	A18	AN54	AT89	ANT29	GU812105	GU812209	MHG6c
	Sunos	57.7041	20.9750	Hotz17323	A19	AN47	AT90	ANT22	GU812106	GU812203	Mildoe
	Skala	38 6675	23 0712	Hotz19483	R10	RE11			AM900662	AJ310329	MHG1
	JKulu	50.0075	23.0712	Hotz19500	R10	RE11			AM900662	AJ310329	WIIIOI
	Lake Stimfalias/ Peloponnese	37.3452	22.1026	ZMB49267	R17	RE11			AM900644	AJ310329	MHG1
		51.3452		ZMB49820	R17	RE11			AM900644	AJ310329	

				ZMB56972	R17	RE11			AM900644	AJ310329	
				Ploetner29.03	R16	RE11			AM749698	AJ310329	
Greece	Zakynthos	37.8052	20.8627	Ploetner30.03	R18	RE11			AM900639	AJ310329	MHG1
				Ploetner31.03	R18	RE11			AM900639	AJ310329	
				Ploetner32.03	R18	RE11			AM900639	AJ310329	
Hungory	Budapest	47.5325	19.0333	ZMB44447		RE1				AJ310338	MHG1
Hungary	Kis Balaton	48.0992	20.3171	ZMB44490		RE1				AJ310338	MHG1
	Phalet village			AMPH\IRA\536		AN60	EU4	EUP6			MHG6d
				AMPH\IRA\508			T7	TE6	ACC	GU812222	
			52.6357	AMPH\IRA\509			T15	TE1	ACC	GU812218	
	Babol Sar	36.6803		AMPH\IRA\510			T15	TE1	ACC	GU812218	MHG7
				AMPH\IRA\511			T15	TE1	ACC	GU812218	
				AMPH\IRA\512			T5	TE2	ACC	GU812219	
	Badal	38.6336	44.7720	AMPH\IRA\351		AN59	EU22	EUP5	ACC	AJ310312	MHG6d
	Bastan	38.9046	44.9386	AMPH\IRA\301		AN59	EU23	EUP5	ACC	AJ310312	MHG6d
	Bavineh	33.6042	47.2026	AMPH\IRA\338		AN56	EU1	EUP2	ACC	GU812210	MHG6d
Iran				AMPH\IRA\307		AN56	EU1	EUP2	ACC	GU812210	MHG6d
				AMPH\IRA\308		AN60	EU5	EUP6	ACC	GU812213	
	Bisotun	34.4038	47.4483	AMPH\IRA\309		AN60	EU6	EUP6	ACC	GU812213	
				AMPH\IRA\310		AN64		EUP10		GU812216	
				AMPH\IRA\311		AN56		EUP2		GU812210	
	Choplu	36.4739	47.0393	AMPH\IRA\302		AN65	EU15	EUP11	ACC	GU812217	MHG6d
				AMPH\IRA\327		AN56		EUP2		GU812210	
	Choga Zanhil	32 0177	48.5451	AMPH\IRA\328		AN61		EUP7		GU812214	MHG6d
	Choqa Zanbil	52.0177		AMPH\IRA\329		AN61		EUP7		GU812214	
				AMPH\IRA\330	1	AN62		EUP8		GU812215	

				AMPH\IRA\331	AN62		EUP8		GU812215	
				AMPH\IRA\332	AN62		EUP8		GU812215	
	Choga Zaphil	32 0177	48 5451	AMPH\IRA\333	AN56		EUP2		GU812210	MHC64
	Choqa Zanon	52.0177	40.5451	AMPH\IRA\334	AN62		EUP8		GU812215	MIIGOu
				AMPH\IRA\335	AN62		EUP8		GU812215	
				AMPH\IRA\336	AN56		EUP2		GU812210	
	Galugan	37 5171	/9 3091	AMPH\IRA\523		T10	TE1	ACC	GU812218	MHG7
	Galugan	57.5171	47.5071	AMPH\IRA\524		Т9	TE1	ACC	GU812218	WIIG/
	Jeiugir	32.9873	47.8106	AMPH\IRA\313	AN59	EU2	EUP5	ACC	AJ310312	MHG6d
	Kenar Darya			AMPH\IRA\117		T12	TE1	ACC	GU812218	
			53.3131	AMPH\IRA\118		T15	TE1	ACC	GU812218	
				AMPH\IRA\119		Т8	TE1	ACC	GU812218	MHG7
Iran		36.8472		AMPH\IRA\120		T15	TE9	ACC	GU812225	
				AMPH\IRA\121		T1	TE4	ACC	GU812221	
				AMPH\IRA\122		T5	TE2	ACC	GU812219	
				AMPH\IRA\123		T15	TE1	ACC	GU812218	
	Lendj Abad	33.4511	49.0293	AMPH\IRA\312	AN60		EUP6		GU812213	MHG6d
				AMPH\IRA\112		T15	TE1	ACC	GU812218	
	Mangol	36.2505	52.3563	AMPH\IRA\113		T5	TE7	ACC	GU812223	MHG7
				AMPH\IRA\114		T6	TE8	ACC	GU812224	
	Marrave Tappe	37.7266	55.9071	AMPH\IRA\521		T1	TE4	ACC	GU812221	MHG7
				AMPH\IRA\513		T15	TE1	ACC	GU812218	
	Now Kandeh	36 7154	53 8836	AMPH\IRA\514		T14	TE1	ACC	GU812218	MHG7
		50.7154	53.8836	AMPH\IRA\515		T5	TE2	ACC	GU812219	101107
				AMPH\IRA\516		T11	TE1	ACC	GU812218	

Table 3.2. (continued).
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				AMPH\IRA\517			T15	TE1	ACC	GU812218	
				AMPH\IRA\518			T15	TE1	ACC	GU812218	
	Now Kandeh	36.7154	53.8836	AMPH\IRA\520			T15	TE1	ACC	GU812218	MHG7
				AMPH\IRA\115			T14	TE3	ACC	GU812220	
				AMPH\IRA\116			T15	TE1	ACC	GU812218	
	Khuzestan Kuli-Ali Reza	31.2499	49.6501	AMPH\IRA\530			T4	TE12	ACC	GU812228	MHG7
			49.4057	AMPH\IRA\316				TE12		GU812228	
Iran				AMPH\IRA\317				TE10		GU812226	
	Simili	31.7108		AMPH\IRA\318				TE11		GU812227	MHG7
				AMPH\IRA\319				TE12		GU812228	
				AMPH\IRA\320				TE12		GU812228	
	Sorkheh Dizaj		48.8247	AMPH\IRA\124		AN56	EU1	EUP2	ACC	GU812210	MHG6d
		38.8278		AMPH\IRA\125		AN56	EU1	EUP2	ACC	GU812210	
				AMPH\IRA\126		AN57	EU3	EUP3	ACC	GU812211	MIIGou
	Zanjan Ab Kenar	37.4474	49.3215	AMPH\IRA\501			Т9	TE1	ACC	GU812218	
	Al Kerak	31.1833	35.7000	J-06-54	B12	BN8			ACC	ACC	MHG2
	Amman	32.0334	35.8091	ZFMK63535	B5	BN3			GU812076	AJ310321	MHG2
				J-01-44	B11	BN5			ACC	GU812158	
	Bah Amman	32 2042	35 8867	J-01-55	B11	BN5			ACC	GU812158	MHG2
	Dab Animan	52.2042	55.0007	J-01-56		BN5				GU812158	WI102
Jordan				J-01-57	B8	BN3			ACC	AJ310321	
				J-05-39		BN7				ACC	
				J-05-45	B7	BN9			ACC	ACC	
	Canyon near Wadi Mujib	31.4492	35.7892	J-05-46	B7	BN9			ACC	ACC	MHG2
				J-05-47	B7	BN7			ACC	ACC	
				J-05-48	B7	BN7			ACC	ACC	

				J-03-35		BN4				AJ310319	
	16 km southward from Jesus	31 /030	35 5861	J-03-36	B9	BN5			ACC	GU812158	MHG2
	Baptisizing site	51.4959	55.5601	J-03-37		BN6				GU812159	WIII02
				J-03-38	B9	BN5			ACC	GU812158	
	King Talal Dam	32.2071	35.8517	ZFMK63514		BN6				GU812159	MHG2
				J-04-30	B10	BN5			ACC	GU812158	
Jordan				J-04-32	B12	BN8			ACC	ACC	
				J-04-33	B12	BN8			ACC	ACC	
	Mulaik Thoba	31.5533	35.7400	J-04-41	B10	BN5			ACC	GU812158	MHG2
				J-04-51	B12	BN8			ACC	ACC	
				J-04-52	B12	BN8			ACC	ACC	
				J-04-53	B12	BN8			ACC	ACC	
	Wadi Mujib	31.4455	35.8175	ZFMK64390	B4	BN5			GU812075	GU812158	MHG2
	Almaty (Alma-Ata)	43.2827	76.8720	ZMB46963	C4		S1	SP1	GU812130	GU812229	MHG8
	Aktoba	50 2007	57 2170	Sp.250		AN27		ANT2		GQ902107	MHC6a
	AKIODE	30.2997	57.2179	Sp.251		AN27		ANT2		GQ902107	MHGoe
				Sp.252	A29	AN27	AT79	ANT2	GU812116	GQ902107	
Kazakhstan				Sp.253	A29	AN27	AT79	ANT2	GU812116	GQ902107	
	Atyrau	47.5442	52.3861	Sp.254	A28	AN27	AT80	ANT2	GU812115	GQ902107	MHG6c
				Sp.255	A29	AN27	AT79	ANT2	GU812116	GQ902107	
				Sp.256		AN27		ANT2		GQ902107	
	Chaganskoy	51.0354	51.7525	Sp.257		AN27		ANT2		GQ902107	MHG6c
Kyrgyzstan	Bishkak	12 8370	74 6360	L1	C5		S2	SP1	GU812131	GU812229	MHG8
	DISHKCK	42.0379	/4.0300	L3	C5		S2	SP1	GU812131	GU812229	мпоо
Libya	Shahhat (Curana)	22 7016	21 4207	AMPH\EGY\001		RE11				AJ310329	MHG1
Libya	Snannat (Cyrene)	52.7910	21.4297	AMPH\LIB\002		RE11				AJ310329	MINUT

				AMPH\LIB\003		RE11				AJ310329	
				AMPH\LIB\004		RE11				AJ310329	
				AMPH\LIB\005		RE11				AJ310329	
				AMPH\LIB\006		RE11				AJ310329	
Libro	Shakhat (Cymona)	22 7016	21 4207	AMPH\LIB\007		RE11				AJ310329	MUC1
Libya	Shainat (Cylene)	32.7910	21.4297	AMPH\LIB\008		RE11				AJ310329	MHOT
				AMPH\LIB\009		RE11				AJ310329	
				AMPH\LIB\010		RE11				AJ310329	
				AMPH\LIB\011		RE11				AJ310329	
				AMPH\LIB\012		RE11				AJ310329	
Magadonia	Lake Dojran	41.2166	22.7333	ZMB46692	R4	RE6			AM900652	AM900653	MHG1
Macedollia	Skopje	42.0121	21.4715	ZMB46744		RE6				AM900653	MHG1
Poland	Poznan	52.3805	16.6674	Hotz18192	R1	RE1			AM900661	AJ310338	MHG1
Pomania	Sfintu Chaorgha straam mila 52	11 6121	26.8504	ZMB47433	R1	RE1			AM900661	AJ310338	MHG1
Romania	Sintu-Oneorgne-stream, inne 52	44.0424	20.8504	ZMB47434	R4	RE11			AM900652	AJ310329	WINGT
	Armavir	44.9887	41.1563	ZISP.6130-Sp.137		AN44		ANT19		GU812201	MHG6c
	Baltiysk	54.6467	19.8820	ZISPSp.273		RE3				GU812152	MHG1
	Chorgay Reservoir	45.5212	44.5678	ZISP.4255-Sp.65		RE10				GU812156	MHG1
	Dakhovskava	44 2251	40 1007	ZMB57380	A31	AN43	AT76	ANT18	GU812117	GU812200	MHG6c
Russia	Dakilovskaya	44.2231	40.1997	ZMB57388	A32	AN43	AT77	ANT18	GU812118	GU812200	WINGOC
Russia	Yekaterinburg	56.8363	60.6379	ZISPSp.244		RE6				AM900653	MHG1
	Ersi	42.0037	47.9904	ZISP.6760-Sp.136		AN27		ANT2		GQ902107	MHG6c
	Gaverdovsky	44.6281	40.0204	ZISP.6644-Sp.131		AN43		ANT18		GU812200	MHG6c
	Kaspiy Sanatorium	42.3504	48.0603	ZISPSp.260		AN40		ANT15		GU812197	MHG6c
	Mochokh	42.6510	46.6347	ZISP.6761-Sp.134		AN27		ANT2		GQ902107	MHG6c

	Moscow	55 7144	37 5857	ZMB46428	R1	RE1			AM900661	AJ310338	MHC1
	Moscow	55.7144	57.5657	ZMB45920-25/1		RE1				AJ310338	MHOT
				ZISP.6319-Sp.85-87		AN27		ANT2		GQ902107	
	Orsk	51.2177	58.6347	ZISP.6319-Sp.85-87		AN27		ANT2		GQ902107	MHG6c
				ZISP.6319-Sp.85-87		AN27		ANT2		GQ902107	
	Psebai	44.1185	40.7853	ZISP.6549-Sp.74		AN27		ANT2		GQ902107	MHG6c
				ZISP.6553-Sp.108		RE9				GU812155	
Russia	Rossosh	50.2153	39.6006	ZISP.6553-Sp.109		RE9				GU812155	MHG1
				ZISP.6553-Sp.110		RE9				GU812155	
	Sochi	43.5707	39.7625	ZISP.6751-Sp.77		AN42		ANT17		GU812199	MHG6c
				ZISPSp.247		AN31		ANT6		GU812188	
	Saint Peterhof Station	59.8853	29.9098	ZISPSp.248		AN31		ANT6		GU812188	MHG6c
				ZISPSp.249		AN43		ANT18		GU812200	
	Step	44.5687	44.8150	ZISP.3280-Sp.68		RE9				GU812155	MHG1
	Volgograd	48.7106	44.4934	ZISP.6659-Sp.126		AN27		ANT2		GQ902107	MHG6c
Serbia	Belgrade	44.8332	20.5019	ZMB46740	R1	RE1			AM900661	AJ310338	MHG1
Slovakia	Bratislava-Devin	48.1579	16.9918	SLOV88	R1	RE1			AM900661	AJ310338	MHG1
SIOvakia	Brodské	48.6941	17.0092	SLOV171	R1	RE1			AM900661	AJ310338	MHG1
Switzerland	Embrach	47.5060	8.6123	Hotz16637		RE6				AM900653	MHG1
				ZFMK61785	A40	AN55	EU11	EUP1	GU812126	AJ310313	
	Abu Kamal	34.4496	40.9386	ZFMK61787	A36	AN63	EU36	EUP9	GU812122	AJ310311	MHG6d
				ZFMK61788	A38	AN59	EU25	EUP5	GU812124	AJ310312	
Syria	Ansari Mountains	34.8166	36.1166	ZFMK60903	B2	BN2			GU812073	AJ310320	MHG2
	As Suwayda	32.6855	36.5525	ZFMK64945	B6	BN4			GU812077	AJ310319	MHG2
	Qalat al-Hisn	31 7789	36 2655	ZFMK57959	B3	BN1			GU812074	AJ310322	MHG2
	(Crac des Chevaliers)	34.7700	36.2655	ZFMK57960		BN1				AJ310322	WI102

				CA1418		CIW3		GU812163	
				CA1419		CIW1		AJ313135	MHG4
	Adana-Ceyhan Cinderesi	37.0353	35.7470	CA1420	CLE13	CIE2	ACC	GU812169	MHG5
				CA1421	CLW13	CIW3	ACC	GU812163	MHG4
				CA1422	CLW10	CIW1	ACC	AJ313135	WING4
				CA1922	CLW15	CIW3	GU812083	GU812163	MHG4
			34.8842	CA1923	CA8	CAR25	GU812089	ACC	MHG6a
	Adana-Pozantı Şekerpınarı	37.4080		CA1924		CIW3		GU812163	MHG4
				CA1925		ANT1		AJ310337	MHG6c
				CA1926		ANT1		AJ310337	WINGOU
				CA1927		CIE2		GU812169	MHG5
	Adana-Pozantı Çakıt river	37.4299	34.8764	CA1928	CLW15	CIW3	GU812083	GU812163	MHG4
Turkey				CA1929		CIW3		GU812163	
Turkey				CA1930		CIW3		GU812163	
				CA1931		CIW3		GU812163	
				CA1540	AT27	ANT1	ACC	AJ310337	MHG6c
				CA1541	AT69	ANT2	ACC	GQ902107	
	Adapazarı-Poyrazlar lake	40.8336	30.4682	CA1542		ANT1		AJ310337	
				CA1543		ANT1		AJ310337	
				CA1544		ANT2		GQ902107	
				CA1545	AT40	ANT53	ACC	ACC	
	Adapazarı Saklıgöl	40.8540	30 3018	CA1546		ANT2		GQ902107	MHG6a
	Adapazan-Sakiigoi	40.0349	50.5018	CA1547	AT69	ANT2	ACC	GQ902107	WITCOC
				CA1548	AT5	ANT1	ACC	AJ310337	1
	Adıyaman-Abuzergaffar river	37 7415	38.3354	CA1270	CLW15	CIW3	GU812083	GU812163	MHG4
		37.7415		CA1271	EU14	EUP1	ACC	AJ310313	MHG6d

				CA1272	E	EU24	EUP5	ACC	AJ310312	
				CA1273			EUP1		AJ310313	
	Adıyaman-Abuzergaffar river	37.7415	38.3354	CA1274			EUP5		AJ310312	MHG6d
				CA1275			EUP5		AJ310312	
				CA1276			EUP1		AJ310313	
				CA1287			EUP5		AJ310312	
				CA1288			EUP5		AJ310312	
Adıyaman-Börgenek Çakal stream				CA1289	E	EU14	EUP1	ACC	AJ310313	MHG6d
	37.7128	38.1663	CA1290	E	EU14	EUP23	ACC	ACC		
			CA1291	E	EU36	EUP5	GU812122	AJ310312		
				CA1292	C	LE12	CIE2	ACC	GU812169	MHG5
				CA1293			EUP1		AJ310313	MHG6d
Turkey	Turkey	37.6983	38.0799	CA1282	A	AT72	ANT1	ACC	AJ310337	MHG6c
Turkey				CA1283	E	EU36	EUP5	GU812122	AJ310312	MHG6d
	Adıyaman-Göksu river			CA1284	E	EU12	EUP1	ACC	AJ310313	MIIGOu
				CA1285			ANT4		GU812186	MHG6C
				CA1286			ANT4		GU812186	windoe
				CA1277			EUP5		AJ310312	
				CA1278			EUP1		AJ310313	
	Adıyaman-Kahta Bircik river	37.7447	38.5060	CA1279	E	EU14	EUP1	ACC	AJ310313	MHG6d
Afyo				CA1280	E	EU36	EUP5	GU812122	AJ310312	
				CA1281			EUP1		AJ310313	
				CA1726		CA9	CAR17	ACC	AJ313132	MHG6a
	Afyonkarahisar-26 Agust Natural	38,7925	30,3816	CA1727	A	T110	ANT21	GU812104	GQ902114	MHG6c
	Park 3	38.7925	30.3816	CA1728			ANT21		GQ902114	
				CA1729			CAR14		AJ313133	MHG6a

Afyonkarahisar-26 Agust Natural Park	38.7925	30.3816	CA1730			CAR14	
			CA1720			ANT21	
			CA1721		AT110	ANT21	GU812104
Cav	38.5923	30.9618	CA1722			CAR14	
, ~y			CA1723		AT41	ANT2	ACC
			CA1724		CA8	CAR17	GU812089
			CA1709		CA8	CAR17	GU812089
			CA1710			CAR14	
Afyonkarahisar-Karamık lake	38.4215	30.8869	CA1711			CAR17	
			CA1712			ANT21	
			CA1713			CAR17	
	20.9191	43.0892	CA679			EUP5	
Ağrı-Aşağı Voldüzü Cuma River			CA680		EU8	EUP12	ACC
Agii-Aşagi Tolduzu Culla River	37.0101		CA681			EUP12	
			CA683			EUP12	
			CA684			EUP12	
			CA685			EUP12	
Ağrı-Çukuralan Village	39.7130	42.9920	CA686		EU24	EUP5	ACC
			CA687		EU8	EUP12	ACC
			CA688			EUP12	
			CA672		EU10	EUP12	ACC
	Afyonkarahisar-26 Agust Natural Park Afyonkarahisar-Cumhuriyet village Çay Afyonkarahisar-Karamık lake Ağrı-Aşağı Yoldüzü Cuma River Ağrı-Çukuralan Village	Afyonkarahisar-26 Agust Natural Park38.7925Afyonkarahisar-Cumhuriyet village Çay38.5923Afyonkarahisar-Cumhuriyet village Qay38.5923Afyonkarahisar-Karamık lake38.4215Ağrı-Aşağı Yoldüzü Cuma River39.8181Ağrı-Çukuralan Village39.7130	Afyonkarahisar-26 Agust Natural Park38.792530.3816Afyonkarahisar-Cumhuriyet village Çay38.592330.9618Afyonkarahisar-Karamık lake38.421530.8869Ağrı-Aşağı Yoldüzü Cuma River39.818143.0892Ağrı-Çukuralan Village39.713042.9920	$ \begin{array}{ c c c c c c } \hline Afyonkarahisar-26 Agust Natural Park & 38.7925 & 30.3816 & CA1730 \\ \hline Park & 38.7925 & 30.3816 & CA1720 \\ \hline CA1721 & CA1721 & CA1722 & CA1723 & CA1723 & CA1723 & CA1723 & CA1724 & CA1724 & CA1724 & CA1724 & CA1724 & CA1709 & CA1714 & CA1712 & CA1710 & 30.8869 & CA1711 & CA1712 & CA1710 & CA1712 & CA1713 & CA1712 & CA1713 & CA1713 & CA679 & CA680 & CA681 & CA683 & CA683 & CA683 & CA683 & CA683 & CA684 & CA683 & CA685 & CA684 & CA685 & CA684 & CA685 & CA688 & CA$	$ \begin{array}{ c c c c c c c } \hline Afyonkarahisar-26 Agust Natural Park & 38.7925 & 30.3816 & CA1730 & & & \\ \hline Park & 38.7925 & 30.3816 & CA1730 & & & \\ \hline Afyonkarahisar-Cumhuriyet village \\ \hline Cay & 38.5923 & 30.9618 & CA1722 & & & \\ \hline CA1721 & & & & \\ \hline CA1722 & & & & \\ \hline CA1723 & & & & \\ \hline CA1724 & & & & \\ \hline CA1724 & & & & \\ \hline CA1724 & & & & \\ \hline CA1709 & & & & \\ \hline CA1710 & & & & \\ \hline CA1710 & & & & \\ \hline CA1710 & & & & \\ \hline CA1710 & & & & \\ \hline CA1710 & & & & \\ \hline CA1712 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA680 & & & & \\ \hline CA681 & & & & \\ \hline CA683 & & & & \\ \hline CA683 & & & & \\ \hline CA683 & & & & \\ \hline CA684 & & & & \\ \hline CA685 & & & & \\ \hline CA685 & & & & \\ \hline CA686 & & & & \\ \hline CA687 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline \end{array} $	$ \begin{array}{ c c c c c c c } \hline Afyonkarahisar-26 Agust Natural Park & 38.7925 & 30.3816 & CA1730 & & & & \\ \hline Park & 38.7925 & 30.3816 & CA1730 & & & & \\ \hline CA1720 & & & & & \\ \hline CA1721 & AT110 & & & \\ \hline CA1722 & & & & & \\ \hline CA1723 & AT41 & & & \\ \hline CA1724 & CA8 & & & \\ \hline CA1724 & CA8 & & & \\ \hline CA1709 & CA8 & & & \\ \hline CA1709 & CA8 & & & \\ \hline CA1710 & & & & \\ \hline CA1710 & & & & \\ \hline CA1711 & & & & & \\ \hline CA1712 & & & & \\ \hline CA1712 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA1713 & & & & \\ \hline CA680 & & & EU8 & \\ \hline CA681 & & & & \\ \hline CA683 & & & & \\ \hline CA683 & & & & \\ \hline CA683 & & & & \\ \hline CA684 & & & & \\ \hline CA683 & & & & \\ \hline CA684 & & & & \\ \hline CA684 & & & & \\ \hline CA685 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline CA688 & & & & \\ \hline \end{array} $	$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$

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Ağrı-Dambat village

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	Ağrı-Dambat village	39.6769	43.0228	CA678			EU24	EUP5	ACC	AJ310312	MHG6d
				CA658				ANT2		GQ902107	MHG6c
				CA659			EU24	EUP5	ACC	AJ310312	
	Ağrı-Doğu Beyazıt Bardaklı village	39.681	44.0655	CA660				EUP12		AJ310313	MHG6d
				CA661			EU24	EUP18	ACC	ACC	
				CA662				ANT2		GQ902107	MHG6c
				Hotz16808	A20	AN50	AT83	ANT25	GU812107	AJ313131	
	Akçapınar	37.1133	28.4656	Ploetner160.03	A20	AN50	AT83	ANT25	GU812107	AJ313131	MHG6c
				Ploetner161.03	A21	AN50	AT85	ANT25	GU812108	AJ313131	WINGOU
				Ploetner165.03	A20	AN50	AT83	ANT25	GU812107	AJ313131	
				CA1225				CAR12		GQ902088	MHG6a
	Aksaray Akin yillaga stream	38.4313	34.0291	CA1226				ANT1		AJ310337	MHG6c
Turkey	Aksaray-Akin vinage sucam			CA1227				ANT1		AJ310337	
Turkey				CA1228				ANT1		AJ310337	
				CA1212				ANT1		AJ310337	
				CA1213				ANT1		AJ310337	MHG6c
				CA1214				ANT1		AJ310337	
	Aksaray-Helvadere	38.1999	34.2103	CA1215				ANT1		AJ310337	
				CA1216				ANT1		AJ310337	
				CA1217				ANT1		AJ310337	
				CA1218				ANT1		AJ310337	
				CA1219				CAR12		GQ902088	MHG6a
				CA1220				ANT1		AJ310337	
	Aksaray-Melendez river	38.2988	34.2662	CA1221				ANT1		AJ310337	MUCG
				CA1222				ANT1		AJ310337	MILGOC
				CA1223				ANT1		AJ310337	

	Aksaray-Melendez river	38.2988	34.2662	CA1224				ANT1		AJ310337	MHG6c
				CA1205				ANT1		AJ310337	
	Alzerov Sultenhan	28 2414	22 5 1 1 9	CA1206			AT5	ANT1	ACC	AJ310337	MHC6a
	Aksaray-Sultannan	30.2414	55.5440	CA1207				ANT1		AJ310337	MHGGe
				CA1208				ANT1		AJ310337	
				CA1209				ANT1		AJ310337	
	Aksaray-Sultanhanı	38.2414	33.5448	CA1210				ANT1		AJ310337	MHG6c
				CA1211				ANT1		AJ310337	
				CBCAST4240		AN17		CAR17		AJ313132	MHG6a
				CBCA03184		AN27		ANT2		GQ902107	
			31.4546	CBCA03185		AN46		ANT21		GQ902114	MHG6c
				CBCA03186		AN27		ANT2		GQ902107	
Turkey				CBCA03187		AN17		CAR17		AJ313132	MHG6a
Turkey	Akşehir-Eber Lakes			CBCA03188		AN36		ANT11		GU812193	MHG6c
		38.4544		CBCA03189		AN46		ANT21		GQ902114	Mildoe
				CBCA03190		AN17		CAR17		AJ313132	MHG6a
				CA1715			CA8	CAR17	GU812089	AJ313132	Milloou
				CA1716				ANT2		GQ902107	MHG6c
				CA1717				CAR14		AJ313133	MHG6a
				CA1718				ANT2		GQ902107	MHG6c
				CA1719				ANT2		GQ902107	Mildoe
				MTHTCA07177		AN1		CAR1		AJ310314	
				MTHTCA07178		AN1		CAR1		AJ310314	
	Alanya	36.6031	32.0694	ZFMK40193		AN1		CAR1		AJ310314	MHG6a
				ZFMK40195	A4	AN1	CA20	CAR1	GU812091	AJ310314	
			-	ZFMK40198	A4	AN1	CA20	CAR1	GU812091	AJ310314	

				CA909				ANT1		AJ310337	
				CA910			AT33	ANT1	ACC	AJ310337	
	Amasya-Boğazköy Tersakan	40.7268	35.7702	CA911				ANT1		AJ310337	MHG6c
				CA912				ANT1		AJ310337	
				CA913				ANT1		AJ310337	
				CA902				ANT35		ACC	
	Amasya Doğantene nond	40.6104	35 5860	CA903			AT1	ANT1	GU812111	AJ310337	MHG6c
	Allasya-Dogantepe politi	40.0104	55.5809	CA904				ANT1		AJ310337	MHG6C
				CA905				ANT1		AJ310337	
			35 5751	CA914			AT1	ANT1	GU812111	AJ310337	
Amasya-Suluoava Yedik	Amasya-Suluoaya Vedikir dam	40 7786		CA915				ANT1		AJ310337	
	rinasya Salabava realkii aani	40.7780	55.5751	CA916				ANT1		AJ310337	
				CA917				ANT1		AJ310337	MHG6c
Turkey	Amasya-Yeşilırmak	40 6735	35 8528	CA907				ANT1		AJ310337	MHG6c
		40.0755	55.0520	CA908				ANT1		AJ310337	Milliot
			33 0453	KKOBCA06229		AN26		ANT1		AJ310337	
	Ankara	39 5793		KKOBCA06230		AN26		ANT1		AJ310337	MHG6c
	, interest	57.5775	55.0155	KKOBCA06231		AN26		ANT1		AJ310337	
				-	A24	AN26	AT1	ANT1	GU812111	AJ310337	
				BCA31291		CI1		CIW1		AJ313135	
	Antakya Samandağı	36.0742	35,9982	BCA31292		CI1		CIW1		AJ313135	MHG4
	i inimity i Summing	00107.12	0000002	BCA31293		CI1		CIW1		AJ313135	
				BCA31294		CI8		CIE1		GU812168	MHG5
				CBCASTO702		AN1		CAR1		AJ310314	
	Antalya	37,2119	30.9415	CBCASTO704		AN1		CAR1		AJ310314	MHG6a
	, interju	37.2119	30.9415	CBCASTO705		AN15		CAR15		GQ902085	
				CBCASTO706		AN1		CAR1		AJ310314	

Table 3.2. (c	ontinued).
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				CBC 407/3	AN1		CAR1		A 1310314	
				HKHSBKC A07111			CARIO		G0902084	
				HKHSBKCA07112	ANI		CARI		A 1310314	
				HKHSBKCA07112	ANI		CAR1		AJ310314	
				HKHSBKCA07114	ANI		CAP1		AJ310314	
				HKHSBKCA07115	ANI		CAR1		AJ310314	
				HKHSBKCA07116	ANI		CAP1		AJ310314	
				HKHSBKCA07110	ANI		CARI		AJ310314	
	Antalya	37 2119	30 9415	HKHSBKCA07117	ANI		CARS		G0002082	MHG6a
	Antaiya	57.2119	50.9115	HKHSBKCA07110	ANJ		CARJ		A 1210214	Milloou
				HKHSBKCA07120	ANI		CARI		AJ510514	
				HKHSBKCA0/120	AIN9		CAR9		GU812178	
				HKHSBKCA07121	AN6		CAR6		GU812176	
Turkey				HKHSBKCA07122	AN1		CAR1		AJ310314	
2				HKHSBKCA07123	AN10		CAR10		GQ902084	
				HKHSBKCA07124	AN1		CAR1		AJ310314	
				ZFMK45348	AN1		CAR1		AJ310314	
				ZFMK45349	AN1		CAR1		AJ310314	
				CA1433			CAR1		AJ310314	
				CA1434		CA20	CAR1	GU812091	AJ310314	
	Antalya-Anamur	36.0429	32.8078	CA1435			CAR1		AJ310314	MHG6a
				CA1436			CAR1		AJ310314	
				CA1437			CAR1		AJ310314	
				CA1862		CA17	CAR1	ACC	AJ310314	
	Antolya Kamar Kasmahağı	26 5054	20 5045	CA1863			CAR1		AJ310314	MUCGo
Aı	Antaiya-Kemer Kesmebogaz river	30.3954	30.5045	CA1864			CAR1		AJ310314	мнбба
				CA1865			CAR1		AJ310314	

	Antalya-Kemer Kesmeboğaz river	36.5954	30.5045	CA1866			CAR1		AJ310314	MHG6a
	Antalya-Gazipaşa	36.1687	32.4467	CA1441		CA18	CAR1	ACC	AJ310314	MHG6a
	Antoliya Kumluga	36.3168	20.2512	CA1870		CA20	CAR24	GU812091	ACC	MHG6a
Turkay	Antarya-Kumuca		30.2512	CA1871			ANT46		ACC	MHG6c
				CA732			ANT12		GU812194	
				CA733			ANT12		GU812194	
	Ardahan Göle Sarma Bridge	40 9087	12 5028	CA734		AT70	ANT12	ACC	GU812194	MHG6c
	Aldanan-Oole Senne Bridge	40.9087	42.5928	CA735			ANT12		GU812194	MHGGe
				CA736			ANT12		GU812194	-
				CA737			ANT12		GU812194	
	Ardahan-Kura River	41.1569		CA743			EUP12		AJ310313	MHG6c
			42.8728	CA744		AT70	ANT12	ACC	GU812194	
				CA745			ANT12		GU812194	
Turkey	Ardahan-Uzunova Village	41.0800	42.4991	CA739			EUP12		AJ310313	MHG6d
				CA740			ANT12		GU812194	MHG6c
				CA741			ANT14		GU812196	
	Ardahan-Uzunova Village	41.0800	42.4991	CA742			ANT14		GU812196	MHG6c
		41.1247	42.0668	CA746			ANT12		GU812194	MHG6c
	Artvin-Ardanue River			CA747			ANT12		GU812194	
				CA748			ANT12		GU812194	
				CA749			ANT12		GU812194	
		41.3649	41.6920	DC08241	AN28		ANT3		GU812185	MHG6c
				DC08242	AN39		ANT14		GU812196	
	Artvin Borçka			DC08243	AN26		ANT1		AJ310337	
				CA757		AT70	ANT50	ACC	ACC	
				CA759			ANT1		AJ310337	

	Artvin Borcka	41.3649	41.6920	CA760			ANT50		ACC	MHG6c
	Aitviii Bolçka			CA761		AT70	ANT50	ACC	ACC	
	Artvin-Hopa Kemalpaşa	41.4870		CA764		AT1	ANT1	GU812111	AJ310337	MHG6c
			41.5269	CA767		AT78	ANT14	ACC	GU812196	
				CA768			ANT1		AJ310337	
				ATDC08191	AN37		ANT12		GU812194	MHG6c
	Artvin Saveat	41 3098	42.4837	ATDC08192	AN37		ANT12		GU812194	
	Artvin Şavşat	41.5098		ATDC08193	AN37		ANT12		GU812194	
				ATDC08194	AN37		ANT12		GU812194	
	Artvin-Yolüstü Village	41.1613	42.0639	CA753			ANT12		GU812194	MHG6c
				CA754		AT70	ANT12	ACC	GU812194	MHG6c
Turker				CA755			EUP12		AJ310313	MHG6d
	Lake Avlan	36.5825	29.9482	HS07165	AN23		CER3		GQ902087	MHG6b
Turkey				HS07166	AN23		CER3		GQ902087	
				HS07167	AN20		CAR20		GQ902086	MHG6a
				HS07168	AN20		CAR20		GQ902086	
				HS07169	AN20		CAR20		GQ902086	
	Aydın-Azap lake	37.5844	27.447	CA1800			ANT21		GQ902114	MHG6c
				CA1801			ANT21		GQ902114	
	Avdın-Azan lake	37.5844	27.447	CA1802		AT110	ANT21	GU812104	GQ902114	MHG6c
	Ayum-Azap lake			CA1803		AT87	ANT25	ACC	AJ313131	
	Aydın-Bıyıkali pond	37.7716		CA1785			ANT21		GQ902114	MHG6c
			27.5741	CA1786			ANT21		GQ902114	
				CA1787		AT107	ANT21	ACC	GQ902114	
				CA1788			ANT21		GQ902114	
				CA1789			ANT21		GQ902114	

		37.5476	27.2371	CA1796	AT87	ANT25	ACC	AJ313131	
	Aydın-Menderes river			CA1797		ANT21		GQ902114	MHG6c
				CA1798	AT107	ANT21	ACC	GQ902114	
				CA1790		ANT21		GQ902114	-
				CA1791	AT87	ANT25	ACC	AJ313131	
	Avdın-Söke water channel	37 6607	27 3087	CA1792		ANT25		AJ313131	MHG6c
	Ayum-soke water channel	57.0007	27.3087	CA1793	AT110	ANT21	GU812104	GQ902114	WINGOU
				CA1794		ANT21		GQ902114	
				CA1795		ANT21		GQ902114	
		40.2968	27.6273	CA1617		ANT21		GQ902114	MHG6c
	Balıkesir-Erdek			CA1618	AT110	ANT21	GU812104	GQ902114	
				CA1619	AT59	ANT2	GU812109	GQ902107	
Turkey				CA1620		ANT2		GQ902107	
Turkey				CA1621		ANT10		GU812192	
	Balıkesir-İkizcetepeler dam	39.4807	27.9274	CA1601		ANT21		GQ902114	MHG6c
				CA1602	AT86	ANT28	ACC	GU812208	
				CA1603	AT108	ANT21	ACC	GQ902114	
				CA1605		ANT10		GU812192	
		40.2206	28.0461	CA1611		ANT21		GQ902114	MHG6c
	Balikesir-Manyas Kus Lake			CA1612	AT96	ANT29	ACC	GU812209	
	Dankesn-Ivianyas Kuş Lake	40.2290		CA1613	AT59	ANT2	GU812109	GQ902107	
				CA1615	AT60	ANT2	ACC	GQ902107	
	Balıkesir-Manyas Kuş Lake	40.2296	28.0461	CA1616		ANT10		GU812192	MHG6c
		39.5295	27.9091	CA1607	AT109	ANT21	ACC	GQ902114	MHG6c
	Balıkesir-Pamukçu stream			CA1608	AT91	ANT29	GU812101	GU812209	
				CA1609		ANT29		GU812209	

	Balıkesir-Pamukçu stream	39.5295	27.9091	CA1610		ANT29		GU812209	MHG6c
	Bartın-Potbaşı river	41.6087		CA981		ANT1		AJ310337	
			22 2727	CA982		ANT1		AJ310337	
			52.5727	CA984	AT12	ANT1	ACC	AJ310337	MIGOC
				CA985		ANT1		AJ310337	
				CA989		ANT1		AJ310337	-
			32.3267	CA990		ANT1		AJ310337	
	Bartın-Bartın stream	41.5721		CA991	AT1	ANT1	GU812111	AJ310337	MHG6c
				CA992		ANT1		AJ310337	
				CA993		ANT1		AJ310337	
	Batman-Dicle river	37.9177	41.0856	CA1355		EUP5		AJ310312	MHG6d MHG6d
				CA1356		EUP5		AJ310312	
Turkey				CA1357		EUP5		AJ310312	
Turkey				CA1358		EUP5		AJ310312	
				CA1359	EU36	EUP5	GU812122	AJ310312	
				CA1360		EUP5		AJ310312	
	Batman-Hasankeyf	37.7356	41.3022	CA1370		EUP5		AJ310312	
				CA1371		EUP5		AJ310312	
		38.0463	41.1834	CA1361	EU33	EUP5	ACC	AJ310312	MHG6d
				CA1362		EUP5		AJ310312	
	Batman-Silvan road Carikli village			CA1363		EUP5		AJ310312	
	Batman-Silvan road Çarikli village			CA1364		EUP5		AJ310312	
				CA1365		EUP5		AJ310312	
				CA1366		EUP5		AJ310312	
	Bayburt-Çamlık District	40 3009	40 2013	CA809		ANT37		ACC	MHG6c
		40.3007	+0.2013	CA810		ANT37		ACC	milliote

	Bayburt-Çamlık District	40.3009	40.2013	CA811				ANT1		AJ310337	
				CA812				ANT37		ACC	MHG6c
				CA813				ANT37		ACC	
				CA814				ANT1		AJ310337	
		40 1156	39.7526	CA802				ANT1		AJ310337	MHG6a
	Bayburt-Gökcedere pond			CA803				ANT1		AJ310337	
	Bayburt-Gokçedere pond	40.1150		CA804				ANT1		AJ310337	WINGOU
				CA805				ANT1		AJ310337	
			40.0626	CA797				ANT1		AJ310337	MHG6c
	Bayburt-Oruçbeyli pond	40.2316		CA798				ANT1		AJ310337	
				CA799				ANT1		AJ310337	
				CA800				ANT37		ACC	
Turkey				CA801				ANT1		AJ310337	
Turkey	Bayburt-Toki	40.2729	40.1343	CA808				ANT1		AJ310337	MHG6c
	Bayramdere	40.2025	28.5003	Hotz17410		AN27		ANT2		GQ902107	MHG6c
				Hotz17421		AN27		ANT2		GQ902107	
		37.6802	31.7180	MEFUCA42195		AN13		CAR13		GU812179	-
				MEFUCA42196		AN12		CAR12		GQ902088	
				MEFUCA42197		AN12		CAR12		GQ902088	
				MEFUCA42198		AN12		CAR12		GQ902088	
	Lake Beysehir			Hotz17310	A2	AN17	CA8	CAR17	GU812089	AJ313132	MHG6a
	Late Deyşenii			Hotz17313	A2	AN14	CA8	CAR14	GU812089	AJ313133	WIIIGoa
				ZFMK40195		AN11		CAR11		AJ310316	
				CA1450				CAR13		GU812179	
				CA1451			CA1	CAR12	GU812088	GQ902088	
				CA1452				CAR21		ACC	
				CA1453		CA5	CAR21	ACC	ACC		
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				CA1454		CA2	CAR22	ACC	ACC		
	Lake Beyşehir	37.6802	31.7180	CA1455			CAR12		GQ902088	MHG6a	
				CA1456		CA6	CAR23	ACC	ACC		
				CA1457		CA8	CAR17	GU812089	AJ313132		
		40.0646		CA1656			ANT13		GU812195		
	Bilecik-Çiğdemlik		30.3142	CA1657		AT1	ANT1	GU812111	AJ310337	MHG6c	
				CA1658		AT69	ANT2	ACC	GQ902107		
	Bilecik-İnhisar Sakarya river			CA1659		AT1	ANT43	GU812111	ACC		
		40.0469		CA1660			ANT2		GQ902107		
			30.4124	CA1661		AT67	ANT13	GU812110	GU812195	MHG6c	
				CA1662			ANT1		AJ310337		
Turkey				CA1663			ANT2		GQ902107		
Turkey				CA1664			ANT1		AJ310337		
				MAYGCA11303	AN27		ANT2		GQ902107	-	
				MAYGCA11304	AN26		ANT1		AJ310337		
				MAYGCA11305	AN26		ANT1		AJ310337		
	Bilecik Söğüt	39.7118	30.0070	MAYGCA11306	AN26		ANT1		AJ310337	MHG6c	
				CA1648			ANT1		AJ310337		
				CA1649		AT1	ANT43	GU812111	ACC		
				CA1651		AT68	ANT13	ACC	GU812195		
				CA570			EUP5		AJ310312		
				CA571		EU35	EUP5	ACC	AJ310312		
	Bingöl-Çeltiksuyu stream	38.8411	40.5654	CA572			EUP5		AJ310312	MHG6d	
				CA573			EUP5		AJ310312		
				CA574		EU16	EUP16	ACC	ACC		

				CA582		EUP5		AJ310312	
	Pingël Cohentee	20.0585	40 7076	CA583		EUP5		AJ310312	МНСеч
	Biligoi-Çobalitaşı	39.0385	40.7970	CA584		EUP5		AJ310312	MHOOd
CA584 EU CA585 EU33 EU CA585 EU33 EU CA575 EU33 EU CA576 EU33 EU CA576 EU33 EU CA576 EU33 EU CA576 EU33 EU CA576 EU33 EU CA576 EU33 EU CA576 EU33 EU CA578 EU33 EU CA579 EU34 EU CA586 EU36 EU CA587 EU EU CA587 EU EU	EUP5	ACC	AJ310312						
				CA575		EUP5		AJ310312	
				CA576		EUP5		AJ310312	
	Bingöl-Sarıçiçek Lake	38.8844	40.5890	CA577	EU36	EUP5	GU812122	AJ310312	MHG6d
				CA578		EUP5		AJ310312	
				CA579		EUP5		AJ310312	
Turker			40.7859	CA586	EU36	EUP5	GU812122	AJ310312	
		38.9184		CA587		EUP5		AJ310312	MHG6d
	Bingöl-Solhan Seref stream			CA588		EUP5		AJ310312	
				CA589		EUP5		AJ310312	
Turkey				CA590		EUP5		AJ310312	
	Bitlis-Ağaçköprü District	38.3328	42.0098	CA630	EU33	EUP5	ACC	AJ310312	MHG6d
				CA618	EU33	EUP5	ACC	AJ310312	
	Bitlis-Gürovmak Water Channel	38 5962	42 0243	CA619	EU17	EUP19	ACC	ACC	MHG6d
	Dittis-Guroymak water Chamier	50.5702	42.0245	CA620		EUP5		AJ310312	MILOOD
				CA621		EUP12		AJ310313	
				CA623	EU33	EUP5	ACC	AJ310312	
	Bitlis-Kemah Stream	38.4407	42.1447	CA624		EUP5		AJ310312	MHG6d
				CA625		EUP5		AJ310312	
				CA631	EU19	EUP5	ACC	AJ310312	MHG6d
	Bitlis-Tatvan Cağlavan	38 4748	42.3084	CA632	EU18	EUP15	ACC	ACC	
	Bitlis-Tatvan Çaglayan	38.4748		CA633		EUP5		AJ310312	
				CA634		EUP5		AI310312	

	Bitlis-Tatvan Çağlayan	38.4748	42.3084	CA635			EUP5		AJ310312	MHG6d
				CA1520			ANT1		AJ310337	
	Bolu-Gölköy	40.7116	31.5314	CA1521		AT1	ANT1	GU812111	AJ310337	MHG6c
				CA1522			ANT1		AJ310337	
				CA1525			ANT1		AJ310337	
	Bolu-Karamanlar lake	40.7619	31.5162	CA1526			ANT1		AJ310337	MHG6c
				CA1527		AT1	ANT1	GU812111	AJ310337	
	Bolu-Karamanlar lake	40.7619	31.5162	CA1528			ANT1		AJ310337	MHG6c
	Bucak		30.5393 -	MTAECA1568	AN1		CAR1		AJ310314	
				MTAECA1569	AN1		CAR1		AJ310314	
Turkey		37.3500		MTAECA1570	AN1		CAR1		AJ310314	
				MTAECA1571	AN1		CAR1		AJ310314	MHG6a
				MTAECA1572	AN14		CAR14		AJ313133	
				MTAECA1573	AN1		CAR1		AJ310314	
				MTAECA1574	AN17		CAR17		AJ313132	
				MTAECA1575	AN14		CAR14		AJ313133	
				MTAECA1576	AN1		CAR1		AJ310314	
				MTAECA1577	AN1		CAR1		AJ310314	
				CBCAST1517	AN1		CAR1		AJ310314	
	Lake Burdur	37.8371	30.3854	CBCAST1519	AN1		CAR1		AJ310314	MHG6a
				CBCAST1520	AN1		CAR1		AJ310314	
				CA1882			CAR1		AJ310314	
	Burdur-Çerçin dam	37.7603	30.4149	CA1883			CAR1		AJ310314	MHG6a
				CA1884			CAR1		AJ310314	<u> </u>
	Burdur-Karataş lake	37 3616	29 9869	CA1877		CA8	CAR17	GU812089	AJ313132	MHG6a
		37.3616	29.9869	CA1878			CAR17		AJ313132	MIIGOa

				CA1879		CAR17		AJ313132	MHG6a
	Burdur-Karataş lake	37.3616	29.9869	CA1880	AT103	ANT21	ACC	GQ902114	MHG6c
				CA1881	CA14	CAR1	ACC	AJ310314	MHG6c
				CA1872		ANT21		GQ902114	
	Purdur Voriali lako	27 5007	20.0562	CA1873		ANT21		GQ902114	MHC6a
	Burdur- i arişir take	37.3907	29.9302	CA1874		ANT21		GQ902114	MINGOC
				CA1876		ANT1		AJ310337	
Turkey	Bursa-Akçalar village			CA1641	AT98	ANT29	ACC	GU812209	
		40.1771	28.7453	CA1642	AT63	ANT52	ACC	ACC	MHG6c
				CA1643	AT67	ANT2	GU812110	GQ902107	
		40 1771	29 7452	CA1644		ANT10		GU812192	MUCC
	Buisa-Akçalal village	40.1771	20.7433	CA1646		ANT2		GQ902107	MHG0e
	Bursa-Boğaz	40.2821	28.4483	CA1622		ANT2		GQ902107	
				CA1623		ANT2		GQ902107	MHG6c
				CA1624	AT59	ANT2	GU812109	GQ902107	Willooc
				CA1626		ANT29		GU812209	
				CA1635	AT67	ANT52	GU812110	ACC	
				CA1636	AT96	ANT29	ACC	GU812209	
	Bursa-Gölyazı	40.1653	28.6795	CA1637		ANT29		GU812209	MHG6c
				CA1638		ANT29		GU812209	
				CA1640	AT59	ANT58	GU812109	ACC	
				CA1628		ANT10		GU812192	
	Bursa-Kemalpaşa			CA1629	AT110	ANT21	GU812104	GQ902114	
		40.0346	28.4104	CA1631		ANT52		ACC	MHG6c
				CA1632		ANT2		GQ902107	
				CA1634	AT110	ANT61	GU812104	ACC	

	Püvükaalmaaa	41.0020	28 6000	Hotz16627		AN27		ANT2		GQ902107	MHC6a
	Buyukçekinece	41.0020	28.0009	Hotz16628		AN27		ANT2		GQ902107	MHOOC
	Ceyhan	37 0880	35 8302	Hotz17303	CL4	CI1	CLW15	CIW1	GU812083	AJ313135	MHG4
	Ceynan	57.0889	55.8502	Hotz17305	CL2	CI1	CLW9	CIW1	GU812081	AJ313135	WI104
				CA1575				ANT2		GQ902107	
	Canakkale-Karacaören	40 1884	26 4330	CA1577				ANT2		GQ902107	MHG6c
	çanakkale-Karacaoren	40.1884	20.4550	CA1578				ANT2		GQ902107	MILOOC
Turkov				CA1580			AT61	ANT2	ACC	GQ902107	
			26.3852	CA1589				ANT2		GQ902107	
				CA1590			AT59	ANT2	GU812109	GQ902107	
	Çanakkale-Kepez	40.0895		CA1591				ANT2		GQ902107	MHG6c
				CA1592				ANT2		GQ902107	ļ
				CA1593				ANT10		GU812192	
Turkey	Çanakkale-Batakova Menderes river	39.9930	26.2078	CA1596				ANT2		GQ902107	
				CA1597				ANT2		GQ902107	MHG6c
				CA1598				ANT2		GQ902107	
				CA1599			AT59	ANT2	GU812109	GQ902107	
				CA1600				ANT2		GQ902107	
				CA1582			AT63	ANT52	ACC	ACC	
	Canakkale-Saricay	40 1393	26 4855	CA1584			AT64	ANT1	ACC	AJ310337	MHG6c
	Çunakkule Buriçûy	40.1575	20.4055	CA1585			AT67	ANT2	GU812110	GQ902107	MIIGOC
				CA1586				ANT2		GQ902107	
				CA1023				ANT1		AJ310337	
	Çankırı-Apsarı stream	40 7018	33 5393	CA1024			AT1	ANT1	GU812111	AJ310337	MHG6c
		10.7010	33.5393	CA1025				ANT1		AJ310337	MIIGOC
				CA1027				ANT1		AJ310337	

Table 3.2.	(continued).
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				CA1028				ANT1		AJ310337	
	Cankırı-Aşağıyanlar district	40 5540	33 5817	CA1029			AT21	ANT1	ACC	AJ310337	MHG6c
	Çankırı-Aşağıyanlar ülsület	+0.55+0	55.5617	CA1030				ANT1		AJ310337	WINGOU
				CA1031				ANT1		AJ310337	
				CA1032				ANT1		AJ310337	
	Cankuri-Terme river	40 4377	33 7/31	CA1033				ANT1		AJ310337	MHG6c
	Çalıkırı-Terme river	40.4377	55.7451	CA1034			AT29	ANT1	ACC	AJ310337	WINGOC
				CA1035				ANT1		AJ310337	
		40.7675		CA1018				ANT1		AJ310337	
Turkey	Çankırı-Yapraklı pond		33,7641	CA1019			AT19	ANT1	ACC	AJ310337	MHG6c
			55.7041	CA1020				ANT1		AJ310337	
				CA1021				ANT1		AJ310337	
	Çevlik District, Antakya	36 1230	35 0281	AMPH\SUR\202	CL3	CI1	CLW8	CIW1	GU812082	AJ313135	MHG4
Turkey		50.1257	35.7201	AMPH\SUR\203	CL1	CI1	CLW7	CIW1	GU812080	AJ313135	WI104
		40.3371	35.0628	CA1041			AT1	ANT1	GU812111	AJ310337	
	Çorum-Alaca stream			CA1042				ANT1		AJ310337	MHG6c
				CA1043				ANT1		AJ310337	
				CA1050				ANT1		AJ310337	
	Corum-Comar dam	40 5856	3/1 0008	CA1051				ANT1		AJ310337	MHG6c
	çorum-çomai tam	40.3850	54.9990	CA1052			AT1	ANT1	GU812111	AJ310337	WINGOC
				CA1054				ANT1		AJ310337	
_				CA1046			AT1	ANT1	GU812111	AJ310337	
	Çorum-Yakacık stream	40.6000	34.9117	CA1047				ANT1		AJ310337	MHG6c
				CA1049				ANT1		AJ310337	
	Çorum-Yılgınözü stream	40 4502	34 3789	CA1037				ANT1		AJ310337	MHG6c
		40.4502	34.3789	CA1038				ANT1		AJ310337	MIIGOC

Table 3.2.	(continued).
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	Corum-Vilginözü stream	40.4502	3/1 3789	CA1039		AT1	ANT1	GU812111	AJ310337	MHG6c
	Çorum- r ngmozu sucam	40.4302	54.5769	CA1040			ANT1		AJ310337	WINGOC
				IS48307	AN50		ANT25		AJ313131	
				IS48308	AN50		ANT25		AJ313131	MHG6c
Turkey	Dalaman	36 7138	28 7856	IS48309	AN17		CAR17		AJ313132	MHG6a
	Dataillait	50.7158	28.7850	IS48310	AN50		ANT25		AJ313131	MHG6c
				IS48311	AN17		CAR17		AJ313132	MHG6a
				IS48312	AN23		CER3		GQ902087	MHG6b
	Denizli-Derbent dam		28.8443 -	CA1814			ANT21		GQ902114	
		38 1532		CA1815		AT110	ANT21	GU812104	GQ902114	
Turkey		36.1332		CA1816			ANT21		GQ902114	
				CA1817			ANT21		GQ902114	MHG6c
				CA1804			ANT21		GQ902114	
Turkey	Denizli-Emmiler district	37 6264	29 2087	CA1805		AT110	ANT21	GU812104	GQ902114	
	Demzii-Limmer distret	37.0204	27.2007	CA1806			ANT21		GQ902114	
				CA1807			ANT21		GQ902114	MHG6c
				CA1809			ANT21		GQ902114	
	Denizli İncirlinmar park	37 7623	20 0072	CA1810			ANT21		GQ902114	
	Demzn-men npmar park	51.1025	27.0772	CA1811			ANT21		GQ902114	
				CA1812			ANT21		GQ902114	MHG6c
	Denizli-İncirlipınar park	37.7623	29.0972	CA1813		AT105	ANT21	ACC	GQ902114	MHG6c
				CA1820			ANT21		GQ902114	
				CA1821			ANT21		GQ902114	
	Denizli-Süleymanlı lake	38.0513	28.7708	CA1822		AT110	ANT21	GU812104	GQ902114	MHG6c
				CA1823			ANT21		GQ902114	
				CA1824			ANT21		GQ902114	

	Divarbakur Batman road	37 8081	40 4055	CA1345	EU36	EUP22	GU812122	ACC	MHG6d
	Diyarbakii-Daunan toad	57.0001	40.4033	CA1346	EU20	EUP5	ACC	AJ310312	MIIOou
				CA1339		EUP5		AJ310312	
				CA1340		EUP5		AJ310312	
	Diyarbakır-Devegeçidi	38.0576	40.0697 -	CA1341		EUP5		AJ310312	MHG6d
		50.0570		CA1342		EUP5		AJ310312	WINGOU
				CA1343	EU36	EUP24	GU812122	ACC	
				CA1344	EU33	EUP5	ACC	AJ310312	
			41.0217	CA1347		EUP5		AJ310312	
		37.8797		CA1348	EU36	EUP5	GU812122	AJ310312	
				CA1349		EUP5		AJ310312	MHG6d
Turkey	Diyarbakır-Dicle river			CA1350		EUP5		AJ310312	
				CA1352		EUP5		AJ310312	
				CA1353		EUP5		AJ310312	
				CA1354		EUP5		AJ310312	
				CA1331	EU36	EUP5	GU812122	AJ310312	
				CA1332		EUP5		AJ310312	
				CA1333		EUP5		AJ310312	
	Divarbakır-Göksu Cınar	37 6916	40 4473	CA1334		EUP5		AJ310312	MHG6d
	Diyarbaklı Ooksu Çıllar	57.0910	+0.++75	CA1335		EUP5		AJ310312	Millou
				CA1336		EUP5		AJ310312	
				CA1337		EUP5		AJ310312	
				CA1338		EUP5		AJ310312	<u> </u>
	Düzce-Asarsu			CA1530	AT1	ANT1	GU812111	AJ310337	MHG6c
		40.8259	31.1840	CA1531	AT66	ANT2	ACC	GQ902107	
				CA1532	AT17	ANT48	ACC	ACC	

	Düzce-Asarsu	40.8259	31 1840	CA1533				ANT1		AJ310337	MHG6c
	Duzte-Asarsu	40.8239	51.1640	CA1534				ANT1		AJ310337	WITCOC
				CA1536			AT17	ANT48	ACC	ACC	
	Düzce-Efteni lake	40 7573	31.0404	CA1537			AT57	ANT1	ACC	AJ310337	MHG6c
	Duzee-Litem lake	40.7575	51.0404	CA1538			AT57	ANT41	ACC	ACC	WINGOU
				CA1539				ANT1		AJ310337	
	Edime			MACA22179		RE1		RE1		AJ310338	
		41.6857	26.4932	MACA22180		RE1		RE1		AJ310338	MHG1
				MACA22181		RE1		RE1		AJ310338	WINGT
				MACA22182		RE1		RE1		AJ310338	
				MACA22183		AN35				GU812192	MHG6c
Turkey	Edremit	39 6119	27.0151	Ploetner179.03	A22	AN27	AT59	ANT2	GU812109	GQ902107	MHG6c
	Editerint	39.0119	27.0151	Ploetner180.03	A23	AN27	AT67	ANT2	GU812110	GQ902107	WITCOC
				CBCAST3233		AN27		ANT2		GQ902107	MHG6c
				CBCAST3234		AN1		CAR1		AJ310314	
				CBCAST3238		AN14		CAR14		AJ313133	
	I ake Eğirdir	38 1393	30 7588	CBCAST3239		AN14		CAR14		AJ313133	
	Lake Eghtin	56.1575	50.7500	CA1890				CAR14		AJ313133	MHG6a
				CA1891			CA8	CAR17	GU812089	AJ313132	
				CA1892			CA20	CAR1	GU812091	AJ310314	
				CA1893				CAR14		AJ313133	
-	Elazığ - Birvan stream	38.7260	38.8375	CA543				EUP5		AJ310312	MHG6d
	Elazığ - Cip			CA535				EUP5		AJ310312	
		29 6910	39.0706	CA536			EU33	EUP5	ACC	AJ310312	MHC64
		36.0019		CA537				EUP5		AJ310312	wiriGou
				CA538				EUP5		AJ310312	

	Elazığ - Cip	38.6819	39.0706	CA539		EUP5		AJ310312	MHG6d
	Flaziă Sivrice	38 4665	30 2753	CA545		EUP5		AJ310312	MHG6d
	Elazig - Sivilce	38.4005	39.2733	CA546		EUP5		AJ310312	MIIGOd
				CA520		ANT1		AJ310337	
		39.6559		CA521		ANT1		AJ310337	MHG6c
	Erzincan - Türkmenoğlu village		39.4996	CA522		ANT1		AJ310337	WINGOU
				CA523	AT1	ANT1	GU812111	AJ310337	
				CA524		CIE2		GU812169	MHG5
Turkey	Erzincan - Ekşisu			CA525		ANT1		AJ310337	MHG6c
				CA526	AT1	ANT1	GU812111	AJ310337	Mildoe
		39.7328	39.6179	CA527	CLW15	CIW8	GU812083	ACC	MHG4
				CA528		ANT1		AJ310337	MHG6c
				CA529		ANT1		AJ310337	MIIGOC
	Erzincan-Sakaltutan	39.8832	39.1954	CA530	AT10	ANT1	ACC	AJ310337	
				CA531	AT31	ANT1	ACC	AJ310337	MHG6c
				CA532		ANT1		AJ310337	
				CA533		ANT1		AJ310337	
				CA695		ANT12		GU812194	
				CA696	AT70	ANT12	ACC	GU812194	
				CA697	AT7	ANT1	ACC	AJ310337	MHG6c
	Erzurum-Ilıca	39.8196	41.1521	CA699		ANT12		GU812194	
				CA700		ANT12		GU812194	
_				CA701	EU7	EUP12	ACC	AJ310313	MHG6d
				CA702		ANT12		GU812194	MHG6c
	Erzurum-Pasinler	39 9611	41 4090	CA704	EU36	EUP5	GU812122	AJ310312	MHG6d
		39.9611	41.4090	CA705		ANT12		GU812194	MHG6c

				CA706	AT70	ANT12	ACC	GU812194	MHG6a
	Erzurum-Pasinler	39.9611	41.4090	CA707		ANT12		GU812194	MHG0e
				CA708	EU7	EUP12	ACC	AJ310313	MHG6d
				CA709		ANT12		GU812194	
				CA710		ANT12		GU812194	MHG6c
	Erzurum-Soğuk Çermik	39.9899	41.3052	CA711	AT70	ANT12	ACC	GU812194	
				CA712		EUP12		AJ310313	MUCGA
				CA713	EU7	EUP12	ACC	AJ310313	MILGOU
				CA689		ANT12		GU812194	
				CA690	AT70	ANT12	ACC	GU812194	
	Erzurum-Teke stream	39.8196	41.1521	CA691		ANT12		GU812194	MHG6c
				CA692		ANT12		GU812194	
Turkey				CA693		ANT12		GU812194	
				CA714	EU9	EUP12	ACC	AJ310313	MHG6d
	Erzurum-Yerlisu Village	40.0425	41.1833	CA715	EU36	EUP5	GU812122	AJ310312	Millood
				CA716	AT8	ANT1	ACC	AJ310337	MHG6c
				CA1672	AT41	ANT53	ACC	ACC	MHG6c
				CA1673	CA8	CAR17	GU812089	AJ313132	MHG6a
	Eskişehir-Alpu road	39.7825	30.6877	CA1674	AT67	ANT2	GU812110	GQ902107	
				CA1675		ANT43		ACC	MHG6c
				CA1676		ANT2		GQ902107	
				CA1677		ANT1		AJ310337	
	Eskisehir-Porsuk river 1	39,7744	30.4511	CA1678	AT1	ANT1	GU812111	AJ310337	7 MHG6c 7 MHG6c 2 MHG6a 17 MHG6c 17 MHG6c 17 MHG6c 17 MHG6c 17 MHG6c 17 MHG6c 17 MHG6c 17 MHG6c
	Eskişehir-Porsuk river 1 3		CA1679 CA1680	CA1679		ANT10		GU812192	
				CA1680		ANT2		GQ902107	

	Eskigshir Porsuk river 2	20 7120	20 4275	CA1683			AT110	ANT29	GU812104	GU812209 MHG6c GU812192	MHC6a
	Eskişenii-Forsuk iivei 2	39.7139 39.7082 38.2296 36.6288 38.6666	30.4275	CA1684				ANT10		GU812192	MHGoc
				CA1666			AT110	ANT21	GU812104	GQ902114	
				CA1667			AT67	ANT2	GU812110	GQ902107	
	Eskişehir-Sarısungur	39.7082	30.5804	CA1668				CAR17		AJ313132	MHG6c
				CA1669			AT1	ANT43	GU812111	ACC	
				CA1670			AT67	ANT2	GU812110	GQ902107	
				TBOECA68170		AN26		ANT1		AJ310337	
				TBOECA68171		AN26		ANT1		AJ310337	
				TBOECA68172		AN26		ANT1		AJ310337	MHG6c
	Eşmekaya	38.2296	33.4876	TBOECA68173		AN26		ANT1		AJ310337	
				TBOECA68174		AN27		ANT2		GQ902107	
Turkey				TBOECA68175		AN17		CAR17		AJ313132	MHG69
Turkey				TBOECA68176		AN17		CAR17		AJ313132	MIIG0a
				RCA48213		AN50		ANT25		AJ313131	MHG6c
	Fethive	36 6288	29 1196	RCA48214	A10	AN23	CE8	CER3	GU812097	GQ902087	
	reunye	50.0200	29.1190	RCA48215	A11	AN25	CE9	CER5	GU812098	GU812184	MHG6b
				RCA48216		AN22		CER2		GU812182	
				Ploetner197.03	A26	AN35	AT56	ANT10	GU812113	GU812192	
	Foca	38 6666	26 7645	Ploetner198.03	A17	AN46	AT110	ANT21	GU812104	GQ902114	MHG6c
	roşu	50.0000	20.7015	Ploetner199.03	A27	AN35	AT52	ANT10	GU812114	GU812192	Milloot
_				Ploetner205.03	A16	AN46	AT106	ANT21	GU812103	GQ902114	
				OBMB27295		AN55		EUP12		AJ310313	MHG6d
	Gazianten	37,0892	37,1735	OBMB27296		CI9		CIE2		GU812169	MHG5
	Gaziantep 37.0892	37.1735	OBMB27297		AN59		EUP5		AJ310312	MHG6d	
				OBMB27298		AN59		EUP5		AJ310312	

				CA1968		CIW	3	GU812163	
				CA1969		CIE	3	GU812170	MHC5
	Gaziantep-Balıkalan village	37.2923	36.8877	CA1970		CIE	2	GU812169	MINUS
				CA1971		CIE	2	GU812169	
				CA1972		CLW15 CIW	3 GU812083	GU812163	MHG4
	Cagionton Kalaurt	27 1260	26.0400	CA1973		CIE	5	ACC	MUC5
	Gaziantep-Kakurt	57.1500	30.9400	CA1974		CIEI	0	ACC	MHG5
				CA1965		CIE	2	GU812169	
	Gaziantep-Sakçagözü	37.1803	36.9285	CA1966		CIE	2	GU812169	MHG5
				CA1967		CIE	3	GU812170	
				CBCAST2021	AN46	ANT	21	GQ902114	
				CBCAST2022	AN46	ANT	21	GQ902114	MUCGo
Turkey				CBCAST2023	AN46	ANT	21	GQ902114	MHOOC
				CBCAST2024	AN46	ANT	21	GQ902114	
	Gemis	37 7902	29 8708	CBCAST2025	AN1	CAR	1	AJ310314	MHG6a
	Gennş	51.1902	29.0700	CBCAST2026	AN46	ANT	21	GQ902114	
				CBCAST2027	AN46	ANT	21	GQ902114	_
				CBCAST2028	AN46	ANT	21	GQ902114	MHG6c
				CBCAST2029	AN46	ANT	21	GQ902114	
				CBCAST2031	AN46	ANT	21	GQ902114	
				CA835		ANT	6	ACC	
				CA836		ANT	1	AJ310337	_
	Giresun-Batlama stream	40.9067	38.3547	CA837		ANT	1	AJ310337	MHG6c
				CA838		ANT	1	AJ310337	
				CA839		ANT	1	A I310337	

				CA854			ANT1		AJ310337		
	Giresun-Bulancak Domuz stream	40.9454	38.1673	CA855			ANT1		AJ310337	MHG6c	
				CA856			ANT1		AJ310337		
				CA849			ANT1		AJ310337		
	Circoun Ecnivo Colivoro Stroom	40.0450	28 7774	CA850			ANT1		AJ310337	MHC6a	
	Gresuit-Espiye Genvera Stream	40.9439	30.7224	CA852			ANT1		AJ310337	WITCOC	
				CA853			ANT1		AJ310337		
				CA844			ANT1		AJ310337		
	Giresun-Kesap Karabulduk stream	40.8988	38.5270	CA847			ANT1		AJ310337	MHG6c	
Turkov				CA848			ANT1		AJ310337		
	Giresun-Yavşan stream	40.8291	38.4584	CA842			ANT1		AJ310337	MHG6c	
	Gödet Dam 37.1076 33.2918 OYCA70204 AN1 CAR1 OYCA70206 AN1 CAR1<		AJ310314								
	Gödet Dam	37.1076	33.2918	OYCA70206	AN1		CAR1		AJ310314	MHG6a	
Turkey				OYCA70207	AN1		CAR1		AJ310314		
				OYCA70208	AN8		CAR8		GU812177		
				CA1900			CAR1		AJ310314		
	Gödet Dam	37.1076	33.2918	CA1901			CAR1		AJ310314	MH6a	
				CA1902		CA15	CAR1	ACC	AJ310314		
				CA1903		CA15	CAR8	ACC	GU812177		
				CCYZCA20153	AN46		ANT21		GQ902114		
				CCYZCA20154	AN46		ANT21		GQ902114		
	Gökpınar Dam	37.7851	29.1306	CCYZCA20155	AN46		ANT21		GQ902114	MHG6c	
					CCYZCA20156	AN46		ANT21		GQ902114]
				CCYZCA20157	AN46		ANT21		GQ902114	<u>] </u>	
	Gümüşhane-Akbaba pond 4	40 1783	39 6521	CA817			ANT1		AJ310337	MHG6c	
		+0.1705	57.0521	CA818			ANT1		AJ310337	MINGOC	

				CA819				ANT1		AJ310337	
	Gümüşhane-Akbaba pond	40.1783	39.6521	CA820				ANT1		AJ310337	MHG6c
				CA821				ANT1		AJ310337	
				CA822				ANT1		AJ310337	
				CA823				ANT1		AJ310337	
	Gümüşhane-Kelkit river	40.1226	39.3737	CA824				ANT1		AJ310337	MHG6c
				CA825				ANT1		AJ310337	
				CA826				ANT1		AJ310337	
				CBCA2047		AN17		CAR17		AJ313132	
				CBCA2048		AN14		CAR14		AJ313133	
Turkey	I ake Isikli	38 2350	29 9605	CBCA2049		AN18		CAR18		GQ902097	MHG6a
	Luke Işikli	50.2550	29.9005	CBCA2050		AN17		CAR17		AJ313132	Millooa
				CBCA2051		AN14		CAR14		AJ313133	
				CBCA2052		AN17		CAR17		AJ313132	
				CA663			AT82	ANT2	HM356084	GQ902107	
	lğdır- Tuzluca Aras River	40.0356	43 6790	CA664			AT82	ANT2	HM356084	GQ902107	MHG6c
	igun Tuzhuou Thus fervor	10.0550	15.0790	CA665				ANT2		GQ902107	initiooe
				CA666				ANT2		GQ902107	
				CA667			AT82	ANT2	HM356084	GQ902107	
				CA668				ANT2		GQ902107	
	Iğdır- Water Channel	39.9513	44.0419	CA669				ANT2		GQ902107	MHG6c
				CA670				ANT2		GQ902107	
				CA671				ANT2		GQ902107	
	İstanbul	41.1340	28.7533	-	A39	AN26	EU27	ANT1	GU812125	AJ310337	MHG6d
-	Istanbul Halkali	41.0432	28.7823	ZISP.6537-Sp.46		RE15				GU812157	MHG1
	Hatay-Deniz	36.0727	35.9508	CA1949			CLW15	CIW11	GU812083	ACC	MHG4

Table 3.2. (continued).
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				CA1950				CIW1		AJ313135	MHG4
				CA1951			CLE16	CIE1	ACC	GU812168	MHG5
	Hatay-Deniz	36.0727	35.9508	CA1952				CIW1		AJ313135	MHG4
				CA1953			CLW5	CIW1	ACC	AJ313135	MIII04
				CA1954			CLE12	CIE2	ACC	GU812169	MHG5
	Hatay Gölbaşı	36 1615	36 1778	CA1955			CLW15	CIW8	GU812083	ACC	MHG4
	Tratay-Goldaşı	50.4045	50.4778	CA1956				CIW3		GU812163	WI104
				CA1403				CIW1		AJ313135	MHG4
				CA1404			CLE17	CIE9	ACC	ACC	MHG5
	Hatay-Hassa	36.8449	36.6504	CA1405				CIW1		AJ313135	
				CA1406			CLW15	CIW11	GU812083	ACC	MHG4
				CA1407				CIW1		AJ313135	
Turkey	Hatay İskandarun Erzin	36 8887	36 1370	CA1416			CLW2	CIW2	ACC	GU812162	MHG4
Turkey		30.8887	50.1570	CA1417			CLW15	CIW10	GU812083	ACC	WI104
				CA1409				CIW1		AJ313135	MHG4
				CA1410			CLE11	CIE2	ACC	GU812169	MHG5
	Hatay-İskenderun Sarışeki	36 6653	36 2157	CA1411			CLW11	CIW3	ACC	GU812163	MHG4
	Tratay-iskenderun Sariseki	50.0055	50.2157	CA1413				CIW1		AJ313135	WI104
				CA1414				CIE2		GU812169	MHG5
				CA1415			CLW1	CIW2	ACC	GU812162	MHG4
				OBMB31341		CI9		CIE2		GU812169	
				OBMB31342		CI9		CIE2		GU812169	
	Hatay Kırıkhan	36 4973	36 4523	OBMB31343	CL6	CI10	CLE4	CIE3	GU812085	GU812170	MHG5
	Thung Kirkhan	50.7775	50.4525	OBMB31344		CI8		CIE1		GU812168	
				OBMB31345		CI10		CIE3		GU812170	
				OBMB31346		CI1		CIW1		AJ313135	MHG4

Table 3.2. (continued).	
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	XX . XZ 11	26.4072	0.6.4500			1) mro (
	Hatay Kırıkhan	36.4973	36.4523	OBMB31347		CI3		CIW3		GU812163	MHG4
				OBMB31336		CI3		CIW3		GU812163	
	Hatay Reyhanlı	36 2379	36 5689	OBMB31338		CI1		CIW1		AJ313135	MHG4
		50.2577	50.5007	OBMB31339		CI3		CIW3		GU812163	
				OBMB31340	(CI10		CIE3		GU812170	MHG5
				OBMB31348	(CI10		CIE3		GU812170	MHG5
	Hatay Tahtalı Dam	36.8515	36.6861	OBMB31349		CI9		CIE2		GU812169	WI105
				OBMB31350		CI1		CIW1		AJ313135	MHG4
				CA1446			AT45	ANT2	ACC	GQ902107	MHG6c
	Isparta Bağıllı	38 1586	31.0825	CA1447			CA20	CAR14	GU812091	AJ313133	MHG6a
Turkey	Isparta-Dagini	36.1360	51.0625	CA1448				ANT2		GQ902107	MHG6c
				CA1449				CAR1		AJ310314	MHG6a
				CA1885			CA13	CAR7	ACC	GQ902112	
Turkey				CA1886				CAR7		GQ902112	
	Isparta-Sevinçbey district	37.8748	30.7760	CA1887				CAR7		GQ902112	MHG6a
				CA1888				CAR7		GQ902112	
				CA1889				CAR1		AJ310314	
				OK42203	А	AN12		CAR12		GQ902088	
	İvriz	27 4409	24 1705	OK42220	А	AN12		CAR12		GQ902088	MHC6a
	IVIIZ	37.4408	54.1705	OK42221	А	AN12		CAR12		GQ902088	WINGOa
				OK42222	А	AN12		CAR12		GQ902088	
				CA1748				ANT21		GQ902114	
				CA1749			AT110	ANT59	GU812104	ACC	
	İzmir-Çandarlı Bakırçay	38.9556	27.0100	CA1750			AT97	ANT29	ACC	GU812209	MHG6c
				CA1751			AT63	ANT52	ACC	ACC	
				CA1752			AT110	ANT21	GU812104	GQ902114	

Table 3.2. (continued).

	1				1					
				CA1759			ANT21		GQ902114	
	İzmir-Gediz river	38 6500	27 0274	CA1761		AT59	ANT2	GU812109	GQ902107	MHG6c
		50.0577	21.0214	CA1762		AT104	ANT21	ACC	GQ902114	MIIGOC
				CA1763			ANT21		GQ902114	
				CA1754			ANT21		GQ902114	
	İzmir-Menemen	38.6955	26.9989	CA1755		AT97	ANT29	ACC	GU812209	MHG6c
				CA1756		AT106	ANT21	GU812103	GQ902114	
				CA1557		AT1	ANT42	GU812111	ACC	
				CA1558		AT41	ANT53	ACC	ACC	
	İzmit-Çayırköy lake	40.8134	29.9910	CA1559			ANT1		AJ310337	MHG6c
Turkay				CA1560		AT94	ANT29	ACC	GU812209	
				CA1561		AT4	ANT1	ACC	AJ310337	
				CA1552			ANT1		AJ310337	
Turkey	İzmit Sanança laka	40.7180	30 1542	CA1553		AT5	ANT1	ACC	AJ310337	MHG6c
	izint-Sapanca lake	40.7100	50.1542	CA1554		AT65	ANT2	ACC	GQ902107	WINGOC
				CA1555			ANT2		GQ902107	
				CA1250		CLW15	CIW3	GU812083	GU812163	MHG4
	Kahramanmaras Göksun	37 0070	36 5211	CA1251			CIW3		GU812163	WIN04
	Kainainainaraş-Ooksun	31.9910	50.5211	CA1252			CIE3		GU812170	MHC5
				CA1253		CLE2	CIE3	ACC	GU812170	WI105
				CA1256		CLE7	CIE3	ACC	GU812170	MHG5
				CA1257		AT2	ANT1	ACC	AJ310337	MHG6c
	Kahramanmaras-Elhistan Gölnınar	38 1050	37.0636	CA1258			CIE3		GU812170	MHG5
	Kumamanaaş-Eloistan Oolpinar	50.1757	57.0050	CA1259			ANT1		AJ310337	MHG6c
				CA1260			CIE3		GU812170	MHG5
				CA1261			CIE3		AJ310337 AJ310337 AJ310337 GQ902107 GQ902107 GU812163 GU812163 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170 GU812170	141105

	Kahramanmaras Elhistan Gölnmar	38 1050	37.0636	CA1262			ANT1		AJ310337	MHG6a
	Kalıramannaraş-Eloistan Gölpinar	36.1939	37.0050	CA1263			ANT1		AJ310337	WINGOU
				CA1264			CIE2		GU812169	
				CA1265		CLE15	CIE2	ACC	GU812169	MHG5
	Kahramanmaraş-Kumaşır lake	37.5102	36.8983	CA1266			CIE3		GU812170	
				CA1268		CLW12	CIW2	ACC	GU812162	MHG4
			CA1269 CI	CIE2		GU812169	MHG5			
				CA965		AT16	ANT44	ACC	ACC	
				CA966			ANT33		ACC	
	Karabük-Cemaller Soğanlı river	41.1313	32.6816	CA967		AT15	ANT1	ACC	AJ310337	MHG6c
				CA968			ANT1		AJ310337	
				CA969			ANT1		AJ310337	
Turkey				CA974			ANT1		AJ310337	
Turkey				CA975			ANT1		AJ310337	
	Karabük-Safranbolu Araç river	41.2165	32.7331	CA976			ANT1		AJ310337	MHG6c
				CA977		AT14	ANT1	ACC	AJ310337	
				CA978	CA978 AT1 ANT31 GU8	GU812111	ACC			
				CBCAST327	AN1		CAR1		AJ310314	
				CBCAST328	AN1		CAR1		AJ310314	
				CBCAST329	AN1		CAR1		AJ310314	
				CBCAST3210	AN1		CAR1		AJ310314	
	Karacaören Dams I-II	37.4031	30.8703	CBCAST3211	AN1		CAR1		AJ310314	MHG6a
				CBCAST3212	AN1		CAR1		AJ310314	
				CBCAST3213	AN1		CAR1		AJ310314	
				CBCA3241	AN1		CAR1		AJ310314	
				CBCA3242	AN2		CAR2		GU812173	1

Table 3.2.	(continued).
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				OBBKCA32103	AN1	7	CAR17		AJ313132	
				OBBKCA32104	AN		CAR1		AJ310314	
	Varaaaäran Dama I II	37.4031	20.0702	OBBKCA32105	ANI		CAR1		AJ310314	MHC6a
	Karacaoten Danis I-II		30.8703	OBBKCA32106	ANI		CAR1		AJ310314	MINGOa
				OBBKCA32107	ANI		CAR1		AJ310314	
				OBBKCA32108	ANI		CAR1		AJ310314	
	Karaman-Belpınarı			CA1909		CA19	CAR1	ACC	AJ310314	
				CA1910			CAR1		AJ310314	
		36.8333	32.5803	CA1911			CAR1		AJ310314	MHG6a
				CA1912			CAR1		AJ310314	
·				CA1913			CAR1		AJ310314	
		37.2047	33.4035	CA1905			CAR8		GU812177	MHG6a
Turkey	Karaman-Vesildere			CA1906		CA15	CAR8	ACC	GU812177	
Turkey				CA1907		CA15	CAR1	ACC	AJ310314	
				CA1908			CAR1		AJ310314	
				CA725			EUP12		AJ310313	MUC64
				CA726			EUP12		AJ310313	
	Kars-Arnacay Akcalar	40 7735	43 2952	CA727			EUP12		AJ310313	
	Kuis / Iipuçuy / Kçulul	40.7755	43.2752	CA728			EUP12		AJ310313	Millood
				CA729			EUP13		ACC	
				CA730			EUP17		ACC	
				CA717			ANT2		GQ902107	
	Kars-Digor Pazarcık	40 5189	43 2690	CA718			ANT2		GQ902107	MHG6c
	Muis Digor i uzuroix	+0.5107	+3.2070	CA719			ANT2		GQ902107	
				CA720			ANT2		GQ902107	
	Kars-Selim	40.4702	42.7904	CA721			EUP17		ACC	MHG6d

				CA722				EUP12		AJ310313	
	Kars-Selim	40.4702	42.7904	CA723				EUP12		AJ310313	MHG6d
				CA724				EUP12		AJ310313	
			1	CA959				ANT1		AJ310337	
	Kastamonu-Araç River	41 2380	33 3774	CA960			AT1	ANT1	GU812111	AJ310337	MHG6c
		41.2307	55.5224	CA961				ANT1		AJ310337	WINGOU
				CA963				ANT1		AJ310337	
	Kastamonu-Beyler dam	41 6903	22 9110	CA952				ANT1		AJ310337	MHG6c
	Kastanionu-Deyler dani	41.0703	55.0117	CA953				ANT1		AJ310337	WINGOU
	Kastamonu-Beyler dam	41.6903	33.8119	CA954			AT44	ANT2	ACC	GQ902107	MHG6c
	Kastanionu-Deyler dani			CA955			AT57	ANT1	ACC	AJ310337	MINGOL
	Kastamonu-Karaçomak dam	41.2852	33.7372	CA939				ANT1		AJ310337	MHG6c
Turkey				CA940			AT20	ANT1	ACC	AJ310337	
Turkey				CA941			AT39	ANT44	ACC	ACC	
				CA942			AT1	ANT49	GU812111	ACC	
				CA943				ANT1		AJ310337	
				CA945			AT43	ANT2	ACC	GQ902107	
				CA946			AT21	ANT1	ACC	AJ310337	
	Kastamonu-Taşlık Dam	41.4018	33.6963	CA947				ANT1		AJ310337	MHG6c
				CA948				ANT1		AJ310337	
				CA949				ANT1		AJ310337	
				CA07217	A9	AN21	CE5	CER1	GU812096	GU812181	
				CA07218		AN23		CER3		GQ902087	MHG6b
	Kaş	36.2769	29.6839	CA07219		AN23		CER3		GQ902087	
				ZFMK21047	A7	AN23	CE6	CER3	GU812094	GQ902087	
				ZFMK29521		AN23		CER3		GQ902087	

Table 3.2. (continued).
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				DCA38313	AN26		ANT1		AJ310337	
			35.4229	DCA38314	AN29		ANT4		GU812186	
	Kayseri	38.7565		DCA38315	AN26		ANT1		AJ310337	MHG6c
				DCA38316	AN26		ANT1		AJ310337	
				DCA38317	AN30		ANT5		GU812187	
				CA1151		AT13	ANT1	ACC	AJ310337	
	Kayseri-Engir lake			CA1152			ANT4		GU812186	
		38.8111	35.5915	CA1154			ANT1		AJ310337	MHG6c
				CA1155			ANT1		AJ310337	
				CA1156			ANT1		AJ310337	
		38.6618	36.104	CA1468		AT1	ANT1	GU812111	AJ310337	
	Kayseri-Pınarbaşı			CA1469			ANT1		AJ310337	MHG6c
Turkov				CA1471			ANT1		AJ310337	
Turkey	Kayseri-Pinarhasi	38 6618	36 104	CA1472			ANT1		AJ310337	MHG6c
	Kaysen-i marbaşı	38.0018	50.104	CA1473			ANT1		AJ310337	
				CA1157			CAR12		GQ902088	MHG6a
				CA1158			ANT1		AJ310337	_
				CA1159			ANT1		AJ310337	
	Kayseri Sultansazlığı	38 3807	35 3657	CA1160			ANT1		AJ310337	
	Kaysen-Sunansazingi	30.3097	55.5057	CA1161			ANT1		AJ310337	MHG6c
				CA1162			ANT1		AJ310337	
				CA1163			ANT1		AJ310337	
				CA1164			ANT1		AJ310337	
				CA1165			CAR12		GQ902088	MHG6a
	Kayseri-Yahyalı	38.1348	35.3641	CA1166			ANT2		GQ902107	MHG6c
				CA1167			ANT2		GQ902107	

			35.3641	CA1168			ANT40		ACC	
	Kayseri-Yahyalı	38.1348		CA1169			ANT1		AJ310337	MHG6c
				CA1170			ANT40		ACC	
				CA1171		AT2	ANT1	ACC	AJ310337	
				CA1172		AT26	ANT4	ACC	GU812186	
	Kayseri-Zamantı Water			CA1173			ANT2		GQ902107	
				CA1174			ANT2		GQ902107	
		38.0345	35.5407	CA1175		CLW3	CIW3	ACC	GU812163	MHG4
				CA1176			ANT1		AJ310337	
Turkey				CA1177			ANT1		AJ310337	MHG6c
				CA1178			ANT2		GQ902107	MIROOC
				CA1179			ANT1		AJ310337	
	Kemer	37.4627	30.1118	MHSACA15137	AN46		ANT21		GQ902114	MHG6c
Turkey				MHSACA15138	AN1		CAR1		AJ310314	MHG6a
				MHSACA15139	AN7		CAR7		GQ902112	
				MHSACA15140	AN1		CAR1		AJ310314	
				MHSACA15141	AN1		CAR1		AJ310314	
	Kemer	37 4627	30 1118	MHSACA15142	AN1		CAR1		AJ310314	MHG6a
		57.4027	50.1110	MHSACA15143	AN1		CAR1		AJ310314	Millooa
				CA1238			ANT1		AJ310337	
	Kırıkkale-Hasandede Kızılırmak	38 7406	33 4879	CA1239			ANT1		AJ310337	MHG6c
	Kinkkale-Hasandede Kizininak	30.7400	33.4879	CA1242			ANT1		AJ310337	MHGoc
				CA1243			ANT1		AJ310337	
				CA1244			ANT1		AJ310337	MHG6c
	Kırıkkale-Yahşiyan Kızılırmak	39.8861	33.4146	CA1245			ANT1		AJ310337	
				CA1246			ANT1		AJ310337	

Table 3.2.	(continued).
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	Kırıkkale-Yahşiyan Kızılırmak	39.8861	33.4146	CA1248			ANT1		AJ310337	MHG6c
				CA1488	RE1				AJ310338	
				CA1489	RE1				AJ310338	
	Kurklarali Eriklica straam	41 7576	27 1914	CA1490	RE16				ACC	MHG1
	Kirklaren-Errkitee sueam	41.7570	27.1014	CA1491	RE1				AJ310338	MIIOI
				CA1492	RE1				AJ310338	
				CA1493	RE1				AJ310338	
				CA1494	RE1					
				CA1495	RE1					MHG1
	Kırklareli-İnece stream	41.6836	27.0731	CA1496	RE1					
				CA1497		AT56	ANT10	GU812113	GU812192	MHG6c
				CA1498	RE1					MHG1
Turkey	Kırklareli-Karakoç pond	41.7817	27.2171	CA1499	RE1				AJ310338	
Turkey				CA1500	RE1				AJ310338	MHG1
				CA1501	RE1				AJ310338	MHG1
			27 2627	CA1484			ANT10		GU812192	MHG6c
	Kırklareli-Sevtandere	41 7104		CA1485	RE1				AJ310338	MHG1
	Kirklaren Şeyandere	41.7104	21.2021	CA1486	RE1				AJ310338	
				CA1487		AT53	ANT10	ACC	GU812192	MHG6c
	Kusehir-Cuyun	39 3168	34 1238	CA1141			ANT4		GU812186	MHG6c
	ixiişenii çuvin	57.5100	54.1250	CA1142		AT1	ANT1	GU812111	AJ310337	Mildoe
	Kusehir-Cuyun	39 3168	34 1238	CA1144			ANT1		AJ310337	MHG6c
		57.5100	37.5100 34.1230	CA1145		AT26	ANT4	ACC	GU812186	Mildoe
				CA1128			ANT1		AJ310337	MHG6c
	Kırşehir-Güneykent	39.0988	34.1600	CA1129		AT1	ANT1	GU812111	AJ310337	
				CA1130		AT26	ANT4	ACC	GU812186	

	Kırşehir-Güneykent	39.0988	34.1600	CA1131				ANT1		AJ310337	MHG6c
				CA1146			AT1	ANT1	GU812111	AJ310337	
	Kırşehir-Kışlapınar	39.2262	34.1338	CA1147				ANT1		AJ310337	MHG6c
				CA1149				ANT1		AJ310337	
				CA1135			AT1	ANT1	GU812111	AJ310337	
	Kırsehir-Sıddıklı	39 1007	33 9073	CA1136				ANT1		AJ310337	MHG6c
		59.1007	55.7075	CA1137				ANT1		AJ310337	milliote
				CA1138				ANT1		AJ310337	
				Hotz18003		AN38		ANT13		GU812195	
Turkey	Kıvıköv	41.7038	28.0514	Hotz18004		AN38		ANT13		GU812195	MHG6c
				Hotz18005		AN38		ANT13		GU812195	
				Hotz18006		AN35		ANT10		GU812192	
	Kilis	36 7641	37.2540	OB79321		CI9		CIE2		GU812169	MHG5
Tunkey				OB79322		CI10		CIE3		GU812170	
				OB79323		AN59		EUP5		AJ310312	MHG6d
	i i i i i i i i i i i i i i i i i i i	50.7011		OB79324		CI1		CIW1		AJ313135	MHG4
				OB79325		AN55		EUP1		AJ310313	MHG6d
				OB79326		AN55		EUP1		AJ310313	Milloou
				OBMB79327		CI10		CIE3		GU812170	MHG5
				OBMB79328		AN59		EUP5		AJ310312	MHG6d
	Kilis-Gazianten road	36.8106	37 3095	OBMB79329		CI10		CIE3		GU812170	
		50.0100	57.5075	OBMB79330		CI10		CIE3		GU812170	MHG5
				OBMB79331		CI11		CIE4		GU812171	
				OBMB79335	CL5	CI10	CLE3	CIE3	GU812084	GU812170	
	Kilis-Polateli	36.7853	37.0588	CA1395			CLE2	CIE7	ACC	ACC	MHG5
			37.0588	CA1396			CLE12	CIE2	ACC	GU812169	

				CA1397		CLE6	CIE3	ACC	GU812170		
				CA1398			CIE2		GU812169	MHG5	
	Kilis-Polateli	36.7853	37.0588	CA1399			CIE3		GU812170		
				CA1400		CLW15	CIW2	GU812083	GU812162	MHG4	
				CA1401			CIE3		GU812170	MHG5	
	Kilis-Töreli Çapalı road		37.3414	CA1390		CLW6	CIW1	ACC	AJ313135	MHG4	
		26.8540		CA1391			CIW1		AJ313135	MIN04	
		30.0349		CA1393		EU13	EUP1	ACC	AJ310313	MHG6d	
				CA1394		CLE2	CIE3	ACC	GU812170	MHG5	
	Kilis Ücgöz	36 80/15	37.0868	CA1957			CIE2		GU812169	MHG5	
	Kins-Oçgoz	50.0045		CA1959			EUP1		AJ310313	MHG6d	
			36.9328	CA1960			CIE3		GU812170	MHG6d	
Tradition				CA1961		CLE5	CIE6	ACC	ACC		
Turkey	Kilis-Üçpınar	36.8669		CA1962			CIE2		GU812169		
				CA1963			CIE2		GU812169		
				CA1964			CIW1		AJ313135	MHG4	
				CA1463			CAR1		AJ310314	MHG6a	
				CA1464			CAR1		AJ310314		
	Konya-Mehmetali dam	37.2124	32.6354	CA1465		CA19	CAR1	ACC	AJ310314		
				CA1466		CA3	CAR12	ACC	GQ902088]	
				CA1467		AT42	ANT2	ACC	GQ902107	MHG6c	
				VDBKCA07125	AN1		CAR1		AJ310314	MHG6a	
				VDBKCA07126	AN46		ANT21		GQ902114	MHG6c	
	Korkuteli	36.9918	29.5279	VDBKCA07127	AN1		CAR1		AJ310314	MHG6a	
				VDBKCA07128	AN7		CAR7		GQ902112		
				VDBKCA07129	AN17		CAR17		AJ313132		

	Koslavtali	26.0019	20,5270	VDBKCA07130	AN1	7	CAR17		AJ313132	MUCK
	Korkuteli	36.9918	29.5279	VDBKCA07131	AN1		CAR1		AJ310314	мнбба
				CBCAST3214	AN1		CAR1		AJ310314	MHG6a
				CBCAST3215	AN2	7	ANT2		GQ902107	MHG6c
				CBCAST3216	AN1	Ļ	CAR14		AJ313133	
				OAOBBKCA1593	AN1		CAR1		AJ310314	MHG6a
				OAOBBKCA1594	AN1		CAR1		AJ310314	Milloou
	Kovada	37.6325	30.8641	OAOBBKCA1596	AN1		CAR1		AJ310314	
				OAOBBKCA1597	AN1		CAR1		AJ310314	
				OAOBBKCA1598	AN3		CAR3		GU812174	
				OAOBBKCA1599	AN1		CAR1		AJ310314	MHG6a
				OAOBBKCA1593101	AN1		CAR1		AJ310314	
Turkov				OAOBBKCA1593102	AN1		CAR1		AJ310314	
Turkey	Kütahya-Enne dam	39.4688	29.8657	CA1685		AT110	ANT61	GU812104	ACC	MHG6c
				CA1686		AT49	ANT10	ACC	GU812192	
				CA1687		AT110	ANT21	GU812104	GQ902114	
				CA1688			ANT61		ACC	
				CA1697		AT110	ANT21	GU812104	GQ902114	
				CA1698		AT45	ANT2	ACC	GQ902107	
	Kütahya-Koçak stream	39.3090	29.9753	CA1699		AT110	ANT61	GU812104	ACC	MHG6c
				CA1700			ANT21		GQ902114	
				CA1701			ANT21		GQ902114	1
				CA1705		AT54	ANT55	ACC	ACC	
	Kütahva-Porsuk dam	39 5740	30.0918	CA1706		AT41	ANT2	ACC	GQ902107	MHG6c
	Kütahya-Porsuk dam	37.3740	50.0918	CA1707		AT56	ANT10	GU812113	GU812192	WITCOC
				CA1708		AT110	ANT21	GU812104	GQ902114	1

			29.9342	CA1690		AT50	ANT10	ACC	GU812192	
				CA1691		AT111	ANT21	ACC	GQ902114	
	Kütahya-Söğütözü stream	39.3374		CA1692			ANT21		GQ902114	MHG6c
				CA1693			ANT61		ACC	-
				CA1694		AT41	ANT2	ACC	GQ902107	
	Malatya	38 1276	38.3505	SO44285	AN59		EUP5		AJ310312	MHG6d
	watatya	30.4270		SO44286	AN59		EUP5		AJ310312	
				CA552		EU33	EUP5	ACC	AJ310312	MHG6d
				CA553			EUP5		AJ310312	MIIGou
Turkay	Malatva-Elemendik lake	38 3229	38 1551	CA554			ANT4		GU812186	MHG6c
	Malatya-Elementik lake	50.5227	56.1551	CA555			EUP5		AJ310312	MHG6d
				CA556			EUP5		AJ310312	Millood
				CA557		AT25	ANT4	ACC	GU812186	MHG6c
Turkey	Malatya-Karakaya dam	38.4882	38.3499	CA547			EUP5		AJ310312	MHG6d
				CA548		AT1		GU812111		MHG6c
				CA549			ANT4		GU812186	
				CA550			EUP5		AJ310312	MHG6d
				CA551		EU33	EUP5	ACC	AJ310312	
				CA558		AT1	ANT1	GU812111	AJ310337	
	Malatya-near Sultansuvu dam	38 3121	38 0/3/	CA559		AT26	ANT4	ACC	GU812186	MHG6c
	Malatya near Sutansuya dam	50.5121	50.0454	CA560			ANT4		GU812186	
				CA561		EU36	EUP5	GU812122	AJ310312	MHG6d
				CA562			ANT4		GU812186	MHG6c
	Malatya-Oluklu district	38 2406	37 9964	CA563			EUP5		AJ310312	MHG6d
	Malatya-Oluklu district	56.2400	57.7704	CA564			ANT4		GU812186	MHG6c
				CA565			ANT4		GU812186	MINGOC

Table 3.2. (continued).
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	Malatya-Oluklu district	38.2406	37.9964	CA566			EUP5		AJ310312	MHG6d
				OA45274	AN49		ANT24		GU812205	
				OA45275	AN46		ANT21		GQ902114	
				OA45276	AN46		ANT21		GQ902114	
	Manica	38 6110	27 1183	OA45277	AN53		ANT28		GU812208	MHG6c
	livianisa	56.0110	27.4405	OA45278	AN46		ANT21		GQ902114	WINGOU
				OA45279	AN46		ANT21		GQ902114	
				OA45280	AN46		ANT21		GQ902114	
				OA45281	AN46		ANT21		GQ902114	
	Manisa	38.6110	27.4483	OA45282	AN46		ANT21		GQ902114	MHG6c
		38.6021		CA1764		AT91	ANT29	GU812101	GU812209	
	Manisa City Forest		27.3867	CA1765		AT109	ANT21	ACC	GQ902114	MHG6c
Turkey				CA1766			ANT21		GQ902114	
Turkey				CA1767			ANT21		GQ902114	
				CA1768			ANT21		GQ902114	
				CA1780			ANT21		GQ902114	
				CA1781			ANT21		GQ902114	
	Manisa-Köseler pond	38.8460	27.2006	CA1782			ANT21		GQ902114	MHG6c
				CA1783			ANT21		GQ902114]
				CA1784		AT110	ANT60	GU812104	ACC	
				CA1774			ANT21		GQ902114	
				CA1775			ANT21		GQ902114	
	Manisa-Örselli pond	38.8698	27.2620	CA1776		AT100	ANT21	ACC	GQ902114	MHG6c
				CA1777		AT51	ANT57	ACC	ACC	
				CA1778		AT52	ANT10	GU812114	GU812192	
	Manisa-Siyekli pond	38.7886	27.2559	CA1769		AT110	ANT21	GU812104	GQ902114	MHG6c

				CA1770		AT91	ANT29	GU812101	GU812209	
	Manisa Sivekli pond	38 7886	27 2550	CA1771			ANT29		GU812209	MHG6c
	Manisa-Siyekii polid	56.7880	21.2337	CA1772			ANT21		GQ902114	WINGOC
				CA1773			ANT21		GQ902114	
	Mardin-Bülbül village	37.3219	40.8364	CA1319		EU33	EUP5	ACC	AJ310312	MHG6d
				CA1313		EU36	EUP5	GU812122	AJ310312	
	Mardin-Kızıltepe	37.1970	40.2767	CA1314			EUP5		AJ310312	MHG6d
				CA1315			EUP5		AJ310312	
				CA1326			EUP5		AJ310312	
	Mardin-Mazıdağ	37 4622	40.6169	CA1327		EU36	EUP22	GU812122	ACC	MHG6d
	Marum-Maziuag	57.4022	40.0107	CA1328			EUP5		AJ310312	MIROOd
				CA1329		EU36	EUP5	GU812122	AJ310312	
Turkey	Mardin-Mazıdağ	37.4622	40.6169	CA1330			EUP5		AJ310312	MHG6d
Turkey	Mardin Same road		40.9552	CA1320		EU28	EUP5	ACC	AJ310312	MHG6d
		27.4406		CA1321			EUP5		AJ310312	
				CA1322			EUP5		AJ310312	
	Wardin-Savur Toad	57.4400	40.0555	CA1323			EUP5		AJ310312	
				CA1324			EUP5		AJ310312	
				CA1325			EUP5		AJ310312	
				AKCA48209	AN50		ANT25		AJ313131	
				AKCA48210	AN51		ANT26		GU812206	
_	Marmaris	36.8467	28.2879	AKCA48211	AN51		ANT26		GU812206	MHG6c
			[AKCA48212	AN52		ANT27		GU812207	
				Hotz16812	AN51		ANT26		GU812206	
	Mersin	36 7969	34 6019	SOCA33224	CI3		CIW3		GU812163	MHG4
		30.7909	54.0017	SOCA33225	CI3		CIW3		GU812163	

				SOCA33226	 CI3		CIW3		GU812163	
	Mersin	36.7969	34.6019	SOCA33227	CI3		CIW3		GU812163	MHG4
				SOCA33228	CI1		CIW1		AJ313135	
				CA1429			CIW8		ACC	
	Mersin-Erdemli	36.5930	34.2873	CA1430		CLW4	CIW3	ACC	GU812163	MHG4
				CA1431		CLW4	CIW3	ACC	GU812163	
	Mersin-Mut Bice stream	36.5896	33.2850	CA1915			CAR1		AJ310314	MHG6a
				CA1916		CA16	CAR1	GU812092	AJ310314	
				CA1917			CAR1		AJ310314	
	Mersin-Mut Bucakışla stream	36.6313	33.3675	CA1918			CAR1		AJ310314	MHG6a
				CA1919			CAR1		AJ310314	
Turker				CA1921			CAR1		AJ310314	
	Mersin-Yenice	36.9673	35.0305	CA1424			CIW3		GU812163	
Turkey				CA1425			CIW3		GU812163	MHG4
				CA1426			CIW1		AJ313135	
	Marcin Vanica	36.9673	35.0305	CA1427		CLW15	CIW3	GU812083	GU812163	MHG4
	With shi- I chiec			CA1428		CLW15	CIW1	GU812083	AJ313135	WI104
				CA1845		CA11	CAR17	ACC	AJ313132	MHG6a
				CA1846		CE7	CER3	ACC	GQ902087	MHG6b
	Muŭla-Dalaman Tersakan	36 7794	28 8252	CA1847			CAR17		AJ313132	MHG6a
	Wugia-Dalaman Tersakan	50.7774	20.0252	CA1848			CAR17		AJ313132	WIIG0a
				CA1849			ANT25		AJ313131	MHG6a
				CA1850		AT83	ANT25	GU812107	AJ313131	WINGOU
				CA1851		CA7	CAR20	ACC	GQ902086	MHG6a
	Muğla-Girdev plateau 3	36.7007	29.6509	CA1852			CAR17		AJ313132	MIIGoa
				CA1853		CE4	CER3	ACC	GQ902087	MHG6b

				CA1854	CA13	CAR1	ACC	AJ310314	
				CA1855		CAR20		GQ902086	
	Mažla Cindan alatam	26 7007	20 6500	CA1856	CA12	CAR17	ACC	AJ313132	MUCC
	Mugia-Ondev plateau	30.7007	29.0309	CA1857		CAR20		GQ902086	MHG0a
				CA1858		CAR17		AJ313132	
				CA1859		CAR20		GQ902086	
				CA1840	CA11	CAR17	ACC	AJ313132	MHG6a
				CA1841	CE3	CER1	ACC	GU812181	MHG6b
	Muğla-Köyceğiz Zaferler village	36.9718	28.6299	CA1842		ANT25		AJ313131	
				CA1843	AT84	ANT62	ACC	ACC	MHG6c MHG6c
				CA1844		ANT25		AJ313131	
	Muğla-Ula Nannan stream	37.0129	28.5105	CA1836	AT83	ANT25	GU812107	AJ313131	
Turkov				CA1837	AT88	ANT26	ACC	GU812206	
Turkey				CA1838		ANT26		GU812206	
				CA1839		ANT26		GU812206	
		27 2750	28.0916	CA1826		ANT21		GQ902114	MHG6c
	Muğla Vatağan Dinsiz stream			CA1827		ANT21		GQ902114	
	Mugia-Tatagan Dipsiz sucam	57.5750		CA1828	AT101	ANT21	ACC	GQ902114	
				CA1829	AT87	ANT25	ACC	AJ313131	
				CA1830	AT87	ANT63	ACC	ACC	
				CA1831	AT102	ANT21	ACC	GQ902114	
	Muğla-Yemişendere	37.2516	28.5810	CA1832		ANT21		GQ902114	MHG6c
_				CA1833		ANT21		GQ902114	-
				CA1834		ANT63		ACC	
	Muş-Çizmeburnu	38 7159	41.6205	CA604	EU24	EUP5	ACC	AJ310312	MHG6d
		38./139		CA605		EUP14		ACC	

			41.6205	CA606			EU32	EUP5	ACC	AJ310312	
				CA607				EUP5		AJ310312	
	Muş-Çizmeburnu	38.7159		CA608				EUP5		AJ310312	MHG6d
				CA609				EUP5		AJ310312	-
				CA610				EUP5		AJ310312	
				CA600			AT81	ANT2	ACC	GQ902107	MHG6c
	Mus Murat Piwar	28.0612	41 5115	CA601				EUP5		AJ310312	
	Muş-Mulat Kivel	38.9013	41.5115	CA602				EUP15		ACC	MHG6d
				CA603				EUP15		ACC	
	Muş-Tigem Drenaj Channel			CA596				EUP5		AJ310312	MHG6d
		38.8017	41.4987	CA597			AT81	ANT2	ACC	GQ902107	MHG6c
				CA598				EUP5		AJ310312	MHG6d
Turkov				CA599			EU31	EUP5	ACC	AJ310312	MHOOd
Turkey		38.7862		CA611			EU32	EUP5	ACC	AJ310312	
	Muş-Tigem Water Channel		41.5969	CA612				EUP5		AJ310312	MHG6c
				CA613				EUP5		AJ310312	
				CA615				EUP5		AJ310312	
				CA616				EUP5		AJ310312	
				OBMBCA46364	CL8	CI9	CLE10	CIE2	GU812087	GU812169	MHG5
				OBMBCA46366		CI10		CIE3		GU812170	WIII05
				OBMBCA46367		CI3		CIW3		GU812163	MHG4
	Narlı	37.3322	37.0425	OBMBCA46369		CI10		CIE3		GU812170	MHG5
				OBMBCA46370		CI10		CIE3		GU812170	MIIOS
				OBMBCA46371		CI6		CIW6		GU812166	MHG4
				OBMBCA46372		CI12		CIE5		GU812172	MHG5
	National Observatory	36.8293	30.3407	VD07132		AN1		CAR1		AJ310314	MHG6a

				VD07134	AN1		CAR1		AJ310314	
	National Observatory	36.8293	30.3407	VD07135	AN1		CAR1		AJ310314	MHG6a
				VD07136	AN1		CAR1		AJ310314	
	Nevsehir-Avanos Kuzulırmak	38 7174	3/ 8575	CA1234		AT26	ANT4	ACC	GU812186	MHG6c
	Nevşenii-Avanos Kizininiak	56.7174	51.0575	CA1235		AT41	ANT2	ACC	GQ902107	mildoe
	Nevsehir-Nar Ilica river	38 6383	34 7089	CA1229			ANT1		AJ310337	MHG6c
	ivevşenii-ivar inca fiver	50.0505	34.7089	CA1230		AT1	ANT1	GU812111	AJ310337	
	Nevsehir-Gülsehir Kızılırmak	38 7564	34 6513	CA1236		AT41	ANT2	ACC	GQ902107	MHG6c
	Nevşenii-Guişenii Kizininiak	36.7304	54.0515	CA1237			ANT5		GU812187	MHG6c
				CA1231		AT1	ANT1	GU812111	AJ310337	
	Nevşehir-Ürgüp	38.5741	34.9157	CA1232			ANT4		GU812186	MHG6c
				CA1233			ANT4		GU812186	
Turkey	Niğde-Çamardı	37.9608	34.3572	CA1181		AT1	ANT1	GU812111	AJ310337	MHG6c
Turkey				CA1182			ANT1		AJ310337	
				CA1183			ANT1		AJ310337	
				CA1185			ANT1		AJ310337	
				CA1186			ANT1		AJ310337	
				CA1187			ANT1		AJ310337	
				CA1195			CAR12		GQ902088	MHG69
				CA1196			CAR12		GQ902088	MIIG0a
				CA1197		AT5	ANT1	ACC	AJ310337	MHG6c
	Niğde-Cifflik	38 1779	34 4629	CA1198			CIE3		GU812170	MHG5
	Nigue-Çinnik	38.1779	34.4629	CA1199			ANT1		AJ310337	
				CA1200			ANT1		AJ310337	MHG6c
				CA1201			ANT1		AJ310337	MINOUC
				CA1202			ANT1		AJ310337	1

			1		1	1					
	Niğde-Ciftlik	38,1779	34 4629	CA1203				ANT1		AJ310337	MHG6c
Turkey		50.1779	51.1025	CA1204				ANT1		AJ310337	
	Niğde-15km to Ulukışla	37.6195	34.5366	CA1193				ANT1		AJ310337	MHG6c
	Ordu-Civil Stream	40.9733	37.9030	CA858				ANT1		AJ310337	MHG6c
				CA859				ANT1		AJ310337	
				CA860				ANT1		AJ310337	
				CA861				ANT46		ACC	
				CA862			AT1	ANT1	GU812111	AJ310337	
	Ordu-Fatsa	41.0753	37.4413	CA871				ANT1		AJ310337	MHG6c
				CA872				ANT1		AJ310337	
				CA873				ANT1		AJ310337	
				CA874			AT1	ANT1	GU812111	AJ310337	
				CA875				ANT1		AJ310337	
	Ordu-Perşembe	41.0251	37.8032	CA867				ANT1		AJ310337	MHG6c
				CA868			AT1	ANT1	GU812111	AJ310337	
				CA869				ANT1		AJ310337	
				CA870			AT9	ANT1	ACC	AJ310337	
	Osmaniye-Aslantaş dam	37.2769	36.2744	CA1933			CLW14	CIW9	ACC	ACC	MHG4
				CA1934				CIW3		GU812163	
				CA1935			CLE14	CIE8	ACC	ACC	MHG5
				CA1936				CIW3		GU812163	MHG4
				CA1937			CLE1	CIE3	ACC	GU812170	MHG5
	Osmaniye-Boru river	37.1780	36.4856	CA1943			CLE8	CIE8	ACC	ACC	MHG5
				CA1944			CLE12	CIE2	ACC	GU812169	
				CA1945				CIE8		ACC	
	Özbaşı	37.6178	27.4327	Hotz17510	A17	AN46	AT110	ANT21	GU812104	GQ902114	MHG6c
				Hotz17511	A17	AN46	AT110	ANT21	GU812104	GQ902114	

Table 3.2. (continued).
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	Özhası	37 6178	27 4327	Hotz17716	A 1 7	AN46	AT110	ANT21	CU812104	C0002114	MHG6c
Turkey	Omerli	41 1003	29.4009	H0tz17710	AI/	AN21	AIII0	ANTC	00812104	GQ902114	MHG6c
		11.1005	29.1009	ODMDCA46254		CIO		AN10		GU812188	Milloot
	Pazarcık	37.2819	37.1249	OBMBCA46354		CI9		CIE2		GU812169	MHG5
				OBMBCA46355		C19		CIE2		GU812169	
				OBMBCA46356	CL7	CI9	CLE9	CIE2	GU812086	GU812169	
				OBMBCA46358		CI10		CIE3		GU812170	
	Rize	41.0110	40.5792	BAYDCA53253		AN26		ANT1		AJ310337	MHG6c
				AA53287		AN26		ANT1		AJ310337	
				AA53288		AN26		ANT1		AJ310337	
				CA771			AT1	ANT39	GU812111	ACC	
	Rize-Çiftekavak	41.0283	40.4858	CA779				ANT1		AJ310337	MHG6c
				CA780				ANT1		AJ310337	
	Rize-Hemşin Stream	41.1554	40.9012	CA769			AT71	ANT2	ACC	GQ902107	MHG6c
	Samsun	41.3621	36.2169	YCDC55244		AN26		ANT1		AJ310337	MHG6c
				YCDC55245		AN26		ANT1		AJ310337	
				YCDC55246		AN26		ANT1		AJ310337	
				YCDC55247		AN26		ANT1		AJ310337	
	Samsun-Bafra	41.5167	36.0158	CA890				ANT1		AJ310337	MHG6c
				CA891				ANT1		AJ310337	
				CA892			AT1	ANT1	GU812111	AJ310337	
				CA893				ANT1		AJ310337	
	Samsun-Kavak	41.1567	36.0927	CA897				ANT1		AJ310337	MHG6c
				CA898				ANT1		AJ310337	
				CA899			AT28	ANT1	ACC	AJ310337	
				CA900				ANT1		AJ310337	
	Samsun-19 Mayıs	41.4674	36.0993	CA884				ANT1		AJ310337	MHG6c
Table 3.2.	(continued).										
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				CA886		AT1	ANT1	GU812111	AJ310337	
	Samsun-19 Mayıs	41.4674	36.0993	CA887		AT1	ANT1	GU812111	AJ310337	MHG6c
				CA888			ANT1		AJ310337	
				CA878			ANT1		AJ310337	
	Samsun-Tekkeköv	41 2126	36 4790	CA879			ANT1		AJ310337	MHG6c
	Samsun-Tekkeköy	41.2120	50.4770	CA880			ANT1		AJ310337	WINGOU
				CA881		AT11	ANT1		AJ310337	
				CBCAST421	AN27		ANT2		GQ902107	MHG6c
				SGCA42199	AN27		ANT2		GQ902107	Mildoe
				SGCA42201	AN12		CAR12		GQ902088	
	Seydişehir	37.4557	31.8156	SGCA42202	AN17		CAR17		AJ313132	MHG6a
Turkey				CA1458		CA4	CAR12	ACC	GQ902088	WIIG0a
				CA1459		CA1	CAR12	GU812088	GQ902088	
				CA1460			ANT1		AJ310337	MHG6c
	Seydişehir	37.4557	31.8156	CA1461		CA6	CAR1	ACC	AJ310314	MHG6a
			41 7806	CA1372		EU33	EUP5	ACC	AJ310312	MHG6d
	Siirt-Baswirt river	37 9726		CA1373			EUP5		AJ310312	
	Shit-Daşyult Hver	51.5120	41.7000	CA1374			EUP5		AJ310312	
				CA1375			EUP5		AJ310312	
				CA1378			EUP5		AJ310312	
				CA1379		EU29	EUP15	ACC	ACC	
_	Siirt-Kezer river	37.9610	41.8573	CA1380		EU30	EUP5	ACC	AJ310312	MHG6d
				CA1383			EUP5		AJ310312]
				CA1384			EUP15		ACC	1
	Silifico	36 3130	33 9595	OBMB33257	AN1		CAR1		AJ310314	MHG62
	Sinike	36.3130	36.3130 33.9595	OBMB33258	AN4		CAR4		GU812175	MIIGUa

Table 3.2.	(continued).
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				OBMB33259		AN1		CAR1		AJ310314	MHG6a	
				OBMB33260		AN1		CAR1		AJ310314	Millooa	
				OBMB33261		CI1		CIW1		AJ313135	MHG4	
				OBMB33263		AN1		CAR1		AJ310314	MHG6a	
				OBMB33264		AN1		CAR1		AJ310314	WIIG0a	
				OBMB33265		CI3		CIW3		GU812163	MHG4	
				OBMB33266		AN1		CAR1		AJ310314		
	C11:4	26 2120	22.0505	OBMB33267		AN1		CAR1		AJ310314		
	Sillike	50.5150	55.9595	OBMB33268		AN1		CAR1		AJ310314		
				OBMB33269		AN1		CAR1		AJ310314		
				OBMB33270		AN1		CAR1		AJ310314	MHG6a	
				OBMB33271		AN1		CAR1		AJ310314		
Tradess				OBMB33272		AN1		CAR1		AJ310314		
Turkey				OBMB33273		AN1		CAR1		AJ310314		
				Hotz19394	A5	AN1	CA16	CAR1	GU812092	AJ310314		
				Hotz19400	A5	AN1	CA16	CAR1	GU812092	AJ310314		
	Sinon Avanally	41.0492	24 7740	CA925				ANT1		AJ310337	MUCGa	
	Smop-Ayancik	41.9485	54.7749	CA926				ANT1		AJ310337	MINGOC	
	Sinop-Ayancık	41.9483	34.7749	CA927			AT18	ANT1	ACC	AJ310337	MHG6c	
				CA930			AT32	ANT34	ACC	ACC		
				CA931				ANT1		AJ310337		
	Sinop-Gerze	41.8567	35.0999	CA932			AT18	ANT1	ACC	AJ310337	MHG6c	
				CA933			AT3	ANT33	ACC	ACC		
				CA934				ANT34		ACC		
	Sinon Erfalak Cabanlar	41 0542	35 0106	CA920			AT6	ANT1	ACC	AJ310337	MHG6c	
	Shiop-Effect Çoballia	41.9543	35.0196	CA921				ANT1		AJ310337		

	Sinon Erfelek Cobenlar	41.0542	25.0106	CA922			ANT1		AJ310337	MHG6a
	Shiop-Effelek Çobahlar	41.9545	33.0190	CA923			ANT1		AJ310337	MINOOC
				CA936			ANT1		AJ310337	
	Sinop-Kabalı village	41.8468	35.0503	CA937			ANT1		AJ310337	MHG6c
				CA938			ANT1		AJ310337	
				CA1090			ANT4		GU812186	
	Sivas-Hafik lake	39 8727	37.3825	CA1091			ANT4		GU812186	MHC6a
	Sivas-Harik lake	57.0727		CA1092		AT1	ANT1	GU812111	AJ310337	WINGOU
				CA1093			ANT1		AJ310337	
				CA1100			ANT1		AJ310337	
	Sivas-Kızılırmak	39 7154	37.0117	CA1101			ANT4		GU812186	MHG6c
Turkey			57.0117	CA1102			ANT1		AJ310337	
				CA1103			ANT1		AJ310337	
Turkey				CA1084			ANT4		GU812186	
		39.6922	37.0034	CA1085			ANT4		GU812186	MHG6c
	Sivas-Tecer river			CA1086			ANT1		AJ310337	
				CA1087		AT1	ANT1	GU812111	AJ310337	
				CA1088		AT24	ANT4	ACC	GU812186	
				CA1096			ANT4		GU812186	
	Sivas-Tödürge lake	39 8713	37 6068	CA1097		AT26	ANT4	ACC	GU812186	MHG6c
	Sivus roduige iake	57.0715	57.0000	CA1098			ANT1		AJ310337	MIIGOC
				CA1099			ANT1		AJ310337	
				CBCA6444	AN4	5	ANT21		GQ902114	
	Sorgun Dam	38.6555	29.3386	CBCA6445	AN4	5	ANT21		GQ902114	MHG6c
	-			CBCA6446	AN4	5	ANT21		GQ902114	
	Şanlıurfa-Bozova	37.3588	38.5284	CA1300			EUP5		AJ310312	MHG6d

				CA1301			EUP6		GU812213	
				CA1302		EU25	EUP7	GU812124	ACC	
	Şanlıurfa-Bozova	37.3588	38.5284	CA1303			EUP21		ACC	MHG6d
				CA1304			EUP5		AJ310312	
				CA1305			EUP5		AJ310312	
				CA1294			EUP5		AJ310312	
		37.1488		CA1295			EUP20		ACC	
	Sanluurfa Harron road		29.0691	CA1296		EU21	EUP5	ACC	AJ310312	MHC64
	Şannuna-Harran Toau		38.9681	CA1297			EUP5		AJ310312	MHOOd
				CA1298			EUP5		AJ310312	
Tester				CA1299			EUP5		AJ310312	
	Şanlıurfa-Mardin road 30. km	37.1467		CA1306			EUP5		AJ310312	
				CA1307			EUP5		AJ310312	
Turkey				CA1308			EUP5		AJ310312	
			38.1144	CA1309			EUP5		AJ310312	MHG6d
				CA1310			EUP5		AJ310312	
				CA1311			EUP5		AJ310312	
				CA1312		EU36	EUP5	GU812122	AJ310312	
				YECA15144	AN20		CAR20		GQ902086	MHG6a
				YECA15145	AN17		CAR17		AJ313132	MIIG0a
				YECA15146	AN46		ANT21		GQ902114	
	Tefenni	37 2333	29 7113	KYECA15147	AN46		ANT21		GQ902114	MHG6c
	Terenin	31.2333	29.7113	KYECA15148	AN46		ANT21		GQ902114	
				KYECA15149	AN17		CAR17		AJ313132	MHG6a
				KYECA15150	AN16		CAR16		GQ902109	
				MAHACA15151	AN48		ANT23		GU812204	MHG6c

Tefenni	37.2333	29.7113	М
Tekirdağ-Bıyıkali lake	41.0114	27.3930	

	Tefenni	37.2333	29.7113	MAHACA15152	AN46		ANT21		GQ902114	MHG6c
				CA1508			ANT10		GU812192	
				CA1509			ANT51		ACC	MHG6c
	Tekirdağ-Bıyıkali lake	41.0114	27.3930	CA1510			ANT10		GU812192	
Turkey				CA1511			RE1		AJ310338	MHG1
				CA1512			ANT13		GU812195	MHG6c
			27.5519	CA1513			ANT10		GU812192	
				CA1514		AT56	ANT10	GU812113	GU812192	
	Takirdağ Kapanakli dam	41.1121		CA1515		AT62	ANT54	ACC	ACC	MHG6a
	Teknuag-Kepenekii uani			CA1516		AT69	ANT2	ACC	GQ902107	WINGOU
Turkey				CA1517			ANT10		GU812192	
				CA1518			ANT2		GQ902107	
	Tekirdağ-Yazır lake	40.9330	27.398	CA1503			ANT10		GU812192	
				CA1504			ANT10		GU812192	MHG6c
				CA1505		AT56	ANT10	GU812113	GU812192	
	Tekirdağ-Vazır lake	40.0220	27 398	CA1506			RE1		AJ310338	MHG1
	Teknuag-Tazin lake	40.7550	27.398	CA1507		AT67	ANT13	GU812110	GU812195	MHG6c
				CA1070		AT1	ANT1	GU812111	AJ310337	MUC6
	Tokat-Aylunlar stream	40 5114	36 7342	CA1071			ANT1		AJ310337	
	Tokat-Aviunai sucan	40.5114	50.7542	CA1072			ANT1		AJ310337	WINGOU
				CA1073			ANT1		AJ310337	
-				CA1079			ANT1		AJ310337	
	Tokat-Cördük plataeu	40 2207	36 5622	CA1080		AT1	ANT1	GU812111	AJ310337	– MHG6c
	Tokat Çorduk planoti	40.2207	50.5022	CA1081			ANT1		AJ310337	
				CA1082			ANT1		AJ310337	
	Tokat-Kömeç Yeşilırmak	40.3396	36.4605	CA1055			ANT1		AJ310337	MHG6c

				CA1057				ANT1		AJ310337	
	Tokat-Kömeç Yeşilırmak	40.3396	36.4605	CA1058			AT23	ANT1	ACC	AJ310337	MHG6c
				CA1059				ANT1		AJ310337	
Turkey				CA1062				ANT1		AJ310337	
	Tokat Valancı stream	40.4106	36 7153	CA1063			AT1	ANT1	GU812111	AJ310337	MHG6a
	Tokat- Talanci sircani	40.4100	50.7155	CA1064				ANT1		AJ310337	MIIGOC
				CA1065				ANT1		AJ310337	
				KO60318		AN26		ANT1		AJ310337	
	Tokat Yusufoğlan	40.3421	36.9325	KO60319		AN29		ANT4		GU812186	MHG6c
Turkey				KO60320		AN32		ANT7		GU812189	
				CA792				ANT1		AJ310337	
	Trabzon Arsin	40.0515	39.9009	CA793				ANT1		AJ310337	
		10.9515		CA794				ANT1		AJ310337	
				CA795				ANT1		AJ310337	MHG6c
			39.5113	CA782				ANT1		AJ310337	
	Trabzon-Darica	41.0718		CA783				ANT1		AJ310337	
	Tradzon-Danca	41.0710		CA787				ANT1		AJ310337	
				CA788				ANT1		AJ310337	
				RYDCA61254		AN26		ANT1		AJ310337	
	Trabzon Derecik	41.0558	39.3900	RYDCA61255		AN26		ANT1		AJ310337	MHG6c
				RYDCA61256		AN26		ANT1		AJ310337	
	Trabzon-Maçka	40.7839	39.6126	CA796				ANT38		ACC	MHG6c
	Trahzon-Söğütlü	41 0064	39 6310	CA790				ANT1		AJ310337	MHG6c
		+1.000+	39.6310	CA791				ANT1		AJ310337	willooc
Turkey	Tuzluca	40.0334	43.6652 -	AMPH\IRA\128	A37	AN58	EU34	EUP4	GU812123	GU812212	MHG6d
	Tuzluca	40.0334		AMPH\IRA\129	A30	AN27	AT82	ANT2	HM356084	GO902107	MHG6c

				OBMBCA46373	CI9		CIE2		GU812169	
	Türkoğlu	37.3808	36.8764	OBMBCA46374	CI9		CIE2		GU812169	MHG5
				OBMBCA46375	CI10		CIE3		GU812170	
				CA1562			ANT45		ACC	
	Valova Altinova stream	40 7084	29 4730	CA1563		AT34	ANT45	ACC	ACC	MHG6c
	Taiova-Attinova sucam	40.7084	29.4750	CA1564			ANT2		GQ902107	MIIGOC
				CA1565		AT48	ANT10	ACC	GU812192	
	Valova-Altinova stream	40 7084	29 4730	CA1566		AT69	ANT2	ACC	GQ902107	MHG6c
		40.7004	27.4750	CA1567			ANT1		AJ310337	WINGOU
		40.6801	29.3907	CA1568		AT95	ANT29	ACC	GU812209	
				CA1569		AT48	ANT56	ACC	ACC	MHG6c
Turkey	Yalova-Taşköprü			CA1570		AT58	ANT2	ACC	GQ902107	
				CA1571			ANT2		GQ902107	
Turkey				CA1572		AT48	ANT10	ACC	GU812192	
				ISCA1578	AN46		ANT21		GQ902114	MHG6c
				ISCA1579	AN17		CAR17		AJ313132	MHG6a
				ISCA1580	AN1		CAR1		AJ310314	
				ISCA1581	AN1		CAR1		AJ310314	
				ISCA1582	AN46		ANT21		GQ902114	-
	Yesilova	37,5348	29.6473	ISCA1583	AN46		ANT21		GQ902114	-
	r byno ru	0110010	20101110	ISCA1584	AN46		ANT21		GQ902114	-
				ISCA1585	AN46		ANT21		GQ902114	MHG6c
				ISCA1586	AN46		ANT21		GQ902114	-
				ISCA1587	AN46		ANT21		GQ902114	
				ISCA1588	AN46		ANT21		GQ902114	
				ISCA1589	AN7		CAR7		GQ902112	MHG6a

				CA1111		AT1	ANT1	GU812111	AJ310337	
	Vozgat-Gedikhasanlı	39 5720	35 1373	CA1112			ANT1		AJ310337	MHG6c
	1 ozgat-Oculkilasalili	39.3720	55.1575	CA1113			ANT1		AJ310337	WINGOU
				CA1114			ANT1		AJ310337	
				CA1117			ANT1		AJ310337	
			35.2215	CA1118		AT1	ANT1	GU812111	AJ310337	
	Yozgat-Mükremin lake	39.8030		CA1119		AT35	ANT47	ACC	ACC	MHG6c
				CA1120			ANT1		AJ310337	
				CA1121			ANT1		AJ310337	
	Vozgat Sorgun	30 8042	35.2035	CA1105		AT37	ANT1	ACC	AJ310337	MHG6c
Turkey	i ozgat-Sorgun	39.0042		CA1106			ANT1		AJ310337	MHG0e
	Vozget Sorgun	20 8042	25 2025	CA1107			ANT1		AJ310337	MHC6a
	102gat-Sorgun	39.0042	35.2035	CA1108			ANT1		AJ310337	MHG0e
Turkey	Vozat Varköv Daliaa rivar	39.6117		CA1124			ANT1		AJ310337	
			34.5031	CA1125			ANT1		AJ310337	MHG6c
	Tozgat-Terkoy Dence fiver			CA1126		AT1	ANT1	GU812111	AJ310337	
				CA1127			ANT1		AJ310337	
				TBCASE0154	CI1		CIW1		AJ313135	
				TBCASE0155	CI1		CIW1		AJ313135	
				TBCASE0156	CI3		CIW3		GU812163	
				TBCASE0157	CI4		CIW4		GU812164	
	Yumurtalık	36.6925	35.6288	TBCASE0158	CI4		CIW4		GU812164	MHG4
				TBCASE0159	CI1		CIW1		AJ313135	-
				TBCASE0160	CI5		CIW5		GU812165	
				TBCASE0161	CI3		CIW3		GU812163	
				TBCASE0162	CI3		CIW3		GU812163	

				TBCASE0163	CI7		CIW7		GU812167	
	Vumurtalık	36 6025	35 6788	TBCASE0164	CI2		CIW2		GU812162	MHG4
	i unutaik	30.0923	55.0288	TBCASE0165	CI2		CIW2		GU812162	WI104
				TBCASE0166	CII		CIW1		AJ313135	
		38.3443		CA638		EU15	EUP1	ACC	AJ310313	
				CA639			EUP1		AJ310313	
	Van-Edremit Çiçekli		43.1881	CA641		EU24	EUP5	ACC	AJ310312	MHG6d
				CA643			EUP1		AJ310313	
				CA644			EUP5		AJ310312	
Turkey	Van-Gevaş		43.1232	CA645			EUP5		AJ310312	
		38.3079		CA646		EU24	EUP5	ACC	AJ310312	
				CA647		EU15	EUP1	ACC	AJ310313	MHG6d
				CA649			EUP5		AJ310312	
				CA650			EUP5		AJ310312	
				CA651			EUP5		AJ310312	
				CA652			EUP5		AJ310312	MHG6d
	Van-Muradiye Bendimahı River	38 9366	43 6611	CA653		EU24	EUP14	ACC	ACC	
	van Maradaye Benemian Ferver	50.7500	15.0011	CA654			EUP5		AJ310312	Milloou
				CA656			EUP5		AJ310312	
				CA1732			ANT21		GQ902114	
				CA1733			ANT21		GQ902114	
				CA1734			ANT21		GQ902114	
	Uşak-Gediz river	38.7878	29.2243	CA1735			ANT29		GU812209	MHG6c
				CA1736			ANT29		GU812209	
				CA1742			ANT21		GQ902114	
				CA1743			ANT21		GQ902114	

	Uşak-Gediz river	38.7878	29.2243	CA1744		АТ	Г110	ANT21	GU812104	GQ902114	MHG6c
	Usak Göğan laka	28 7207	20 5562	CA1745				ANT21		GQ902114	MHC6a
	Uşak-Oogen lake	38.7207	29.3303	CA1746				ANT21		GQ902114	MINGOC
				CA1737		АТ	Г110	ANT21	GU812104	GQ902114	
				CA1738				ANT29		GU812209	
	Uşak-Karakoç lake	38,6542	29,3336	CA1739				ANT13		GU812195	MHG6c
				CA1740		A	T55	ANT10	ACC	GU812192	
				CA1741				ANT21		GQ902114	
				FGCA67248	AN	133		ANT8		GU812190	
		41.4271	31.7268	FGCA67249	AN	134		ANT9		GU812191	
Turkey	Zonguldak			FGCA67250	AN	134		ANT9		GU812191	MHG6c
				FGCA67251	AN	126		ANT1		AJ310337	
				FGCA67252	AN	126		ANT1		AJ310337	
		41.1515	31.3521	CA1013		A	T47	ANT9	ACC	GU812191	
	Zonguldak-Alanlı			CA1014		A	T38	ANT1	ACC	AJ310337	MHG6c
	Zonguldak-Alaph			CA1015				ANT1		AJ310337	
				CA1016				ANT1		AJ310337	
				CA1002		А	AT1	ANT1	GU812111	AJ310337	
				CA1003		A	T46	ANT9	ACC	GU812191	
	Zonguldak-Kilimli	41.4005	31.6837	CA1004				ANT1		AJ310337	MHG6c
				CA1005				ANT1		AJ310337	
				CA1006				ANT9		GU812191	
				CA996		А	T 1	ANT30	GU812111	ACC	
	Zonguldak-Sirinköy	41 5065	31 9726	CA997		A	T36	ANT1	ACC	AJ310337	MHG6c
	Zonguldak-Şirinköy 4	41.5005	31.9726	CA998		А	T 1	ANT30	GU812111	ACC	MINGOC
				CA1000				ANT1		AJ310337	

Turkmenistan	Ashkhabad	38 3214	57 9/98	ZMB45638	C1		T1	TE5	GU812127	AJ310317	MHG7
Turkmenistan	Asiikiiabau	30.3214	57.9490	ZMB45494				TE5		AJ310317	WIIG/
Ukraine	Alushta	11 7528	34 4058	ZISP.6050-Sp.88		AN31		ANT6		GU812188	MHG6c
Ukraine	Alusita	44.7528	54.4058	ZISPSp.245		AN31		ANT6		GU812188	MHGoe
	Chervlyonaya Gusirovka	49.4704	36.8705	ZISPSp.305		RE9		RE9		GU812155	MHG1
	Palagorsk	45 0727	24 5072	Czech-U3-1		AN42		ANT17		GU812199	MHC6a
	Belogorsk	45.0757	54.5975	Czech-U3-2		AN42		ANT17		GU812199	MHGoe
	Delekamannaa	11 6917	22 8020	Czech-U19-1		AN45		ANT20		GU812202	MUCGa
	Belokamennoe	44.0847	55.8950	Czech-U19-2		AN45		ANT20		GU812202	MINGOC
Libracia a				ZISP.5611-Sp.92		RE6		RE6		AM900653	
Ukraine	Chernobyl	51.2696	30.2335	ZISP.5611-Sp.93		RE6		RE6		AM900653	MHG1
				ZISP.5611-Sp.94		RE6		RE6		AM900653	
	Dolinnoe	44.7538	33.7673	Czech-U18		AN45		ANT20		GU812202	MHG6c
				ZISP.6641-Sp.96		RE6		RE6		AM900653	
	Golaya Pristan	46.5344	32.5230	ZISP.6641-Sp.97		RE6		RE6		AM900653	MHG1
				ZISP.6641-Sp.98		RE6		RE6		AM900653	
	Irpen	50.5465	30.2773	Czech-U54		RE6		RE6		AM900653	MHG1
	Kerch	45.3607	36.4761	ZISP.6642-Sp.143		AN43		ANT18		GU812200	MHG6c
Libracia a	Kiev	50.4170	30.5106	ZMB25779	R4	RE6	R4	RE6	AM900652	AM900653	MHG1
Ukraine	Korostyshiv	50.3211	29.0726	Czech-U53		RE6		RE6		AM900653	MHG1
	Lavanda	44.7532	34.3696	ZISPSp.245		AN31		ANT6		GU812188	MHG6c
	Luchistoe	44.7513	34.4045	ZISP.6050-Sp.88		AN31		ANT6		GU812188	MHG6c
Uzbakistar	Control Nurotou	40.0812	66 1177	-	C2		T2	TE13	GU812128	AJ310318	MHC7
OZUCKISTAII	Central Inulatau	40.0012	00.4177	-	C3		T3	TE13	GU812129	AJ310318	MINU/

Outgroups										
Italy	Metaponto	40.3737	16.8028	No.6311	BER1	Ber1		GU812133	GU812231	
itary	Tarsia	39.6166	16.2666	Hotz19586	BER2	Ber2		GU812134	GU812232	P. bergeri
Corsica	Solenzara	41.8500	9.3999	-	BER3	Ber3		GU812135	GU812233	
	Skinias/Crete	35.0632	25.3083	Hotz19852/ZMB56959	CRE1	Cre1		GU812136	GU812234	
Greece				Hotz19854/ZMB56942	CRE2	Cre2		GU812137	AJ310336	P. cretensis
	Demati/Crete	35.0333	25.2833	Hotz19856/ZMB56960	CRE3	Cre2		GU812138	AJ313136	
	Igoumenitsa	39.5000	20.2666	Hotz19421	EPE1	Epe1		GU812139	GU812235	
Greece				Ploetner181.03	EPE2	Epe2		GU812140	GU812236	P. epeiroticus
	Lechena/Peloponnese	37.9174	21.2667	Hotz19552	EPE3	Epe3		GU812141	GU812237	
Italy	Carbonare	45.9333	11.2166	-	LES1	Les1		AM887975	AM887975	
Poland	Rogaczewo	52.0666	16.8166	Hotz18472	LES2	Les2		AM887976	AM887976	P. lessonae
Romania	Caraorman	45.1000	29.3333	ZMB47456	LES3	Les1		AM887968	AM887975	
France	Ligagnaeu	43.5333	4.7500	Lib11	PER1	Per1		GU812142	GU812238	
Trailee	Narbonne	43.1833	3.0166	-	PER2	Per2		GU812143	GU812239	P. perezi
Spain	Bajamar/Tenerife	28.5333	16.3333	-	PER3	Per3		GU812144	GU812240	
Algeria	Ain Salah	27.0000	2.4666	Hotz16724	SAH1	Sah1		GU812145	GU812241	
Morocco	Ait Boukha	28.9833	10.0666	Hotz16647	SAH2	Sah2		GU812146	GU812242	P. saharicus
Tunis	Tunis	c. 36.8001	10.17020	-	SAH3	Sah3		GU812147	GU812243	
Montenegro	Lake Skutari/Virpazar	42.2333	19.1000	ZMB47496	SHQ1	Shq1		GU812148	GU812244	P shainariaus
Wontenegro				Hotz17523	SHQ2	Shq2		GU812149	GU812245	1. snqipericus
North Korea	unknown			-	NIG1	Nig1		GU812150	GU812246	<i>P</i> .
Japan	unknown			-	NIG2	Nig2		AB043889	AB043889	nigromaculatus

Table 3.3. Locality information of nuclear SAI-1 alleles. Locality, collection or laboratory ID, nuclear allele composition for SAI-1 marker of the investigated individuals. Dark grey coloured fields show samples used in the first time in this study, the rest was taken from former publications. Sequences beginning with FN and HE were obtained from EMBL database. They refer to the publications of Plötner et al. 2009 and 2012. SAI-1: intron of the serum albumin gene containing a 5' truncated chicken repeat 1 like long interspersed nuclear element called as *Rana*CR1 by Plötner et al. (2009).

Country	Locality	Locality Latitude Longitude Collection/ Laboratory ID	Collection / Laboratory ID	SAI-1 allele-	SAI-1	Accession Number		Allele Group/	
Country	Locality	Lautuue	Longhude	Conection/ Laboratory ID	Α	allele-B	Allele-A	Allele-B	Species
	Ararat Yeraskh	39.7357	44.8326	Ilona1	ant11-7	ant11-8	ACC	ACC	ANT
	Armavir Vanand village	40.1144	43.8235	Ilona2	ant6-2	ant6-2	ACC	ACC	ANT
Armenia	Gegharkunik Martuni town	40.1091	45.3000	Ilona3	ant6-1	ant6-2	ACC	ACC	ANT
Timenia	Kotaik Hrazdan	40.4961	44.7662	Ilona4	ant11-7	ant11-7	ACC	ACC	ANT
	Kotaik-Bjni town	40.3002	44.5047	Ilona5	ant6-1	ant11-8	ACC	ACC	ANT
	Vayots' Dzor -Yeghegnadzor	39.7539	45.3270	Ilona6	ant11-8	ant11-8	ACC	ACC	ANT
Karabakh Region	Nagorno-Sharifan	39.2649	46.9714	Ilona7	ant7-3	ter-2	ACC	ACC	ANT/TER
Karabakii Kegioli	Nagorno-Kubatlı	39.5559	46.9836	Ilona8	ant11-8	ant11-8	ACC	ACC	ANT
	between Ayia Napa and Cape Greco	34.9774	34.0327	P36	cyp-2	cyp-2	HE858241	HE858241	СҮР
	Northern Cyprus	35.1855	33.5674	Hotz19410	cyp-2	ant5-6	HE858241	HE858230	CYP/ANT
				Hotz19411	cyp-2	cyp-3	HE858241	ACC	CYP
				Hotz19412	cyp-2	cyp-2	HE858241	HE858241	
				Hotz19413	cyp-2	cyp-2	HE858241	HE858241	
Cyprus				Hotz19416	cyp-2	ant5-5	HE858241	ACC	CYP/ANT
	Cape Greco (water reservoir in the west)	34.9794	34.0548	P24	cyp-1	cyp-2	HE858244	HE858241	СҮР
	Stream close to Gialia	35.0769	32.5690	P32	cyp-1	cyp-1	HE858244	HE858244	CYP
				P33	cyp-1	cyp-2	HE858244	HE858241	
	Gönyeli dam	35.2332	33.2937	Р3	cyp-2	cyp-2	HE858241	HE858241	CYP
	Maroullenas-1	35.0084	33.1477	Р5	cyp-2	cyp-2	HE858242	HE858242	СҮР

between Mathiatis and Kataliontas	34.9782	33.3321	P38	cyp-2	cyp-2	HE858241	HE858241	
Nicosia-2	35.1951	33.3189	CA1985	cyp-3	cyp-3	ACC	ACC	CYP
Panagia (forest station at stream Peristerona)	35.0156	33.0833	P4	cyp-2	cyp-2	HE858243	HE858243	СҮР
Lake Paralimni	35.0358	33.9753	P20	cyp-2	cyp-2	HE858241	HE858241	CYP
Paristarona 1	34 9770	33 0012	P34	cyp-2	cyp-2	HE858241	HE858241	CVP
Tensterona-1	54.9770	55.0912	P35	cyp-2	cyp-2	HE858241	HE858241	СП
			Р9	cyp-1	cyp-1	HE858244	HE858244	
Pyrgos Dam	35.1495	32.6549	P11	cyp-1	cyp-1	HE858244	HE858244	CYP
			P13	cyp-1	cyp-2	HE858244	HE858241	
Troodos-2	34.9161	32.9003	ZISP10531	cyp-2	cyp-2	HE858245	HE858245	CYP
			PL01-2012	rid2-3	rid2-3	FN432365	FN432365	RID
			PL02-2012	kur-3	kur-3	ACC	ACC	KUR
Citoy	50 3751	14 4415	PL03-2012	rid2-3	rid2-3	FN432365	FN432365	RID
Chov	50.5751	14.4415	PL06-2012	kur-3	kur-3	ACC	ACC	KIID
			PL07-2012	kur-3	kur-3	ACC	ACC	KOK
			PL11-2013	ant7-3	ant7-3	ACC	ACC	ANT
Lebus near Frankfurt/O.	52.4141	14.5422		rid2-4	rid2-4	FN432364	FN432364	RID
			PL82-2012	rid2-3	rid2-3	FN432365	FN432365	
			PL83-2012	rid2-3	rid2-3	FN432365	FN432365	RID
Oder River	52.4213	14.5333	PL84-2012	rid2-3	rid2-3	FN432365	FN432365	
			PL85-2012	rid2-3	ant7-3	FN432365	ACC	RID/ANT
			PL86-2013	rid2-3	rid2-3	FN432365	FN432365	RID
Aliartos	38 3688	23 08/11	Hotz17357	rid2-2	rid2-2	HE857212	HE857212	RID
Allatos	56.5066	23.0041	Hotz17360	rid2-2	rid2-2	HE857213	HE857213	NID
Archipolis/Rhodos	36.3117	28.1360	Hotz17220	cer-1	cer-1	HE850228	HE850228	CER
	between Mathiatis and Kataliontas Nicosia-2 Panagia (forest station at stream Peristerona) Lake Paralimni Peristerona-1 Pyrgos Dam Troodos-2 Citov Lebus near Frankfurt/O. Oder River Aliartos Archipolis/Rhodos	between Mathiatis and Kataliontas34.9782Nicosia-235.1951Panagia (forest station at stream Peristerona)35.0156Lake Paralimni35.0358Peristerona-134.9770Pyrgos Dam35.1495Troodos-234.9161Citov50.3751Lebus near Frankfurt/O.52.4141Oder River52.4213Aliartos38.3688Archipolis/Rhodos36.3117	between Mathiatis and Kataliontas 34.9782 33.3321 Nicosia-2 35.1951 33.3189 Panagia (forest station at stream Peristerona) 35.0156 33.0833 Lake Paralimni 35.0358 33.9753 Peristerona-1 34.9770 33.0912 Pyrgos Dam 35.1495 32.6549 Troodos-2 34.9161 32.9003 Citov 50.3751 14.4415 Lebus near Frankfurt/O. 52.4141 14.5422 Oder River 52.4213 14.5333 Aliartos 38.3688 23.0841 Archipolis/Rhodos 36.3117 28.1360	between Mathiatis and Kataliontas 34.9782 33.3321 P38 Nicosia-2 35.1951 33.3189 CA1985 Panagia (forest station at stream Peristerona) 35.0156 33.0833 P4 Lake Paralimni 35.0358 33.9753 P20 Lake Paralimni 35.0358 33.9753 P20 Peristerona-1 34.9770 33.0912 P34 Pyrgos Dam 35.1495 32.6549 P11 P13 32.9003 ZISP10531 P10 P13 32.9003 ZISP10531 P10.2012 PL01-2012 PL02-2012 PL02-2012 PL02-2012 PL03-2012 PL03-2012 Citov 52.4141 14.5422 PL06-2012 PL07-2012 PL11-2013 PL82-2012 PL83-2012 Oder River 52.4213 14.5333 PL82-2012 PL85-2012 PL85-2012 PL85-2012 PL85-2013 PL86-2013 Hotz17357 Aliartos 36.3117 28.1360 Hotz17220	between Mathiatis and Kataliontas 34.9782 33.3321 $P38$ $cyp-2$ Nicosia-2 35.1951 33.3189 $CA1985$ $cyp-3$ Panagia (forest station at stream Peristerona) 35.0156 33.0833 $P4$ $cyp-2$ Lake Paralimni 35.0358 33.9753 $P20$ $cyp-2$ Peristerona-1 34.9770 33.0912 $P34$ $cyp-2$ Peristerona-1 34.9770 33.0912 $P34$ $cyp-2$ Pyrgos Dam 35.1495 32.6549 $P11$ $cyp-1$ Pyrgos Dam 35.1495 32.6549 $P11$ $cyp-2$ Troodos-2 34.9161 32.9003 $ZISP10531$ $cyp-2$ Citov 50.3751 14.4415 $PL02.2012$ kur-3 PL07-2012 kur-3 $PL07-2012$ kur-3 Oder River 52.4141 14.5422 $PL82-2012$ $rid2-3$ PL3 $cyp-1$ $rid2-3$ $PL8-2012$ $rid2-3$ Oder River 52.4	between Mathiatis and Kataliontas 34.9782 33.321 $P38$ $cyp-2$ $cyp-3$ Nicosia-2 35.1951 33.3189 $CA1985$ $cyp-3$ $cyp-3$ Panagia (forest station at stream Peristerona) 35.0156 33.0833 $P4$ $cyp-2$ $cyp-2$ Lake Paralimni 35.0358 33.9753 $P20$ $cyp-2$ $cyp-2$ Peristerona-1 34.9770 33.0912 $P34$ $cyp-2$ $cyp-2$ Pyrgos Dam 35.1495 32.6549 P11 $cyp-1$ $cyp-1$ Pyrgos Dam 35.1495 32.6549 P11 $cyp-1$ $cyp-1$ Pyrgos Dam 35.1495 32.6549 P11 $cyp-1$ $cyp-2$ Troodos-2 34.9161 32.9003 ZISP10531 $cyp-2$ $cyp-2$ Citov 50.3751 14.4415 PL02-2012 $kur-3$ $kur-3$ PL07-2012 $kur-3$ $kur-3$ $kur-3$ $kur-3$ $kur-3$ Dder River 52.4213	between Mathiatis and Kataliontas 34.9782 33.3321 $P38$ $cyp-2$ $cyp-2$ HE858241 Nicosia-2 35.1951 33.3189 CA1985 $cyp-3$ $cyp-3$ ACC Panagia (forest station at stream Peristerona) 35.0156 33.0833 P4 $cyp-2$ $cyp-2$ HE858243 Lake Paralimni 35.0358 33.9753 P20 $cyp-2$ $cyp-2$ HE858241 Peristerona-1 34.9770 33.0912 P34 $cyp-2$ $cyp-2$ HE858241 Peristerona-1 34.9770 33.0912 P35 $cyp-2$ $cyp-2$ HE858241 Pyrgos Dam 35.1495 32.6549 P11 $cyp-1$ $cyp-2$ HE858244 Troodos-2 34.9161 32.9003 ZISP10531 $cyp-2$ $cyp-2$ HE858245 Citov 50.3751 14.4415 P1.01-2012 $rid-3$ $rid-3$ FN432365 PL07-2012 kur-3 kur-3 ACC P1.11-2013 $an7-3$ ACC	between Mathiatis and Kataliontas 34.9782 33.3321 P38 cyp-2 cyp-2 HE858241 HE858241 Nicosia-2 35.1951 33.3189 CA1985 cyp-3 cyp-3 ACC ACC Panagia (forest station at stream Peristerona) 35.0156 33.0833 P4 cyp-2 cyp-2 HE858243 HE858243 Lake Paralimni 35.0358 33.9753 P20 cyp-2 cyp-2 HE858241 HE858241 Peristerona-1 34.9770 33.0912 P34 cyp-2 cyp-2 HE858241 HE858241 Pyrgos Dam 35.1495 32.6549 P11 cyp-1 cyp-1 HE858244 HE858244 Pyrgos Dam 35.1495 32.6549 P13 cyp-2 HE858244 HE858244 Pyrgos Dam 35.1495 32.0654 P13 cyp-2 HE858244 HE858244 Pyrgos Dam 52.4191 32.003 ZISP10531 cyp-2 HE858244 HE858244 Pyrgos Dam 50.3751

	Archipolis/Rhodos	36.3117	28.1360	Hotz17221	cer-1	cer-1	HE850228	HE850228	CER
	Kayagilag	27 9762	21 2854	Hotz19510	kur-5	kur-5	HE858214	HE858214	VID
	Kavasiias	37.8702	21.2654	Hotz19528	kur-2	kur-2	FN432366	FN432366	KUK
	Neo Manolada	38.0316	21 3611	Hotz19532	kur-1	kur-4	ACC	HE858215	KIID
	ivea Wanolada	38.0310	21.3011	Hotz19538	kur-4	kur-4	FN432367	FN432367	KUK
	Olympos/Karpathos	35 7427	27 1685	Hotz17207	cer-1	cer-1	HE858227	HE858227	CEP
Graaca	Orympos/Karpanios	33.7427	27.1085	Hotz17208	cer-1	cer-1	HE858227	HE858227	CEK
Gleece	Skala	38 6675	23.0712	Hotz19483	kur-4	kur-4	HE858215	HE858215	KIID
	Skala	38.0075	23.0712	Hotz19485	kur-1	kur-1	ACC	ACC	KUK
	Skala	38 6675	23.0712	Hotz19500	kur-1	kur-4	ACC	HE858215	KIID
	Skala	38.0075	23.0712	Hotz19501	kur-1	kur-1	ACC	ACC	KUK
				Ploetner30.03	kur-4	kur-4	HE858215	HE858215	
	Zakynthos	37.8052	20.8627	Ploetner31.03	kur-4	kur-4	HE858215	HE858215	KUR
				Ploetner32.03	kur-4	kur-4	HE858215	HE858215	
				AMPH\IRA\509	ter-2	ter-2	ACC	ACC	
	Babol Sar	36.6803	52.6357	AMPH\IRA\510	ter-2	ter-2	ACC	ACC	TER
				AMPH\IRA\511	ter-2	ter-2	ACC	ACC	
	Bavineh	33.6042	47.2026	AMPH\IRA\338	ter-5	ter-2	ACC	ACC	TER
				AMPH\IRA\308	ter-2	ter-2	ACC	ACC	
Iran	Bisotun	34.4038	47.4483	AMPH\IRA\309	ter-5	ter-2	ACC	ACC	TER
iran				AMPH\IRA\310	ter-2	ter-2	ACC	ACC	
	Choplu	36.4739	47.0393	AMPH\IRA\302	ter-2	ter-2	ACC	ACC	TER
				AMPH\IRA\328	ter-3	ter-2	ACC	ACC	
	Choqa Zanbil	32 0177	48.5451	AMPH\IRA\331	ter-2	ter-2	ACC	ACC	TER
		52.0177		AMPH\IRA\332	ter-2	ter-2	ACC	ACC	
				AMPH\IRA\333	ant11-7	ant3-5	ACC	HE858238	ANT

Iran	Choqa Zanbil	32.0177	48.5451	AMPH\IRA\336	ter-3	ter-6	ACC	ACC	TER
li ali	Now Kandeh	36.7154	53.8836	AMPH\IRA\520	ter-2	ter-2	ACC	ACC	TER
	Al Kerak	31.1833	35.7000	J-06-54	bed1-3	cil3-3	HE858221	ACC	BED1/CIL
	Al shouna Al Shamaliya	32.6155	35.6388	J-02-50	bed1-3	bed1-3	HE858221	HE858221	BED1
				J-01-31	bed1-3	bed1-4	HE858221	ACC	BED1
				J-01-44	bed2-3	bed2-3	ACC	ACC	
	Bab Amman	32.2042	35.8867	J-01-55	bed2-3	bed2-2	ACC	ACC	BED2
				J-01-56	bed2-1	bed2-1	HE858239	HE858239	
				J-01-57	bed2-2	bed2-2	ACC	ACC	
				J-05-39	bed2-1	bed2-1	HE858240	HE858240	
				J-05-45	bed2-1	bed2-1	HE858240	HE858240	
Iordan	Canyon near Wadi Mujib	31.4492	35.7892	J-05-46	bed2-1	bed2-1	HE858240	HE858240	BED2
				J-05-47	bed2-4	bed2-1	ACC	HE858240	
				J-05-48	bed2-1	bed2-1	HE858240	HE858240	
Jordan				J-03-35	bed1-3	bed1-3	HE858221	HE858221	BED1
	16 km southward from Jesus Baptisizing	31 /030	35 5861	J-03-36	bed1-3	cil3-2	HE858221	ACC	BED1/CIL
	site	51.4757	55.5601	J-03-37	bed1-3	bed1-3	HE858221	HE858221	BED1
				J-03-38	bed1-3	cil3-2	HE858221	ACC	BED1/CIL
				J-04-30	bed2-1	bed2-1	HE858218	HE858218	
				J-04-31	bed1-3	bed1-3	HE858218	HE858218	
				J-04-32	bed2-1	bed2-1	HE858218	HE858218	
	Mulaik Thoba	31 5533	35 7400	J-04-33	bed2-1	bed2-1	HE858219	HE858219	BED2
		0110000	2217 100	J-04-41	bed2-1	bed2-1	HE858219	HE858219	
				J-04-51	bed2-1	bed2-1	HE858219	HE858219	
				J-04-52	bed2-1	bed2-1	HE858220	HE858220	
				J-04-53	bed2-1	bed2-1	HE858220	HE858220	

				Sp.253	rid1-5	rid1-5	FN432363	FN432363	RID
				Sp.254	ant8-4	rid1-4	HE858222	HE858211	RID/ANT
Kazakhstan Poland Syria Turkey				Sp.255	ant8-4	ant8-4	HE858222	HE858222	ANT
				41-1	ant7-3	rid1-5	ACC	FN432363	
	Atyrau	47.5442	52.3861	41-1L	ant7-3	rid1-5	ACC	FN432363	
				41-2	ant7-3	rid1-5	ACC	FN432363	KID/AN1
				41-2L	ant7-3	rid1-5	ACC	FN432363	
				41-3	ant8-2	ant8-2	ACC	ACC	ANT
				41-5	ant7-3	rid1-5	ACC	FN432363	RID/ANT
Kazakhstan	Aturau Kulsary taun	46.9611	54 0091	40-3	ant7-3	ant7-3	ACC	ACC	ANT
Kazaklistali	Atyrau-Kuisary town	40.9011	54.0091	40-5	ant7-3	ant6-2	ACC	ACC	ANI
				39-03	rid1-6	rid1-6	ACC	ACC	
	Mangistau-The Saura spring	44.2319	50.803	39-19	rid1-5	rid1-5	ACC	ACC	RID
				39-24	rid1-5	rid1-5	ACC	ACC	
				38-03	rid1-5	rid1-5	ACC	ACC	
				38-06	rid1-5	rid1-5	ACC	ACC	BID
	Mangistau-The Tymshaly spring	44.6014	50.5966	38-07	rid1-5	rid1-5	ACC	ACC	KID
				38-08	rid1-5	rid1-5	ACC	ACC	
				38-09	ant8-2	ant8-2	ACC	ACC	ANT
	Dzhambul-Taraz city	42.8838	71.4122	36-20	sp nov-1	sp nov-1	ACC	ACC	SP NOV
Poland	Poznan	52.3805	16.6674	Hotz18192	rid2-3	rid2-3	FN432365	FN432365	RID
Syria	As Suwayda	32.6855	36.5525	ZFMK64945	bed1-1	bed1-1	FN432368	ACC	BED1
				CA1418	cil1-4	cil1-4	HE858225	HE858225	CII
Turkey	Adana-Ceyhan Cinderesi	37 0353	35 7470	CA1421	cil1-6	cil1-5	ACC	ACC	CIL
Turkey	Adana-Ceynan Cinderesi	51.0555	33.1410	CA1422	cil1-4	ant3-5	HE858225	HE858238	CIL/ANT
ž				CA1423	cil1-4	cil1-4	HE858225	HE858225	CIL

				CA1922	cil1-4	ant11-7	HE858225	ACC	CIL/ANT
Turkey	Adana-Pozantı Şekerpınarı	37.4080	34.8842	CA1924	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA427	cil1-4	ant7-3	HE858225	ACC	CIL/ANT
				CA1927	cil1-4	cil1-4	HE858225	HE858225	CII
				CA1928	cil1-4	cil1-4	HE858225	HE858225	CIL
	Adama Darante Calut niven	27 4200	24.9764	CA1929	cil1-4	ant11-7	HE858225	ACC	
	Adana-Pozanti Çakit river	37.4299	34.8704	CA1930	cil1-4	ant3-5	HE858225	HE858238	CIL/ANT
				CA1931	cil1-4	ant11-7	HE858225	ACC	
				CA1932	rid3-5	rid3-6	ACC	ACC	RID
	A demogram Desmoglam laise	40.9226	20,4692	CA1541	rid2-1	ant11-7	ACC	ACC	RID/ANT
	Adapazari-Foylaziai lake	40.8330	30.4082	CA1542	ant7-2	ant7-3	ACC	ACC	ANT
				CA1545	ant7-2	ant11-7	ACC	ACC	ANT
Turker	Adapazarı-Saklıgöl	40.8549	30.3018	CA1547	rid2-1	ant7-2	ACC	ACC	RID/ANT
Turkey				CA1548	ant11-7	ant11-7	ACC	ACC	ANT
				CA1270	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1271	ant11-7	cil1-4	ACC	HE858225	CIL/ANT
	Adıyaman-Abuzergaffar river	37.7415	38.3354	CA1273	ant11-9	ant11-9	ACC	ACC	
				CA1275	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA1276	ant11-7	ant11-9	ACC	ACC	
				CA1287	rid3-8	rid3-8	ACC	ACC	RID
				CA1289	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Adıyaman-Börgenek Çakal stream	37.7128	38.1663	CA1290	ant10-3	cil2-1	ACC	ACC	CIL/ANT
				CA1292	rid3-8	ant11-7	ACC	ACC	
				CA1293	rid3-7	ant3-5	ACC	HE858238	KID/ANT
	Adıyaman-Göksu river	37 6983	38.0799	CA1282	rid3-8	cil2-1	ACC	ACC	CIL/RID
		37.6983		CA1283	rid3-8	ant11-7	ACC	ACC	RID/ANT

	Adwaman Gökey river	27 6082	28.0700	CA1284	ant11-7	cil2-1	ACC	ACC	CIL/ANT
	Aufyanian-Ooksu niver	37.0983	38.0799	CA1286	rid3-8	cil1-4	ACC	HE858225	CIL/RID
				CA1277	rid3-5	ant11-9	ACC	ACC	RID/ANT
	Adwaman Kahta Piraik rivar	27 7447	38 5060	CA1278	rid3-8	rid3-8	ACC	ACC	PID
	Aufyanian-Kanta Bircik fiver	37.7447	38.3000	CA1279	rid3-8	rid3-8	ACC	ACC	KID
				CA1280	ant11-9	cil1-4	ACC	HE858225	CIL/ANT
				CA1725	ant10-1	ant10-1	ACC	ACC	ANT
				CA1726	ant3-2	ant7-3	ACC	ACC	ANI
	A frienkarshicar 26 A guat Natural Bark	38.7925	20 2816	CA1727	rid2-1	ant3-2	ACC	ACC	RID/ANT
	Aryonkaranisar-20 Agust Naturai Park		50.5810	CA1728	ant1-4	ant10-4	ACC	ACC	ANT
Turkey				CA1729	ant10-1	ant10-4	ACC	ACC	
				CA1731	ant2-1	ant8-5	ACC	ACC	
				CA1720	ant11-7	ant10-1	ACC	ACC	ANT
				CA1721	ant10-1	ant10-4	ACC	ACC	ANI
	Afyonkarahisar-Cumhuriyet village Çay	38.5923	30.9618	CA1722	rid2-1	ant10-4	ACC	ACC	RID/ANT
				CA1723	ant10-1	ant10-4	ACC	ACC	
				CA1724	ant1-3	ant8-3	ACC	ACC	ANT
				CA1710	ant10-4	ant10-4	ACC	ACC	
				CA1711	ant10-4	ant10-4	ACC	ACC	
	Afyonkarahisar-Karamık lake	38.4215	30.8869	CA1712	ant10-1	ant10-4	ACC	ACC	ANT
				CA1713	ant8-3	ant10-4	ACC	ACC	
				CA1714	ant8-3	ant10-4	ACC	ACC	
				CA679	ant11-8	ant11-8	ACC	ACC	ANT
	Ağrı-Aşağı Voldüzü Cuma Piyer	39 8181	43 0892	CA680	rid3-8	ant11-8	ACC	ACC	PID/ANT
	Ağrı-Aşağı Yoldüzü Cuma River 39	37.0101	43.0892	CA681	ant11-7	rid3-8	ACC	ACC	KID/ANT
				CA683	ant11-8	ant11-8	ACC	ACC	ANT

				CA684	rid3-8	rid3-8	ACC	ACC	RID
	Ağrı-Çukuralan Village	39.7130	42.9920	CA686	rid3-8	ant11-8	ACC	ACC	RID/ANT
				CA688	ant11-8	ant11-8	ACC	ACC	ANT
				CA672	ant11-8	ant11-8	ACC	ACC	ANT
				CA673	ant6-1	rid3-8	ACC	ACC	
	Ağrı Damhat yıllaga	39 6769	43 0228	CA674	rid3-8	ant11-8	ACC	ACC	KID/ANT
	Agn-Damoat vinage	39.0709	43.0228	CA675	ant11-8	ant11-8	ACC	ACC	ANT
				CA676	rid3-8	ant11-8	ACC	ACC	RID/ANT
				CA678	ant11-8	ant11-8	ACC	ACC	ANT
				CA658	ant11-8	ant11-8	ACC	ACC	ANT
	Ağrı Doğu Poyezit Perdeklı villago	20.681	44.0655	CA659	rid3-8	ant11-8	ACC	ACC	RID/ANT
Turkey -	Agir-Dogu Deyazit Datuakii vinage	39.081	44.0655	CA660	ant11-8	ant11-8	ACC	ACC	ANT
				CA661	rid3-8	ant11-8	ACC	ACC	RID/ANT
Turkey	Aksaray-Akın village stream		34.0291	CA1225	rid3-5	rid3-5	ACC	ACC	PID
		38.4313		CA1226	rid3-5	rid3-5	ACC	ACC	KID
				CA1227	ant11-7	rid2-1	ACC	ACC	RID/ANT
				CA1212	ant11-7	ant11-7	ACC	ACC	ANT
				CA1213	rid3-5	rid3-5	ACC	ACC	RID
				CA1214	rid3-5	ant3-5	ACC	HE858238	
	Aksaray-Helvadere	38.1999	34.2103	CA1215	ant11-6	ant3-5	ACC	HE858238	KID/ANT
				CA1216	rid3-5	rid3-5	ACC	ACC	
				CA1217	rid3-5	rid3-5	ACC	ACC	RID
				CA1218	rid3-5	rid3-5	ACC	ACC	
				CA1219	rid3-5	rid3-5	ACC	ACC	RID
	Aksaray-Melendez river	38.2988	34.2662	CA1220	ant1-4	rid3-5	ACC	ACC	RID/ANT
		30.2700	5112002	CA1221	ant1-1	ant11-7	ACC	ACC	ANT

				CA1222	rid2-1	ant11-7	ACC	ACC	RID/ANT
	Aksaray-Melendez river	38.2988	34.2662	CA1223	rid3-5	rid3-5	ACC	ACC	RID
				CA1224	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1205	rid3-5	rid3-5	ACC	ACC	
				CA1206	rid3-5	rid3-5	ACC	ACC	
				CA1207	rid3-5	rid3-5	ACC	ACC	PID
	Aksaray-Sultanhanı	38.2414	33.5448	CA1208	rid3-5	rid3-5	ACC	ACC	KID
				CA1209	rid3-5	rid3-5	ACC	ACC	
				CA1210	rid3-5	rid3-5	ACC	ACC	
				CA1211	rid3-5	ant11-7	ACC	ACC	RID/ANT
				CBCA03184	ant10-4	ant10-4	ACC	ACC	
Turkey				CBCA03186	ant7-3	ant10-4	ACC	ACC	
				CBCA03187	ant1-4	ant10-4	ACC	ACC	ANT
Turkey				CBCA03188	ant10-1	ant10-4	ACC	ACC	
				CBCA03189	ant10-1	ant10-1	ACC	ACC	
	Akşehir-Eber Lakes	38.4544	31.4546	CBCA03190	ant10-4	rid2-1	ACC	ACC	
				CA1715	ant10-4	rid2-1	ACC	ACC	RID/ANT
				CA1716	ant10-4	ant8-2	ACC	ACC	
				CA1717	ant1-6	ant10-4	ACC	ACC	ANT
				CA1718	ant10-4	ant10-4	ACC	ACC	ANI
				CA1719	ant8-3	ant10-4	ACC	ACC	
				MTHTCA07178	ant3-5	ant5-2	HE858238	ACC	ANT
	Alanva	36 6031	32 0694	CA1442	ant5-6	ant5-6	ACC	ACC	
	Alanya	50.0051	52.0074	CA1443	ant5-6	ant5-1	ACC	ACC	
				CA1444	ant5-6	ant3-5	ACC	HE858238	
	Amasya-Boğazköy Tersakan	40.7268	35.7702	CA909	ant11-7	ant11-7	ACC	ACC	ANT

				CA910	ant11-7	ant11-6	ACC	ACC	
	Amasya-Boğazköy Tersakan	40.7268	35.7702	CA911	ant3-5	ant11-7	HE858238	ACC	ANT
				CA912	ant3-2	ant11-7	ACC	ACC	
				CA902	ant11-7	ant11-7	ACC	ACC	ANT
	Amaswa Doğantana nond	40.6104	35 5860	CA903	ant3-5	rid2-1	HE858238	ACC	RID/ANT
	Amasya-Dogantepe pond	40.0104	33.3809	CA904	ant11-7	ant11-7	ACC	ACC	ANT
				CA905	rid2-1	ant11-7	ACC	ACC	RID/ANT
	Amasya-Suluoaya Vedikir dam	40 7786	35 5751	CA915	ant7-3	rid2-1	ACC	ACC	RID/ANT
	Amasya-Suluoava Teurkii dam	40.7780	55.5751	CA917	ant11-7	ant7-3	ACC	ACC	ANT
	Amasya-Yeşilırmak	40.6735	35.8528	CA907	ant11-7	ant11-7	ACC	ACC	ANT
		39.5793		KKOBCA06230	ant11-7	rid2-1	ACC	ACC	
Turkey			33.0453	KKOBCA06231	ant11-7	rid2-1	ACC	ACC	KID/ANT
	Ankara			KKOBCA06232	ant11-7	ant10-4	ACC	ACC	ANT
Turkey				KKOBCA06235	ant3-5	ant11-7	HE858238	ACC	AN
				KKOBCA06237	rid2-1	rid2-1	ACC	ACC	RID
	Antakya Samandağı	26.0742	35 0082	BCA31292	cil1-4	cil1-4	HE858225	HE858225	CIL
		50.0742	55.7762	BCA31293	cil1-4	cil1-7	HE858225	ACC	
				CBCASTO704	ant5-3	ant5-6	ACC	HE858230	
				CBCASTO705	ant1-2	ant3-4	ACC	ACC	
				CBCASTO706	ant4-3	ant8-3	ACC	ACC	
				CBCA0743	ant5-5	ant5-2	ACC	ACC	
	Antalya	37.2119	30.9415	HKHSBKCA07111	ant5-5	ant5-2	ACC	ACC	ANT
				HKHSBKCA07112	ant5-6	ant5-6	HE858230	HE858230	
				HKHSBKCA07113	ant5-6	ant5-6	HE858230	HE858230	
				HKHSBKCA07114	ant7-3	ant5-6	ACC	HE858230	
				HKHSBKCA07115	ant5-6	ant3-4	HE858230	ACC	

				HKHSBKCA07116	ant5-6	ant5-6	HE858230	HE858230	
				HKHSBKCA07117	ant5-5	ant5-2	ACC	ACC	
				HKHSBKCA07118	ant5-6	ant5-2	HE858230	ACC	
	Antolivo	27 2110	20.0415	HKHSBKCA07119	ant5-6	ant5-5	HE858230	ACC	ANT
	Antaiya	57.2119	30.9413	HKHSBKCA07120	ant2-1	ant10-1	ACC	ACC	ANI
				HKHSBKCA07121	ant5-6	ant5-6	HE858230	HE858230	
				HKHSBKCA07122	ant5-6	ant5-5	HE858230	ACC	
				HKHSBKCA07124	ant5-4	ant5-5	ACC	ACC	
				CA1433	ant5-6	ant5-1	HE858230	ACC	ANT
	Antalya-Anamur	36.0429	32 8078	CA1434	cil1-4	ant5-6	HE858225	HE858230	CIL/ANT
	Antarya-Antaniu	30.042)	52.0070	CA1435	ant5-6	ant5-2	HE858230	ACC	ANT
Turkey				CA1437	ant5-6	ant5-1	HE858230	ACC	AN
	Antalya-Kemer Kesmeboğaz river	36.5954		CA1863	ant8-3	ant8-3	ACC	ACC	
Turkey			30.5045	CA1865	ant8-3	ant5-6	ACC	HE858230	ANT
				CA1866	ant8-3	ant8-3	ACC	ACC	
	Antalya-Gazipaşa	36.1687	32.4467	CA1441	ant5-2	ant3-5	ACC	HE858238	ANT
	Antalya-Kumluca	36.3168	30.2512	CA1871	ant8-3	ant8-3	ACC	ACC	ANT
				CA732	ant11-7	rid2-5	ACC	ACC	
				CA733	ant6-1	rid3-7	ACC	ACC	RID/ANT
	Ardahan-Göle Serme Bridge	40.9087	42.5928	CA734	ant6-1	rid2-5	ACC	ACC	
				CA735	ant6-1	ant6-1	ACC	ACC	ANT
_				CA736	ant6-1	ant6-1	ACC	ACC	
				CA743	ant6-1	ant11-8	ACC	ACC	ANT
	Ardahan-Kura River	41.1569	42.8728	CA744	ant6-1	rid3-8	ACC	ACC	RID/ANT
				CA745	ant6-1	ant11-8	ACC	ACC	ANT
	Ardahan-Uzunova Village	41.0800	42.4991	CA739	rid3-8	rid3-8	ACC	ACC	RID

	Ardehen Lizunova Villago	41.0800	42 4001	CA740	rid3-8	rid3-8	ACC	ACC	RID
	Aldanan-Ozunova vinage	41.0800	42.4991	CA741	ant6-1	ant6-1	ACC	ACC	ANT
				CA746	ant6-1	ant6-1	ACC	ACC	
	Artvin Ardanuc Piver	41 1247	42.0668	CA747	ant6-1	ant6-1	ACC	ACC	ANT
	Artvin-Ardanuç Kiver	41.1247	42.0008	CA748	ant6-1	ant6-1	ACC	ACC	
				CA749	ant6-1	rid2-5	ACC	ACC	RID/ANT
				DC08241	rid2-5	rid2-5	ACC	ACC	RID
				DC08242	ant11-7	ant11-7	ACC	ACC	
	Artvin Borçka	41.3649	41.6920	CA757	ant6-1	ant11-7	ACC	ACC	ANT
				CA759	ant6-1	ant11-7	ACC	ACC	
				CA761	ant6-1	ant11-7	ACC	ACC	
Turkey	Artvin-Hopa Kemalpaşa	41.4870		CA764	ant11-7	rid2-5	ACC	ACC	
			41.5269	CA765	ant6-1	rid2-5	ACC	ACC	KID/AIT
				CA767	rid2-5	rid2-5	ACC	ACC	RID
				ATDC08191	rid2-5	rid2-5	ACC	ACC	RID
	Artvin Şavşat	41.3098	42.4837	ATDC08193	ant6-1	ant6-1	ACC	ACC	ANT
				ATDC08194	ant6-1	rid2-5	ACC	ACC	RID/ANT
				CA751	ant6-1	ant6-1	ACC	ACC	ANT
				CA752	ant6-1	rid3-8	ACC	ACC	
	Artvin-Yolüstü Village	41.1613	42.0639	CA753	ant6-1	rid2-5	ACC	ACC	RID/ANT
				CA754	ant6-1	ant6-1	ACC	ACC	ANT
				CA755	ant6-1	ant6-1	ACC	ACC	AN
				HS07165	ant3-5	ant3-5	HE858238	HE858238	
	Lake Avlan	36 5825	29 9482	HS07166	ant4-2	ant3-5	ACC	HE858238	ANT
	Lake Avian	50.5025	29.9482	HS07167	ant3-5	ant9-2	HE858238	ACC	
				HS07168	ant4-3	ant3-5	ACC	HE858238	

	Lake Avlan	36.5825	29.9482	HS07169	ant3-5	ant2-1	HE858238	ACC	ANT
				CA1800	ant9-2	ant10-1	ACC	ACC	
	Aydın-Azap lake	37.5844	27.447	CA1801	ant10-1	ant10-4	ACC	ACC	ANT
				CA1803	ant11-7	ant10-1	ACC	ACC	
	Aydın Byakali nond	27 7716	27 5741	CA1787	ant7-3	ant10-1	ACC	ACC	ANT
	Ayuni-Biyikan ponu	37.7710	27.3741	CA1789	ant7-3	ant7-3	ACC	ACC	ANI
				CA1796	ant8-1	ant4-3	ACC	ACC	
	Aydın-Menderes river	37.5476	27.2371	CA1797	ant9-2	ant9-2	ACC	ACC	ANT
				CA1798	ant10-1	ant10-4	ACC	ACC	
				CA1790	ant7-3	ant11-7	ACC	ACC	
Turkey	Aydın-Söke water channel	37.6607		CA1792	ant7-3	ant11-7	ACC	ACC	
			27.3087	CA1793	ant9-2	ant10-1	ACC	ACC	ANT
				CA1794	ant10-1	ant10-1	ACC	ACC	
Turkey				CA1795	ant4-3	ant4-3	ACC	ACC	
			27 6272	CA1618	ant11-7	ant11-7	ACC	ACC	ANT
	Balıkesir-Frdek	40 2968		CA1619	ant7-3	ant11-7	ACC	ACC	
	Duikesh Lidek	40.2900	27.0275	CA1620	ant11-7	ant11-7	ACC	ACC	
				CA1621	ant11-7	ant11-7	ACC	ACC	
				CA1601	ant11-7	ant10-1	ACC	ACC	
	Balıkesir-İkizceteneler dam	39 4807	27 9274	CA1602	ant11-5	ant11-7	ACC	ACC	ANT
	Bunkesh ikizetepelei dum	57.4007	21.9214	CA1603	ant11-7	ant11-7	ACC	ACC	21111
_				CA1605	ant11-7	ant11-7	ACC	ACC	
				CA1612	ant7-3	ant11-7	ACC	ACC	ANT
	Balıkesir-Manyas Kus Lake	40 2296	28.0461	CA1613	ant11-7	ant11-7	ACC	ACC	
	Sancon maryas reay lake	10.2290	20.0101	CA1615	ant7-2	ant11-7	ACC	ACC	
				CA1616	ant7-2	ant11-7	ACC	ACC	

				CA1608	ant11-7	ant11-7	ACC	ACC	
	Balıkesir-Pamukçu stream	39.5295	27.9091	CA1609	ant11-7	ant11-7	ACC	ACC	ANT
				CA1610	ant7-3	ant11-7	ACC	ACC	
				CA982	ant12-1	ant11-3	ACC	ACC	
				CA983	ant11-6	ant11-6	ACC	ACC	
				CA984	ant3-2	ant11-3	ACC	ACC	
	Bartın-Potbaşı river	41.6087	32.3727	CA985	ant11-6	ant12-1	ACC	ACC	ANT
				CA986	ant11-6	ant12-1	ACC	ACC	
				CA987	ant3-2	ant12-1	ACC	ACC	
				CA988	ant7-3	ant12-1	ACC	ACC	
		41.5721		CA989	ant11-6	ant12-1	ACC	ACC	
	Bartın-Bartın stream		32.3267	CA992	ant3-2	ant11-6	ACC	ACC	ANT
Turkey				CA993	ant11-7	ant11-6	ACC	ACC	
				CA995	ant10-1	ant7-3	ACC	ACC	
				CA1355	ant11-9	ant11-9	ACC	ACC	ANT
				CA1356	rid3-8	ant11-7	ACC	ACC	RID/ANT
	Batman-Dicle river	37 9177	41.0856	CA1357	rid3-8	rid3-8	ACC	ACC	RID
	Bathan-Dicie fiver	57.9177	41.0050	CA1358	rid3-8	ant11-7	ACC	ACC	
				CA1359	rid3-7	ant11-7	ACC	ACC	KID/AIVI
				CA1360	ant11-7	ant11-9	ACC	ACC	ANT
	Batman-Hasankeyf	37 7356	41 3022	CA1370	rid3-8	ant11-9	ACC	ACC	RID/ANT
		51.1550	41.5022	CA1371	rid3-7	ant11-7	ACC	ACC	
				CA1361	ant11-7	ant11-9	ACC	ACC	ANT
	Batman-Silvan road Carıklı village	38.0463	41,1834	CA1362	ant11-7	ant3-5	ACC	HE858238	ANI
		2010102	41.1834	CA1363	rid3-8	rid3-7	ACC	ACC	RID
				CA1364	rid3-8	rid3-7	ACC	ACC	

	Batman Silvan road Carikli village	38 0463	41 1834	CA1365	ant11-7	ant11-7	ACC	ACC	ANT
	Baunan-Sirvan toau Çarıklı vinage	58.0405	41.1054	CA1366	rid3-8	ant11-9	ACC	ACC	RID/ANT
	Bayburt-Camlık District	40 3009	40 2013	CA809	rid3-8	rid3-8	ACC	ACC	RID
	Baybart Çannık Disalet	40.5009	40.2015	CA812	rid3-8	rid2-5	ACC	ACC	KID
				CA802	ant11-7	ant11-7	ACC	ACC	ANT
	Bayburt-Gökçedere pond	40.1156	39.7526	CA804	ant3-5	ant11-7	HE858238	ACC	
				CA805	ant11-6	rid2-5	ACC	ACC	RID/ANT
				CA797	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Bayburt-Oruçbeyli pond	40.2316	40.0626	CA799	rid3-8	rid2-5	ACC	ACC	RID
				CA801	ant11-7	ant11-7	ACC	ACC	ANT
	Bayburt-Toki	40.2729	40.1343	CA808	rid3-8	ant11-7	ACC	ACC	RID/ANT
Turkey				MEFUCA42195	ant1-4	ant1-5	ACC	ACC	
				MEFUCA42196	ant1-4	ant1-5	ACC	ACC	
Turkey				MEFUCA42197	ant3-5	ant1-5	HE858238	ACC	ANT
				MEFUCA42198	ant1-5	ant10-4	ACC	ACC	
				CA1450	ant1-5	ant3-2	ACC	ACC	
	Lake Beyşehir	37.6802	31.7180	CA1451	ant1-4	ant1-5	ACC	ACC	
				CA1452	rid2-1	ant10-4	ACC	ACC	RID/ANT
				CA1453	ant1-5	ant3-2	ACC	ACC	ANT
				CA1454	ant1-4	ant10-4	ACC	ACC	
-				CA1456	ant1-5	rid2-1	ACC	ACC	RID/ANT
				CA1457	ant1-5	ant10-4	ACC	ACC	ANT
	Bilecik-Ciğdemlik	40.0646	30 3142	CA1657	ant7-3	ant7-2	ACC	ACC	ANT
	Direck-Çigucinik	+0.00+0	50.5172	CA1658	ant7-3	ant7-3	ACC	ACC	ANI
	Bilecik-İnhisar Sakarya river	40.0469	30.4124	CA1660	ant7-3	ant11-7	ACC	ACC	ANT
	Bilecik-İnhisar Sakarya river	40.0469	30.4124 -	CA1662	ant7-3	ant7-3	ACC	ACC	ANT

	Bilecik-İnhisar Sakarya river	40.0469	30.4124	CA1663	ant7-2	ant11-7	ACC	ACC	ANT
				MAYGCA11303	ant11-7	ant10-4	ACC	ACC	
				MAYGCA11304	ant7-3	ant11-7	ACC	ACC	ANT
	Bilecik Söğüt	39 7118	30.0070	MAYGCA11305	ant7-3	ant10-1	ACC	ACC	
	Blicelk Sögut	39.7110	30.0070	MAYGCA11306	ant11-7	rid2-1	ACC	ACC	RID/ANT
				CA1649	rid2-1	ant7-2	ACC	ACC	Kib//iitti
				CA1652	ant7-2	ant7-3	ACC	ACC	ANT
				CA570	rid3-8	ant11-8	ACC	ACC	RID/ANT ANT
	Bingöl-Çeltiksuyu stream	38.8411	40.5654	CA571	ant11-8	ant3-5	ACC	HE858238	
				CA572	ant3-5	ant3-5	HE858238	HE858238	
		39.0585	40.7976	CA582	ant11-7	rid3-7	ACC	ACC	
	Bingöl-Çobantaşı			CA583	rid3-7	ant11-9	ACC	ACC	RID/ANT
Turkey				CA584	rid3-7	ant3-5	ACC	HE858238	
				CA585	rid3-8	rid3-8	ACC	ACC	RID
				CA575	ant11-7	ant3-5	ACC	HE858238	ANT
	Bingöl-Saricicek Lake	38 8844	40 5890	CA576	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Billgor-Barlylyck Lake	30.0044	40.5670	CA578	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA579	ant3-5	ant3-5	HE858238	HE858238	2111
				CA586	rid3-8	ant11-9	ACC	ACC	
				CA587	ant11-7	rid3-8	ACC	ACC	
	Bingöl-Solhan Seref stream	38.9184	40.7859	CA588	rid3-8	ant3-5	ACC	HE858238	RID/ANT
				CA589	ant11-7	rid3-8	ACC	ACC	
				CA590	rid3-8	ant11-9	ACC	ACC	
-	Bitlis-Ağaçköprü District	38.3328	42.0098	CA630	rid3-8	rid3-8	ACC	ACC	RID
	Bitlis-Güroymak Water Channel	38.5962	42.0243	CA618	rid3-8	ant11-8	ACC	ACC	RID/ANT

				CA619	rid3-8	ant11-8	ACC	ACC	ANT
	Bitlis-Güroymak Water Channel	38.5962	42.0243	CA620	ant3-5	ant3-5	HE858238	HE858238	ANI
				CA621	rid3-8	rid3-8	ACC	ACC	RID
				CA623	rid3-7	ant11-9	ACC	ACC	RID/ANT
	Bitlis-Kemah Stream	38.4407	42.1447	CA624	ant11-9	ant11-9	ACC	ACC	ANT
				CA625	rid3-7	ant3-5	ACC	HE858238	RID/ANT
				CA631	ant11-8	ant3-5	ACC	HE858238	ANT
	Bitlis-Tatvan Cağlavan	38 4748	42 3084	CA632	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Bitlis-Tatvali Çaglayalı	38.4748	42.3084	CA634	ant11-9	ant11-8	ACC	ACC	ANT
				CA635	ant11-9	ant11-8	ACC	ACC	ANI
	Bolu-Gölköy			CA1520	ant11-7	ant10-4	ACC	ACC	ANT
Turkey		40.7116	31 5314	CA1521	ant7-2	ant7-2	ACC	ACC	ANI
		40.7110	51.5514	CA1522	ant7-2	rid1-1	ACC	ACC	RID/ANT
Turkey				CA1524	ant11-7	rid1-7	ACC	ACC	KID/ANT
			21 5162	CA1525	ant7-2	ant7-2	ACC	ACC	ANT
	Bolu-Karamanlar lake	40 7619		CA1526	ant7-3	ant7-3	ACC	ACC	
	Bolu-Karamamar lake	40.7019	51.5102	CA1527	ant11-7	ant3-2	ACC	ACC	
				CA1528	ant7-2	ant11-7	ACC	ACC	
				MTAECA1568	ant5-6	ant10-1	HE858230	ACC	
				MTAECA1569	ant3-5	ant10-1	HE858238	ACC	
				MTAECA1570	ant3-5	ant10-1	HE858238	ACC	
	Bucak	37 3500	30 5393	MTAECA1571	ant5-7	ant3-5	ACC	HE858238	ANT
	Bucak	57.5500	50.5575	MTAECA1572	ant3-5	ant2-1	HE858238	HE858238	AN
				MTAECA1573	ant5-6	ant10-1	HE858230	ACC	
				MTAECA1574	ant3-5	ant4-3	HE858238	HE858238	
				MTAECA1575	ant3-5	ant3-5	HE858238	HE858238	

	Bucak	37.3500	30.5393	MTAECA1576	ant3-5	ant5-6	HE858238	HE858230	ANT
	Lake Burdur	37.8371	30.3854	CBCAST1518	ant4-3	ant10-5	ACC	ACC	ANT
	Burdur-Çerçin dam	37.7603	30.4149	CA1883	ant3-2	ant3-2	ACC	ACC	ANT
	Burdur Karatas lake	37 3616	20.0860	CA1877	ant10-1	ant4-3	ACC	ACC	ANT
		37.3010	29.9809	CA1880	ant10-1	ant10-1	ACC	ACC	ANI
	Burdur-Varisli lake	37 5907	29.9562	CA1872	ant3-5	ant3-5	HE858238	HE858238	ANT
		37.3907	29.9502	CA1873	ant10-1	ant10-1	ACC	ACC	ANI
				CA1642	ant11-7	ant11-7	ACC	ACC	
	Bursa-Akçalar village	40.1771	28.7453	CA1644	ant7-3	ant10-4	ACC	ACC	ANT
				CA1646	ant7-2	ant10-1	ACC	ACC	
Turkey –				CA1622	ant11-7	ant10-1	ACC	ACC	
	Bursa-Boğaz	40.2821	28.4483	CA1623	ant7-3	ant11-7	ACC	ACC	ANT
				CA1625	ant7-3	ant11-7	ACC	ACC	
	Bursa-Gölvazı		28.6795	CA1636	ant7-2	ant11-7	ACC	ACC	
		40.1653		CA1637	ant7-2	ant11-7	ACC	ACC	ANT
	Duisu Goryuzi			CA1638	ant11-7	ant11-7	ACC	ACC	
				CA1640	ant7-3	ant11-7	ACC	ACC	
				CA1628	ant11-7	ant11-7	ACC	ACC	
	Bursa-Kemalnasa	40.0346	28 4104	CA1629	ant7-3	ant7-3	ACC	ACC	ANT
	Dursu Komulpuşu	10.0510	20.1101	CA1630	ant7-3	ant11-7	ACC	ACC	
				CA1634	ant11-7	ant11-7	ACC	ACC	
				CA1576	ant11-7	ant11-7	ACC	ACC	
				CA1577	ant11-7	ant11-7	ACC	ACC	
	Çanakkale-Karacaören	40.1884	26.4330	CA1578	ant11-7	ant11-7	ACC	ACC	ANT
				CA1580	ant7-3	ant11-7	ACC	ACC	
				CA1581	ant11-7	ant11-7	ACC	ACC	

				CA1590	ant11-7	ant11-7	ACC	ACC	
	Canaldrala Kanag	40.0805	26 2952	CA1591	ant7-3	ant11-7	ACC	ACC	ANT
	Çanakkale-Kepez	40.0893	20.3832	CA1592	ant11-7	ant11-7	ACC	ACC	ANI
				CA1595	ant11-7	ant11-7	ACC	ACC	
				CA1596	ant11-7	ant11-7	ACC	ACC	
				CA1597	ant7-3	ant11-7	ACC	ACC	
	Çanakkale-Batakova Menderes river	39.9930	26.2078	CA1598	ant7-3	ant11-7	ACC	ACC	ANT
				CA1599	ant7-3	ant11-7	ACC	ACC	
				CA1600	ant11-7	ant11-7	ACC	ACC	
			26.4855 -	CA1582	ant7-3	ant11-7	ACC	ACC	
	Canakkale Saricay	40.1393		CA1584	ant7-3	ant11-7	ACC	ACC	ANT
Turkey	Çanakkai¢-Sariçay			CA1585	ant11-7	ant11-7	ACC	ACC	ANI
				CA1586	ant7-3	ant11-7	ACC	ACC	
Turkey	Cankuri-Ansari stream		33.5393	CA1023	rid2-1	ant7-3	ACC	ACC	
		40.7018		CA1024	ant11-7	rid2-1	ACC	ACC	RID/ANT
	Çalıkıri-Apsalı sucalı			CA1025	ant11-7	rid2-1	ACC	ACC	
				CA1026	ant11-7	rid2-1	ACC	ACC	
				CA1028	ant11-7	ant11-6	ACC	ACC	ANT
	Çankırı-Aşağıyanlar district	40.5540	33.5817	CA1029	ant3-2	rid1-7	ACC	ACC	RID/ANT
				CA1030	ant11-6	ant3-2	ACC	ACC	ANT
				CA1033	ant11-7	ant3-2	ACC	ACC	
_	Çankırı-Terme river	40.4377	33.7431	CA1034	ant11-7	ant3-2	ACC	ACC	ANT
				CA1035	ant10-4	ant10-4	ACC	ACC	
	Cankuri-Yanraklı pond	40 7675	33 7641	CA1019	ant11-7	rid2-1	ACC	ACC	RID/ANT
	Çalıkır- rapiakii pond	40.7075	33.7641	CA1020	ant11-7	ant3-5	ACC	HE858238	ANT
	Çorum-Alaca stream	40.3371	35.0628	CA1041	ant11-7	ant11-7	ACC	ACC	ANT

	Çorum-Alaca stream	40.3371	35.0628	CA1042	ant11-7	ant11-6	ACC	ACC	
				CA1043	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1044	ant11-7	ant3-2	ACC	ACC	
	Çorum-Çomar dam	40.5856		CA1052	ant11-7	rid2-1	ACC	ACC	RID/ANT
			34.9998	CA1053	ant11-7	ant10-4	ACC	ACC	ANT
				CA1054	ant11-6	ant3-2	ACC	ACC	
	Corum Vakacık stream	40,6000	24.0117	CA1046	ant11-7	ant11-7	ACC	ACC	ANT
		40.0000	54.9117	CA1047	rid2-1	ant3-2	ACC	ACC	RID/ANT
			34.3789	CA1037	ant3-2	ant3-2	ACC	ACC	
	Çorum-Yılgınözü stream	40.4502		CA1038	ant11-7	ant11-6	ACC	ACC	ANT
				CA1039	ant11-7	ant3-5	ACC	HE858238	
	Dalaman	36.7138	28.7856	IS48307	ant11-7	ant9-2	ACC	ACC	ANT
Turkey				IS48308	ant4-3	ant4-3	ACC	ACC	
Turkey				IS48309	ant9-3	ant10-3	ACC	ACC	
				IS48310	ant4-3	ant10-1	ACC	ACC	
				IS48312	ant4-3	ant10-3	ACC	ACC	
		38.1532	28.8443	CA1815	ant9-2	ant7-3	ACC	ACC	
	Denizli-Derbent dam			CA1817	ant11-7	ant11-7	ACC	ACC	
				CA1819	ant10-4	ant4-3	ACC	ACC	
				CA1804	ant11-7	ant10-1	ACC	ACC	ANT
	Denizli-Emmiler district	37 6264	29 2087	CA1805	ant7-3	ant11-7	ACC	ACC	
		37.0204	29.2087	CA1806	ant7-3	ant11-7	ACC	ACC	
				CA1807	ant11-7	ant11-7	ACC	ACC	
				CA1810	ant11-7	ant11-7	ACC	ACC	ANT
	Denizli-İncirlipınar park	37.7623	29.0972	CA1812	ant7-3	ant7-3	ACC	ACC	
				CA1813	ant11-7	ant11-7	ACC	ACC	

Table 3.3.	(continued).
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	Denizli-Süleymanlı lake	38.0513	28.7708	CA1820	ant11-7	ant11-7	ACC	ACC	
				CA1821	ant10-4	ant8-1	ACC	ACC	
				CA1822	ant10-1	ant10-1	ACC	ACC	ANT
				CA1823	ant9-2	ant10-4	ACC	ACC	
				CA1824	ant11-7	ant4-3	ACC	ACC	
				CA1825	ant7-3	ant10-1	ACC	ACC	
	Divarbakır Batman road	37 8081	40.4055	CA1345	rid3-8	rid3-7	ACC	ACC	RID
	Diyarbakii-Batiliali Ibad	57.8081		CA1346	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1339	rid3-8	rid3-7	ACC	ACC	RID
	Diyarbakır-Devegeçidi	38.0576	40.0697	CA1340	ant11-7	ant11-7	ACC	ACC	ANT
				CA1341	rid3-8	ant11-7	ACC	ACC	RID/ANT
				CA1342	rid3-8	ant11-7	ACC	ACC	
				CA1343	rid3-7	ant3-5	ACC	HE858238	
Turkey				CA1344	rid3-8	rid3-7	ACC	ACC	RID
		37.8797	41.0217	CA1347	rid3-8	rid3-7	ACC	ACC	RID
				CA1348	rid3-8	ant11-7	ACC	ACC	RID/ANT
				CA1349	rid3-8	rid3-9	ACC	ACC	RID
	Diyarbakır-Dicle river			CA1350	ant11-7	ant11-7	ACC	ACC	ANT
				CA1352	rid3-8	rid3-7	ACC	ACC	RID
				CA1353	ant11-9	ant11-9	ACC	ACC	ANT
				CA1354	rid3-7	ant11-9	ACC	ACC	RID/ANT
				CA1331	rid3-7	ant11-7	ACC	ACC	RID/ANT
		37.6916		CA1332	rid3-8	ant11-7	ACC	ACC	
	Diyarbakır-Göksu Çınar		40.4473	CA1333	rid3-8	rid3-7	ACC	ACC	RID
				CA1334	rid3-8	rid3-7	ACC	ACC	
				CA1335	rid3-7	ant11-7	ACC	ACC	RID/ANT

	Divarbakır-Göksu Cınar	37.6916	40 4473	CA1337	rid3-8	rid3-8	ACC	ACC	RID
	Diyarbakii-Goksu Çillar		40.4473	CA1338	ant11-7	ant11-9	ACC	ACC	ANT
		40.8259		CA1530	ant7-2	ant7-2	ACC	ACC	ANT
				CA1531	ant7-3	ant11-7	ACC	ACC	
	Düzce-Asarsu		31.1840	CA1532	ant3-5	ant11-7	HE858238	ACC	
				CA1533	ant7-2	rid1-7	ACC	ACC	RID/ANT
				CA1534	ant7-2	ant11-7	ACC	ACC	ANT
			31.0404	CA1536	ant7-2	ant7-2	ACC	ACC	
	Düzce-Efteni lake	40.7573		CA1538	ant11-7	ant11-7	ACC	ACC	ANT
				CA1539	ant11-7	ant11-7	ACC	ACC	
	Edime	41.6857	26.4932	MACA22180	ant7-3	rid1-8	ACC	ACC	RID/ANT
Turker				MACA22181	ant7-3	ant10-4	ACC	ACC	ANT
				MACA22182	ant7-3	ant3-2	ACC	ACC	
Turkey				MACA22183	ant7-3	rid2-1	ACC	ACC	ANT/RID
				MA497	ant11-7	ant7-3	ACC	ACC	ANT
				MA500	ant11-7	ant11-7	ACC	ACC	
				MA501	rid3-5	rid3-5	ACC	ACC	RID
				MA503	ant11-7	rid1-3	ACC	ACC	RID/ANT
		38.1393	30.7588	CBCAST3234	ant5-6	ant10-4	HE858230	ACC	ANT
	Lake Eğirdir			CBCAST3238	ant4-3	ant8-3	ACC	ACC	
				CBCAST3239	ant3-2	ant1-2	ACC	ACC	
				CA535	rid3-8	rid3-7	ACC	ACC	RID
	Flaziă Cin	38.6819	39.0706	CA536	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Elazig - Cip			CA537	ant11-7	ant3-5	ACC	HE858238	ANT
				CA539	ant11-7	rid3-7	ACC	ACC	RID/ANT
	Elazığ - Sivrice	38.4665	39.2753	CA545	ant11-7	rid3-8	ACC	ACC	RID/ANT

	Elazığ - Sivrice	38.4665	39,2753	CA546	rid3 8	rid3 8	ACC	ACC	RID
	Erzincan - Türkmenoğlu village	39.6559	39.4996	CA520	rid2 9	rid2 5	ACC	ACC	KID
				CA520	1103-8	-: 12 0	ACC	ACC	RID
				CA521	nu3-8	103-8	ACC	ACC	
				CA522	ant11-/	ant3-5	ACC	HE858238	ANT
				CA523	ant11-7	ant3-5	ACC	HE858238	
				CA524	rid3-8	cil1-5	ACC	ACC	CIL/RID
				CA525	ant11-7	ant3-5	ACC	HE858238	
			39.6179	CA526	ant3-5	ant3-5	HE858238	HE858238	
	Erzincan - Ekşisu	39.7328		CA527	ant11-7	ant11-7	ACC	ACC	ANT
				CA528	ant11-7	ant3-5	ACC	HE858238	-
				CA529	ant11-7	ant11-7	ACC	ACC	
	Erzincan-Sakaltutan	39.8832	39.1954	CA530	ant11-7	ant11-7	ACC	ACC	ANT
Tustray				CA531	ant11-7	ant3-5	ACC	HE858238	
Turkey				CA532	ant11-7	ant3-5	ACC	HE858238	
				CA533	ant11-7	ant3-5	ACC	HE858238	
	Erzurum-Ilıca	39.8196	41.1521	CA696	ant11-7	rid3-8	ACC	ACC	RID/ANT
				CA697	rid3-8	ant3-5	ACC	HE858238	
				CA699	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA701	rid3-8	rid3-7	ACC	ACC	RID
				CA702	ant11-8	ant3-5	ACC	HE858238	ANT
				CA704	ant3-5	ant3-5	HE858238	HE858238	ANT
	Fraurum Desinler	39.9611	41 4000	CA706	rid3-8	rid3-8	ACC	ACC	RID
	Eizurum-Fasimer		41.4090	CA707	rid3-7	ant3-5	ACC	HE858238	
				CA708	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Erzurum-Soğuk Cermik	39.9899	41 3052	CA709	ant3-5	ant3-5	HE858238	HE858238	ANT
	Eizurum-Soguk Çennik		41.5052	CA710	rid3-8	rid3-8	ACC	ACC	RID

	Erzurum Soğuk Cermik	39.9899	41.3052	CA711	rid3-8	ant3-5	ACC	HE858238	PID/ANT	
	Erzurum-Söguk Çennik			CA712	rid3-7	ant3-5	ACC	HE858238	KID/AN1	
		39.8196	41.1521	CA689	rid3-8	rid3-8	ACC	ACC	RID	
				CA690	rid3-8	rid3-8	ACC	ACC		
	Erzurum-Teke stream			CA691	rid3-8	ant3-5	ACC	HE858238	RID/ANT	
				CA692	rid3-8	rid3-8	ACC	ACC	PID	
				CA693	rid3-8	rid3-8	ACC	ACC	KID	
	Erzurum-Verlisu Village	40.0425	41.1833	CA714	rid3-8	rid3-8	ACC	ACC	RID	
	Erzurum-Ternsu vinage	40.0425		CA715	rid3-8	ant3-5	ACC	HE858238	RID/ANT	
			30.6877	CA1672	ant7-3	ant10-1	ACC	ACC		
	Eskişehir-Alpu road	39.7825		CA1673	ant7-3	ant10-4	ACC	ACC	ANT	
				CA1674	ant7-2	ant11-7	ACC	ACC		
Turkey				CA1675	ant3-2	rid1-7	ACC	ACC	RID/ANT	
Turkey				CA1676	ant11-7	ant3-2	ACC	ACC	ANT	
	Eskişehir-Porsuk river 1	39.7744	30.4511	CA1677	ant11-7	ant11-7	ACC	ACC	ANT	
				CA1678	ant11-7	ant10-4	ACC	ACC		
				CA1679	ant11-7	ant10-4	ACC	ACC		
				CA1680	ant11-7	ant10-4	ACC	ACC		
			30.4275	CA1681	ant11-7	ant11-7	ACC	ACC	ANT	
	Eskişehir-Porsuk river 2	39.7139		CA1683	ant7-3	ant11-7	ACC	ACC		
				CA1684	ant10-4	ant10-4	ACC	ACC		
				CA1666	ant7-3	ant11-7	ACC	ACC	ANT	
		39.7082	30.5804	CA1667	ant7-2	ant11-7	ACC	ACC		
	Eskişehir-Sarısungur			CA1668	ant11-7	ant11-7	ACC	ACC		
				CA1669	ant9-1	ant11-7	ACC	ACC		
				CA1670	ant7-3	ant10-1	ACC	ACC		
	Giresun-Batlama stream	40.9067	38.3547	CA840	ant3-2	ant7-3	ACC	ACC	ANT	
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				CA854	ant3-2	ant7-3	ACC	ACC		
	Giresun-Bulancak Domuz stream	40.9454	38.1673	CA855	ant3-2	ant3-2	ACC	ACC	ANT	
				CA856	ant3-2	ant11-7	ACC	ACC		
				CA849	ant3-5	ant7-3	HE858238	ACC		
	Giragun Espiza Calivara Straam	40.0450	28 7004	CA850	ant7-3	ant7-3	ACC	ACC	ANT	
	Gnesun-Espiye Genvera Suean	40.9439	38.7224	CA852	ant3-2	ant3-2	ACC	ACC	ANI	
				CA853	ant11-7	ant11-6	ACC	ACC		
				OYCA70204	ant5-6	ant5-6	HE858230	HE858230		
				OYCA70205	ant5-6	ant5-6	HE858230	HE858230		
	Gödet Dam	37.1076		OYCA70207	ant11-7	ant11-7	ACC	ACC		
			33 2018	OYCA70208	ant5-6	ant11-7	HE858230	ACC	ANT	
Turkey			33.2918	CA1900	ant5-6	ant5-6	HE858230	HE858230		
				CA1901	ant1-4	ant11-7	ACC	ACC		
				CA1902	ant5-6	ant11-7	HE858230	ACC		
				CA1903	ant1-4	ant5-6	ACC	HE858230		
				CCYZCA20153	ant5-6	ant5-6	HE858230	HE858230		
	Gökninar Dam	37 7851	29 1306	CCYZCA20154	ant9-2	ant11-7	ACC	ACC	ANT	
	Gokphiai Dani	57.7651	29.1500	CCYZCA20155	ant5-6	ant5-6	HE858230	HE858230		
				CCYZCA20157	ant4-3	ant11-7	ACC	ACC		
	Gümüshane-Akhaha pond	40 1783	39 6521	CA818	ant3-5	ant11-7	HE858238	ACC	ANT	
		40.1705	57.0521	CA821	ant11-7	ant11-7	ACC	ACC	71111	
				CA822	ant11-7	rid2-5	ACC	ACC	RID/ANT	
	Gümüşhane-Kelkit river	40.1226	39.3737	CA824	ant3-5	ant3-5	HE858238	HE858238	ANT	
				CA826	ant3-5	ant3-5	HE858238	HE858238		
	Lake Işıklı	38.2350	29.9605	CBCA2047	ant11-10	ant11-7	ACC	ACC	ANT	

				CBCA2048	ant8-3	ant5-2	ACC	ACC	
				CBCA2049	ant2-1	ant10-4	ACC	ACC	
	Lake Işıklı	38.2350	29.9605	CBCA2050	ant1-4	ant1-5	ACC	ACC	ANT
				CBCA2051	ant1-4	ant3-2	ACC	ACC	
				CBCA2052	ant11-10	ant11-10	ACC	ACC	
				CA663	ant6-1	ant6-1	ACC	ACC	ANT
	Iğdır- Tuzluca Aras River	40.0356	43.6790	CA664	ant6-2	rid3-7	ACC	ACC	RID/ANT
				CA665	ant6-2	ant6-2	ACC	ACC	ANT
				CA667	ant11-8	ant11-8	ACC	ACC	_
				CA668	ant6-1	ant11-8	ACC	ACC	
	Iğdır- Water Channel	39.9513	44.0419	CA669	ant6-2	ant11-8	ACC	ACC	ANT
				CA670	ant11-8	ant11-8	ACC	ACC	
Turkey				CA671	ant11-8	ant11-8	ACC	ACC	
Turkey			35.9508	CA1949	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1950	cil1-4	cil1-4	HE858225	HE858225	
	Hatay-Deniz	36.0727		CA1951	cil1-4	cil1-4	HE858225	HE858225	
				CA1953	cil1-4	cil1-3	HE858225	ACC	
				CA1954	cil1-4	cil1-4	HE858225	HE858225	
				CA1403	cil1-4	cil1-1	HE858225	ACC	CII
				CA1404	cil1-4	cil1-4	HE858225	HE858225	СШ
	Hatay-Hassa	36 8//19	36 6504	CA1405	cil1-4	ant3-5	HE858225	HE858238	CIL/ANT
	Thury Thussa	50.0449	50.0504	CA1406	cil1-4	cil2-1	HE858225	ACC	CII
				CA1407	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1408	cil1-4	ant3-5	HE858225	HE858238	CIL/ANT
	Hatav-İskenderun Frzin	36 8887	36.1370	CA1416	cil1-4	cil2-1	HE858225	ACC	CIL
	Hatay-İskenderun Erzin	36.8887		CA1417	cil1-6	cil1-4	ACC	HE858225	

302

				CA1409	ant3-5	cil1-4	HE858238	HE858225	CIL/ANT
				CA1411	cil1-4	cil1-4	HE858225	HE858225	
	Hatay-İskenderun Sarıseki	36.6653	36.2157	CA1413	cil1-4	cil1-8	HE858225	ACC	CIL
				CA1414	cil1-4	cil1-4	HE858225	HE858225	
				CA1415	ant11-7	cil1-4	ACC	HE858225	CIL/ANT
				OBMB31342	cil3-1	cil3-1	ACC	ACC	CIL
	Hatay Kurkhan	26 4072	26 4522	OBMB31343	ant3-5	cil1-4	HE858238	HE858225	CIL/ANT
	Hatay Kirkhan	30.4973	50.4525	OBMB31345	cil1-4	cil1-4	HE858225	HE858225	CII
				OBMB31346	cil1-4	cil1-4	HE858225	HE858225	CIL
				OBMB31336	ant3-5	cil1-4	HE858238	HE858225	CII /ANT
	Hatay Reyhanlı	36.2379	36.5689	OBMB31338	ant3-5	cil1-4	HE858238	HE858225	CIL/AIVI
Turkey				OBMB31340	cil1-4	rid3-5	HE858225	ACC	CIL/RID
	Hatay Tahtalı Dam	36.8515	36 6861	OBMB31348	cil1-4	cil1-4	HE858225	HE858225	CIL
Turkey		50.8515	50.0801	OBMB31349	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1446	ant3-5	ant10-4	HE858238	ACC	
	Isparta-Bağıllı	38.1586	31.0825	CA1447	ant1-4	ant10-1	ACC	ACC	ANT
				CA1449	ant10-4	ant10-4	ACC	ACC	
	Isparta-Sevinçbey district	37.8748	30.7760	CA1885	ant4-3	ant3-2	ACC	ACC	ANT
				OK42203	cil1-4	ant1-4	HE858225	ACC	CIL/ANT
	İvriz	37.4408	34.1705	OK42221	ant1-4	ant1-2	ACC	ACC	ANT
				OK42222	ant1-5	ant11-7	ACC	ACC	AN
				CA1748	ant11-7	ant10-1	ACC	ACC	
				CA1749	ant11-7	ant11-7	ACC	ACC	
	İzmir-Çandarlı Bakırçay	38.9556	27.0100	CA1751	ant7-3	ant11-7	ACC	ACC	ANT
				CA1752	ant11-7	ant2-1	ACC	ACC	
				CA1753	ant11-7	ant11-7	ACC	ACC	

	Kahramanmaraş-Elbistan Gölpınar	38.1959	37.0636	CA1263	rid3-8	cil2-1	ACC	ACC	CIL/RID
				CA1264	cil2-1	ant3-5	ACC	HE858238	CIL/ANT
				CA1266	cil1-4	cil2-1	HE858225	ACC	СП
				CA1267	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1268	cil1-1	bed1-5	ACC	ACC	CIL/BED1
	Kahramanmaraş-Kumaşır lake	37.5102	36.8983	CA1269	cil1-4	cil3-1	HE858225	ACC	
				OB379	cil1-6	cil1-1	ACC	ACC	
				OB381	cil1-4	cil3-1	HE858225	ACC	CIL
				OB383	cil1-4	cil3-1	HE858225	ACC	
				OB384	cil1-4	cil1-4	HE858225	HE858225	
	Karabük-Cemaller Soğanlı river	41.1313	32.6816	CA965	ant11-3	ant11-6	ACC	ACC	
Turkey				CA966	ant10-4	ant11-6	ACC	ACC	
				CA967	ant11-7	ant7-2	ACC	ACC	ANT
				CA968	ant3-2	rid2-1	ACC	ACC	RID/ANT
				CA969	ant10-4	ant12-1	ACC	ACC	ANT
				CA970	ant10-4	rid1-7	ACC	ACC	RID/ANT
				CA971	ant3-2	ant10-4	ACC	ACC	ANT
				CA972	ant10-4	rid1-7	ACC	ACC	PID/ANT
				CA973	rid2-1	ant12-1	ACC	ACC	KID/ANT
				CA974	ant11-6	ant3-1	ACC	ACC	
				CA975	ant3-2	ant12-1	ACC	ACC	ANT
				CA976	rid2-1	ant12-1	ACC	ACC	RID/ANT
	Karabük-Safranbolu Araç river	41.2165	32.7331	CA977	ant3-2	ant12-1	ACC	ACC	
				CA978	ant11-6	ant12-1	ACC	ACC	ANT
				CA979	ant10-4	rid1-7	ACC	ACC	RID/ANT

				CBCAST328	ant3-2	ant8-3	ACC	ACC	
				CBCAST329	ant5-3	ant5-7	ACC	ACC	
				CBCAST3213	ant7-3	ant5-2	ACC	ACC	
				CBCA3241	ant5-6	ant5-6	ACC	HE858230	
	Karaaaäran Dama L II	27 4021	20,8702	CBCA3242	ant5-4	ant7-3	ACC	ACC	ANT
	Karacaolen Danis I-II	37.4031	30.8703	OBBKCA32103	ant3-5	ant5-2	HE858238	ACC	ANI
				OBBKCA32104	ant5-7	ant10-4	ACC	ACC	
				OBBKCA32106	ant7-3	ant5-6	ACC	HE858230	
				OBBKCA32107	ant10-1	ant5-7	ACC	ACC	
				OBBKCA32108	ant3-3	ant3-2	ACC	ACC	
				CA1909	ant5-6	ant5-6	HE858230	HE858230	
Turkey	Karaman-Belpınarı	36.8333	32.5803	CA1912	ant5-6	ant5-6	HE858230	HE858230	ANT
				CA1913	ant5-6	ant5-6	HE858230	HE858230	
				CA1906	ant5-6	ant11-7	HE858230	ACC	
	Karaman-Yeşildere	37.2047	33.4035	CA1907	ant11-7	ant11-7	ACC	ACC	ANT
				CA1908	ant5-6	ant11-7	HE858230	ACC	
				CA725	ant11-8	ant11-8	ACC	ACC	
				CA726	ant6-1	ant11-8	ACC	ACC	
	Kars-Arpaçay Akçalar	40.7735	43.2952	CA727	ant11-8	ant11-8	ACC	ACC	ANT
				CA729	ant11-8	ant11-8	ACC	ACC	
				CA730	ant6-1	ant11-8	ACC	ACC	
				CA717	ant11-8	ant11-8	ACC	ACC	
	Kars-Digor Pazarcık	40 5189	43 2690	CA718	ant11-8	ant11-8	ACC	ACC	ANT
	Kars Digor i azarcık	40.5109	+5.2070	CA719	ant6-2	ant11-8	ACC	ACC	21111
				CA720	ant11-8	ant11-8	ACC	ACC	
	Kars-Selim	40.4702	42.7904	CA721	ant11-8	ant11-8	ACC	ACC	ANT

				CA722	ant11-8	ant11-8	ACC	ACC	ANT
	Kars-Selim	40.4702	42.7904	CA723	ant11-7	ant11-8	ACC	ACC	AINI
				CA724	ant11-8	ant11-8	ACC	ACC	ANT
				CA959	ant11-3	ant11-6	ACC	ACC	ANT
	Kastamonu Aras Diver	41 2380	33 3774	CA960	ant3-2	ant11-7	ACC	ACC	AINI
	Kastanionu-Araç Kiver	41.2309	33.3224	CA963	ant3-5	rid2-1	HE858238	ACC	RID/ANT
				CA964	ant11-6	ant12-1	ACC	ACC	ANT
				CA952	rid2-1	ant12-1	ACC	ACC	RID/ANT
	Kastamonu-Beyler dam	41.6903	33.8119	CA954	ant3-2	ant3-2	ACC	ACC	ANT
				CA955	ant11-7	ant11-7	ACC	ACC	AINI
				CA939	ant7-2	rid2-1	ACC	ACC	RID/ANT
Turkey	Kastamonu-Karaçomak dam	41.2852	22 7272	CA941	ant3-2	ant12-1	ACC	ACC	ANT
			33.1312	CA942	ant3-2	ant3-2	ACC	ACC	
				CA943	ant3-1	ant3-2	ACC	ACC	
				CA945	ant11-7	ant11-7	ACC	ACC	ANT
				CA946	ant3-2	ant3-2	ACC	ACC	AINI
	Kastamonu-Taşlık Dam	41.4018	33.6963	CA947	ant7-3	rid2-1	ACC	ACC	RID/ANT
				CA948	ant3-2	ant11-6	ACC	ACC	
				CA950	ant3-2	ant11-6	ACC	ACC	AINI
	Vas	26 2760	20.6820	CA07217	ant4-4	ant10-1	ACC	ACC	ANT
	Kaş	30.2709	29.0839	CA07218	ant4-4	ant10-1	ACC	ACC	AINI
-	Kawari	29 7565	25 4220	DCA38315	ant11-6	rid2-1	ACC	ACC	PID/ANT
	Kaysen	38.7505	33.4229	YC38445	rid3-5	ant11-7	ACC	ACC	KID/ANT
				CA1151	rid2-1	ant3-2	ACC	ACC	RID/ANT
	Kayseri-Engir lake	38.8111	35.5915	CA1153	rid3-8	rid3-8	ACC	ACC	RID
				CA1154	ant11-7	ant3-5	ACC	HE858238	ANT

	Kaysari Engir laka	39 9111	35 5015	CA1155	ant11-7	ant3-5	ACC	HE858238	ANT
	Kaysen-Engir lake	56.6111	33.3913	CA1156	ant11-7	ant3-2	ACC	ACC	ANI
				CA1468	ant3-5	ant11-7	HE858238	ACC	
				CA1469	ant3-5	ant11-7	HE858238	ACC	
	Kayseri-Pınarbaşı	38.6618	36.104	CA1471	ant11-7	ant3-2	ACC	ACC	ANT
				CA1472	ant11-7	ant11-7	ACC	ACC	
				CA1473	ant11-7	ant11-7	ACC	ACC	
				CA1157	cil1-7	ant3-5	ACC	HE858238	CIL/ANT
				CA1158	ant3-5	rid3-8	HE858238	ACC	RID/ANT
				CA1159	cil1-4	ant3-5	HE858225	HE858238	CIL/ANT
	Kaycari Sultançazlığı	38.3897	35 3657	CA1160	cil1-7	rid3-6	ACC	ACC	CIL/RID
	Kaysen-Sunansazingi		33.3037	CA1161	ant1-4	ant1-4	ACC	ACC	ANT
Turkey			-	CA1162	cil1-4	rid3-6	HE858225	ACC	CIL/RID
Turkey				CA1163	cil1-4	ant3-5	HE858225	HE858238	CII /ANT
				CA1164	cil1-7	ant3-5	ACC	HE858238	CIL/ANI
				CA1165	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1166	cil1-4	ant11-7	HE858225	ACC	CIL/ANT
	Kawari Vahyalı	38 13/8	35 3641	CA1167	ant3-5	ant3-5	HE858238	HE858238	ANT
	Kaysen-Tanyan	56.1546	55.5041	CA1168	cil1-4	ant3-5	HE858225	HE858238	CII /ANT
				CA1169	cil1-7	ant3-5	ACC	HE858238	CIL/ANI
				CA1170	cil1-4	rid3-6	HE858225	ACC	CIL/RID
				CA1171	ant11-7	ant3-5	ACC	HE858238	
				CA1172	ant11-7	ant3-5	ACC	HE858238	ANT
	Kayseri-Zamantı Water	38.0345	35.5407	CA1173	ant11-7	ant11-7	ACC	ACC	
				CA1174	ant11-7	ant3-5	ACC	HE858238	
				CA1175	ant11-7	ant11-7	ACC	ACC	

				CA1176	ant11-7	ant11-7	ACC	ACC	
				CA1177	ant11-7	ant11-7	ACC	ACC	
	Kayseri-Zamantı Water	38.0345	35.5407	CA1178	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1179	ant11-6	ant11-7	ACC	ACC	
				CA1180	ant11-7	ant11-7	ACC	ACC	
				MHSACA15137	ant3-5	ant3-5	HE858238	HE858238	
				MHSACA15138	ant3-5	ant9-2	HE858238	ACC	
	Kemer	37.4627	30.1118	MHSACA15139	ant8-3	ant10-1	ACC	ACC	ANT
				MHSACA15140	ant3-6	ant3-6	ACC	ACC	
				MHSACA15142	ant2-1	ant4-3	ACC	ACC	
	Kemer	37.4627	30.1118	MHSACA15143	ant4-3	ant10-5	ACC	ACC	ANT
				CA1238	ant11-7	ant11-7	ACC	ACC	ANT
Turkey	Kırıkkale-Hasandede Kızılırmak	38.7406	33.4879	CA1239	ant11-7	ant3-2	ACC	ACC	21111
				CA1242	rid2-1	ant3-2	ACC	ACC	RID/ANT
			33.4146	CA1244	ant11-7	ant3-5	ACC	HE858238	
	Kırıkkale-Yahşiyan Kızılırmak	39.8861		CA1245	ant11-7	ant10-4	ACC	ACC	ANT
				CA1247	ant11-7	ant7-3	ACC	ACC	
				CA1488	ant11-7	rid1-8	ACC	ACC	
				CA1489	ant11-7	rid1-2	ACC	ACC	KID/AIVI
	Kurklareli-Eriklice stream	41 7576	27 1814	CA1490	ant7-3	ant11-7	ACC	ACC	ANT
	Kirklaren-Erikhee sireani	41.7570	27.1014	CA1491	ant11-7	rid1-7	ACC	ACC	RID/ANT
				CA1492	ant7-3	ant11-7	ACC	ACC	ANT
				CA1493	rid3-5	rid1-8	ACC	ACC	RID
				CA1494	ant7-2	rid2-1	ACC	ACC	RID/ANT
	Kırklareli-İnece stream	41.6836	27.0731	CA1495	ant7-3	rid2-6	ACC	ACC	
				CA1496	ant11-7	ant3-2	ACC	ACC	ANT

	Kurklarali İnaaa straam	41 6926	27.0721	CA1497	ant11-7	ant11-7	ACC	ACC	ANT
		41.0850	27.0731	CA1498	ant7-2	ant10-4	ACC	ACC	ANI
				CA1499	rid3-5	ant11-7	ACC	ACC	
	Kurklarali Karakaa nond	41 7917	27 2171	CA1500	ant11-7	rid1-2	ACC	ACC	RID/ANT
	Kirkiaren-Karakoç ponu	41.7017	27.2171	CA1501	ant7-3	ant11-7	ACC	ACC	ANT
				CA1502	ant11-7	ant11-7	ACC	ACC	ANT
				CA1484	ant11-7	rid1-8	ACC	ACC	RID/ANT
	Kırklareli-Sevtandere	41 7104	27 2627	CA1485	ant7-3	rid1-8	ACC	ACC	KID/ANT
	Kirklaren-şeytandere	41.7104	27.2027	CA1486	ant11-7	ant11-7	ACC	ACC	ANT
				CA1487	ant3-2	rid2-7	ACC	ACC	RID/ANT
				CA1141	rid2-1	ant11-7	ACC	ACC	RID/ANT
	Kırşehir-Çuvın	39.3168	34.1238	CA1142	ant7-3	ant3-5	ACC	HE858238	ANT
Turkey				CA1145	ant11-6	rid2-1	ACC	ACC	RID/ANT
Turkey —	Kusehir-Güneykent		34.1600	CA1128	rid2-1	rid2-1	ACC	ACC	RID
		39.0988		CA1129	ant11-7	rid2-1	ACC	ACC	RID/ANT
	Kinşenir Guneykent	37.0700		CA1130	ant11-6	rid2-1	ACC	ACC	
				CA1131	ant11-7	ant11-7	ACC	ACC	ANT
				CA1146	rid2-1	ant7-3	ACC	ACC	RID/ANT
	Kırsehir-Kışlanınar	39 2262	34 1338	CA1147	ant11-7	rid2-1	ACC	ACC	KID// III II
	Kişemi Kişiapinar	37.2202	54.1550	CA1148	rid2-1	ant3-5	ACC	HE858238	RID/ANT
				CA1149	ant11-7	ant10-4	ACC	ACC	ANT
				CA1135	rid2-1	ant3-5	ACC	HE858238	RID/ANT
	Kırşehir-Sıddıklı	39.1007	33.9073	CA1136	ant11-7	ant11-7	ACC	ACC	ANT
				CA1138	ant11-7	ant3-2	ACC	ACC	AINI
-	Kilis	36 7641	37.2540 —	OB79322	cil1-4	cil1-4	HE858225	HE858225	CIL
	Kilis	36.7641		OB79326	rid3-8	cil1-4	ACC	HE858225	CIL/RID

				OBMB79327	cil1-4	cil1-4	HE858225	HE858225	CIL
	Kilis Cogionton road	26.9106	27 2005	OBMB79328	cil1-4	rid3-5	HE858225	HE858238	CIL/RID
	Kins-Gaziantep road	50.8100	37.3093	OBMB79329	rid3-8	ant11-7	ACC	ACC	RID/ANT
				OBMB79330	cil1-4	rid3-5	HE858225	ACC	CIL/RID
				CA1395	ant3-5	cil1-4	HE858238	HE858225	CIL/ANT
				CA1396	ant3-5	ant3-5	HE858238	HE858238	ANT
	Kilis-Polateli	36.7853	37.0588	CA1397	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1398	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA1400	ant11-7	cil1-4	ACC	HE858225	CIL/ANT
				CA1390	ant3-5	cil1-4	HE858238	HE858225	CIL/ANT
	Kilis Täreli Capalı road	36 85/10	27.2414	CA1391	rid3-7	ant3-5	ACC	HE858238	RID/ANT
Turkey -	Kins-Toten Çapan toau	50.8549	37.3414	CA1393	rid3-8	cil1-4	ACC	HE858225	CIL/RID
				CA1394	ant11-7	cil1-4	ACC	HE858225	CIL/ANT
Turkey	Kilis-Üçpınar	36 8660	36.9328	CA1961	cil1-4	cil1-4	HE858225	HE858225	СП
		30.8009		CA1964	cil1-1	cil1-1	ACC	ACC	CIL
				CA1463	ant11-7	ant10-4	ACC	ACC	ANT
	Konya-Mehmetali dam	37.2124	32.6354	CA1465	ant5-6	ant3-5	HE858230	HE858238	
				CA1466	ant1-5	ant1-5	ACC	ACC	
	Korkuteli	36 9918	29 5279	VDBKCA07125	ant3-5	ant2-1	HE858238	ACC	ANT
		50.5510	29.3219	VDBKCA07130	ant3-5	ant3-5	HE858238	HE858238	71111
				CBCAST3215	ant5-7	ant3-2	ACC	ACC	
				OAOBBKCA1593	ant5-6	ant2-1	HE858230	ACC	
	Koyada	37 6325	30 8641	OAOBBKCA1596	ant4-3	ant2-1	ACC	ACC	ANT
		51.0525	50.0041	OAOBBKCA1597	ant8-3	ant5-2	ACC	ACC	ANI
				OAOBBKCA1598	ant1-5	ant7-3	ACC	ACC	
				OAOBBKCA1599	ant5-4	ant5-7	ACC	ACC	

	Koyada	37 6325	30 8641	OAOBBKCA1593101	ant5-6	ant10-1	HE858230	ACC	ANT
	Kovaua	37.0323	30.8041	OAOBBKCA1593102	ant10-1	ant10-1	ACC	ACC	ANI
				CA1685	ant10-4	ant8-3	ACC	ACC	
	Kütahya-Enne dam	39.4688	29.8657	CA1686	ant1-6	ant10-4	ACC	ACC	ANT
				CA1687	ant7-3	ant7-3	ACC	ACC	
				CA1697	ant11-7	ant10-4	ACC	ACC	
	Kütahva-Kocak stream	39 3090	29 9753	CA1698	ant7-3	ant10-4	ACC	ACC	ANT
	Kutanya-Koçak sucam	37.3070	29.9133	CA1699	ant11-7	ant10-4	ACC	ACC	
				CA1701	ant11-7	ant11-7	ACC	ACC	
				CA1705	ant7-3	ant10-1	ACC	ACC	
	Kütahya-Porsuk dam	39.5740	30.0918	CA1707	ant7-3	ant10-4	ACC	ACC	ANT
Turkey				CA1708	ant10-4	ant10-4	ACC	ACC	
	Kütahya-Söğütözü stream	39.3374	29.9342 -	CA1690	ant10-4	ant10-4	ACC	ACC	
Turkey				CA1691	ant10-1	ant10-4	ACC	ACC	ANT
				CA1692	ant10-4	ant10-4	ACC	ACC	
				CA1694	ant10-1	ant10-4	ACC	ACC	
	Malatya	38.4276	38.3505	SO44286	ant11-7	ant11-7	ACC	ACC	ANT
				CA552	ant11-9	ant3-5	ACC	HE858238	
	Malatya-Elemendik lake	38.3229	38.1551	CA554	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA555	ant3-5	ant3-5	HE858238	HE858238	
				CA547	ant11-7	rid3-8	ACC	ACC	RID/ANT
_	Malatya-Karakaya dam	38.4882	38.3499	CA548	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA551	rid3-8	ant11-9	ACC	ACC	RID/ANT
				CA558	ant11-7	ant11-7	ACC	ACC	ANT
	Malatya-near Sultansuyu dam	38.3121	38.0434	CA559	rid3-8	rid3-8	ACC	ACC	RID
		50.5121		CA560	rid3-8	ant3-5	ACC	HE858238	RID/ANT

	Malatya-near Sultansuyu dam	38.3121	38.0434	CA561	ant11-9	ant11-7	ACC	ACC	ANT
				CA563	ant3-5	ant3-5	HE858238	HE858238	ANT
	Malatya-Oluklu district	38.2406	37.9964	CA564	rid3-8	ant11-9	ACC	ACC	PID/ANT
				CA565	rid3-8	ant3-5	ACC	HE858238	KID/ANT
				OA45274	ant7-3	ant10-4	ACC	ACC	
				OA45275	ant11-7	ant11-7	ACC	ACC	
				OA45276	ant4-3	ant10-1	ACC	ACC	
	Manisa	38 6110	27 1183	OA45278	ant7-3	ant10-1	ACC	ACC	ANT
	Wallisa	58.0110	27.4463	OA45279	ant11-7	ant10-4	ACC	ACC	
				OA45280	ant7-3	ant4-3	ACC	ACC	
				OA45281	ant10-4	ant10-4	ACC	ACC	
Turkey				OA45282	ant10-4	ant10-4	ACC	ACC	
	Manisa City Forest	38.6021		CA1764	ant7-3	ant11-7	ACC	ACC	
Turkey			27.3867	CA1766	ant10-1	ant10-4	ACC	ACC	ANT
				CA1768	ant7-3	ant7-3	ACC	ACC	
				CA1780	ant7-3	ant10-1	ACC	ACC	ANT
	Manisa-Köseler pond	38.8460	27.2006	CA1781	ant11-7	ant11-7	ACC	ACC	
				CA1784	ant7-3	ant7-3	ACC	ACC	
				CA1774	ant11-7	ant11-7	ACC	ACC	
	Manisa-Örselli nond	38 8608	27 2620	CA1776	ant11-7	ant11-7	ACC	ACC	ANT
	Wallisa-Orsell polici	56.6076	27.2020	CA1777	ant11-7	ant11-7	ACC	ACC	
				CA1778	ant7-3	ant11-7	ACC	ACC	
				CA1769	ant7-3	ant4-3	ACC	ACC	1
	Manisa-Siyekli pond	38 7886	27 2559	CA1770	ant7-3	ant11-7	ACC	ACC	ANT
	Manisa-Biyokn pond	50.7000	21.2337	CA1771	ant7-3	ant11-7	ACC	ACC	
				CA1772	ant11-7	ant10-1	ACC	ACC	

	Manisa-Siyekli pond	38.7886	27.2559	CA1773	ant7-3	ant11-7	ACC	ACC	ANT
	Mardin-Bülbül village	37.3219	40.8364	CA1319	rid3-8	rid3-8	ACC	ACC	RID
				CA1313	rid3-8	rid3-8	ACC	ACC	RID
	Mardin-Kızıltepe	37.1970	40.2767	CA1314	rid3-8	rid3-8	ACC	ACC	KID
				CA1315	ant11-9	ant11-9	ACC	ACC	ANT
				CA1326	ant11-9	ant11-9	ACC	ACC	ANT
				CA1327	rid3-8	ant11-7	ACC	ACC	RID/ANT
	Mardin-Mazıdağ	37.4622	40.6169	CA1328	rid3-7	ant3-5	ACC	HE858238	
				CA1329	rid3-8	ant11-7	ACC	ACC	KID/ANT
				CA1330	rid3-8	ant3-5	ACC	HE858238	
				CA1320	rid3-8	rid3-7	ACC	ACC	RID
Turkey	Mardin-Savur road	37.4406	40.8553	CA1321	ant3-5	ant3-5	HE858238	HE858238	ANT
				CA1322	ant11-7	ant3-5	ACC	HE858238	
Turkey				CA1323	rid3-8	ant11-7	ACC	ACC	RID/ANT
				CA1324	rid3-8	rid3-8	ACC	ACC	RID
				AKCA48209	ant9-2	ant4-1	ACC	ACC	
	Marmaris	36.8467	28 2879	AKCA48210	ant10-2	ant9-2	ACC	ACC	
	Wannans	50.0407	20.2077	AKCA48211	ant7-3	ant4-3	ACC	ACC	AN
				AKCA48212	ant9-2	ant4-3	ACC	ACC	
	Mersin	36.7969	34.6019	SOCA33226	cil1-4	cil2-1	HE858225	ACC	CIL
				CA1429	cil1-5	cil1-5	ACC	ACC	
	Mersin-Erdemli	36.5930	34.2873	CA1430	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1431	cil1-4	ant5-6	HE858225	HE858230	CIL/ANT
	Mersin-Mut Bice stream	36.5896	33.2850	CA1915	ant5-6	ant5-6	HE858230	HE858230	ANT
	Mersin-Mut Bucakışla stream	36 6313	33 3675	CA1916	ant5-6	ant5-6	HE858230	HE858230) ANT
	Mersin-Mut Bucakışla stream	36.6313	33.3675	CA1917	ant5-6	ant5-6	HE858230	HE858230	

	Marsin Mut Bucakışla stream	36 6313	33 3675	CA1919	ant5-6	ant5-6	HE858230	HE858230	ANT
	Mersiii-Mut Bucakişia su'calli	30.0313	55.5075	CA1921	ant5-6	ant5-6	HE858230	HE858230	ANI
				CA1424	cil2-2	cil1-8	ACC	ACC	
				CA1425	cil1-6	cil1-6	ACC	ACC	CIL
	Mersin-Yenice	36.9673	35.0305	CA1426	cil1-6	cil1-2	ACC	ACC	
				CA1427	cil1-4	ant11-7	HE858225	ACC	CIL/ANT
				CA1428	cil1-1	cil1-8	ACC	ACC	CIL
				CA1845	ant10-1	ant4-3	ACC	ACC	
				CA1846	ant11-7	ant4-3	ACC	ACC	
	Muğla-Dalaman Tersakan	36.7794	28.8252	CA1847	ant9-2	ant4-3	ACC	ACC	ANT
				CA1849	ant11-7	ant4-3	ACC	ACC	
Turkey				CA1850	ant10-1	ant3-5	ACC	HE858238	
	Muğla-Girdev plateau	36.7007		CA1851	ant10-1	ant2-1	ACC	ACC	
Turkey			29.6509	CA1852	ant2-1	ant2-1	ACC	ACC	
				CA1853	ant2-1	ant10-1	ACC	ACC	ANT
				CA1857	ant3-5	ant3-5	HE858238	HE858238	
				CA1861	ant3-5	ant2-1	HE858238	ACC	
				CA1840	ant10-1	ant3-5	ACC	HE858238	
	Muğla-Kövceğiz Zaferler village	36 9718	28 6299	CA1841	ant9-2	ant11-7	ACC	ACC	ANT
		50.9710	20.0277	CA1842	ant9-2	ant10-1	ACC	ACC	21111
				CA1844	ant10-1	ant4-3	ACC	ACC	
				CA1836	ant9-2	ant10-1	ACC	ACC	
	Muğla-Ula Nannan stream	37.0129	28 5105	CA1837	ant4-3	ant10-1	ACC	ACC	ANT
		57.0127	20.5105	CA1838	ant11-7	ant11-7	ACC	ACC	AN1
				CA1839	ant11-7	ant10-1	ACC	ACC	
	Muğla-Yatağan Dipsiz stream	37.3750	28.0916	CA1826	ant7-3	ant11-7	ACC	ACC	ANT

				OBMBCA46365	cil1-4	cil1-4	HE858225	HE858225	CIL
				OBMBCA46366	cil1-4	ant3-5	HE858225	HE858238	CIL /ANT
				OBMBCA46367	cil1-4	ant11-7	HE858225	ACC	CIL/ANI
	Narlı	37.3322	37.0425	OBMBCA46369	cil1-4	cil1-4	HE858225	HE858225	
				OBMBCA46370	cil1-4	cil1-4	HE858225	HE858225	CII
				OBMBCA46371	cil1-1	cil1-1	ACC	ACC	CIL
				OBMBCA46372	cil1-4	cil1-6	HE858225	ACC	
				VD07132	ant5-6	ant5-5	HE858230	ACC	
	National Observatory	36 8293	30 3407	VD07134	ant5-6	ant5-5	HE858230	ACC	ANT
	National Observatory	50.8275	50.5407	VD07135	ant5-6	ant5-5	HE858230	ACC	
				VD07136	ant5-6	ant5-5	HE858230	ACC	
Turkey —	Nevşehir-Avanos Kızılırmak	38 7174	34.8575	CA1234	rid2-1	rid2-1	ACC	ACC	RID
	Nevşenii-Avanos Kızınınak	50.7174	51.0575	CA1235	ant3-5	rid3-8	HE858238	ACC	RID/ANT
Turkey	Nevşehir-Nar Ilıca river	38.6383	34.7089	CA1229	ant11-7	rid2-1	ACC	ACC	RID/ANT
			54.7009	CA1230	ant11-7	ant11-7	ACC	ACC	ANT
	Nevsehir-Gülsehir Kızılırmak	28 7561	24 6512	CA1236	rid2-1	ant3-5	ACC	HE858238	RID/ANT
		50.7504	54.0515	CA1237	ant11-7	rid2-1	ACC	ACC	RID// IIII
				CA1231	ant11-7	rid2-1	ACC	ACC	RID/ANT
	Nevşehir-Ürgüp	38.5741	34.9157	CA1232	ant11-7	rid2-1	ACC	ACC	RID// IIII
				CA1233	ant11-7	ant11-7	ACC	ACC	ANT
				CA1181	ant3-5	rid3-6	HE858238	ACC	RID/ANT
				CA1182	ant11-7	rid3-6	ACC	ACC	RID// IIII
	Niğde-Camardı	37 9608	34 3572	CA1183	cil1-4	ant3-5	HE858225	HE858238	CIL/ANT
	ingue çunurur	51.9000	54.5572	CA1185	ant3-5	rid3-6	HE858238	ACC	RID/ANT
				CA1186	cil1-4	rid2-1	HE858225	ACC	CIL/ANT
				CA1187	ant11-7	rid3-6	ACC	ACC	RID/ANT

				CA1195	rid3-5	rid3-5	ACC	ACC	RID
				CA1196	ant11-7	ant11-7	ACC	ACC	ANT
				CA1197	rid3-5	rid3-5	ACC	ACC	RID
				CA1198	cil1-4	cil2-1	HE858225	ACC	CIL
	Niăda Cifflik	28 1770	24 4620	CA1199	rid3-5	rid3-5	ACC	ACC	RID
	Nigue-Çinnik	56.1779	54.4029	CA1200	ant11-7	rid3-5	ACC	ACC	
				CA1201	ant1-4	rid3-5	ACC	ACC	RID/AIVI
				CA1202	ant11-7	ant11-7	ACC	ACC	ANT
				CA1203	rid3-5	rid3-5	ACC	HE858238	RID
				CA1204	ant11-7	rid3-5	ACC	HE858238	RID/ANT
	Niğde-15km to Ulukışla	37.6195	34.5366	CA1193	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA858	ant11-7	ant7-3	ACC	ACC	
Turkey	Ordu-Civil Stream	40.9733	37.9030	CA860	ant3-2	ant11-7	ACC	ACC	ANT
				CA861	ant11-7	ant7-3	ACC	ACC	
	Ordu-Civil Stream	40.9733	37.9030	CA862	ant11-7	ant7-3	ACC	ACC	ANT
				CA871	ant3-2	ant11-7	ACC	ACC	
	Ordu Estas	41.0752	27 4412	CA872	ant11-7	ant11-1	ACC	ACC	
	Ordu-Patsa	41.0733	37.4415	CA873	ant3-2	ant11-7	ACC	ACC	ANI
				CA874	ant3-2	ant11-7	ACC	ACC	
				CA867	ant11-7	ant11-7	ACC	ACC	
	Ordu-Perşembe	41.0251	37.8032	CA869	ant3-5	ant11-6	HE858238	ACC	ANT
				CA870	ant3-5	ant11-6	HE858238	ACC	
				CA1933	cil1-4	cil2-1	HE858225	ACC	
	Osmaniye-Aslantaş dam	37.2769	36.2744	CA1935	cil1-4	cil1-6	HE858225	ACC	CIL
				CA1936	cil1-4	cil2-1	HE858225	ACC	
	Osmaniye-Boru river	37.1780	36.4856	CA1943	cil1-4	cil1-8	HE858225	ACC	CIL

	Osmaniya Poru rivar	27 1780	26 1956	CA1944	cil1-4	cil1-4	HE858225	HE858225	CII
	Osmaniye-Boru nver	37.1780	30.4830	CA1945	cil1-4	cil1-5	HE858225	ACC	CIL
				OBMBCA46353	cil1-4	cil1-4	HE858225	HE858225	CIL
				OBMBCA46354	ant3-5	cil1-4	HE858238	HE858225	CII /ANT
				OBMBCA46355	ant3-5	cil1-4	HE858238	HE858225	CIL/ANI
				OBMBCA46356	rid3-8	cil2-1	ACC	ACC	CIL/RID
	Pazarojk	37 2810	37 1240	OBMBCA46357	cil1-4	cil1-4	HE858225	HE858225	
	Fazaicik	37.2819	37.1249	OBMBCA46358	cil1-4	cil1-4	HE858225	HE858225	CIL
				OBMBCA46359	cil1-4	cil1-4	HE858225	HE858225	
				OBMBCA46360	rid3-8	cil1-4	ACC	HE858225	CIL/RID
				OBMBCA46361	cil1-4	cil1-1	HE858225	ACC	
				OBMBCA46362	cil1-4	cil1-4	HE858225	HE858225	CIL
Turkey	Pize	41.0110	40 5792	CA771	ant11-7	ant11-7	ACC	ACC	ANT
Turkey	Rize	41.0110	40.3792	CA772	ant11-7	ant11-7	ACC	ACC	ANI
	Rize-Çiftekavak	41.0283	40.4858	CA779	ant11-7	ant11-6	ACC	ACC	ANT
	Dizo Homain Stroom	41 1554	40.0012	CA769	ant11-7	ant11-7	ACC	ACC	ANT
	Kize-Hellişili Suealli	41.1554	40.9012	CA770	ant3-5	ant11-7	HE858238	ACC	
				CA891	ant3-2	ant11-7	ACC	ACC	
	Samsun-Bafra	41.5167	36.0158	CA892	ant3-2	ant7-3	ACC	ACC	ANT
				CA893	ant3-2	ant7-3	ACC	ACC	
				CA897	ant3-2	ant3-2	ACC	ACC	
	Samsun-Kavak	41.1567	36.0927	CA899	ant11-7	ant11-1	ACC	ACC	ANT
	Sullsul Kuvuk			CA900	ant11-7	ant11-6	ACC	ACC	
				CA886	ant11-6	ant7-3	ACC	ACC	ANT
	Samsun-19 Mayıs	41.4674	36.0993	CA887	ant3-5	ant7-3	HE858238	ACC	
	Sunsul 17 Mayls			CA888	ant3-2	ant11-7	ACC	ACC	

	Samsun-19 Mayıs	41.4674	36.0993	CA889	ant3-2	ant11-6	ACC	ACC	ANT
				CA878	ant3-2	ant11-7	ACC	ACC	
	Someun Takkakäy	41 2126	26 4700	CA879	ant3-2	ant11-7	ACC	ACC	ANT
	Sanisui-Tekkeköy	41.2120	30.4790	CA881	ant11-7	ant11-7	ACC	ACC	ANI
				CA882	ant3-2	ant7-3	ACC	ACC	
				SGCA42202	ant1-4	ant1-5	ACC	ACC	
	Seydisehir	37 4557	31 8156	CA1458	ant11-7	ant10-4	ACC	ACC	ANT
	Seydişemi	37.4337	51.8150	CA1459	ant10-4	ant8-3	ACC	ACC	AN
				CA1460	ant1-4	ant8-3	ACC	ACC	
		37.9726		CA1372	rid3-8	ant11-7	ACC	ACC	RID/ANT
				CA1373	rid3-8	ant11-7	ACC	ACC	
Turkey	Siirt-Başyurt river		41 7806	CA1374	rid3-8	rid3-7	ACC	ACC	RID
			41.7000	CA1375	ant11-9	ant11-9	ACC	ACC	ANT
Turkey				CA1376	ant11-7	ant11-7	ACC	ACC	
				CA1377	ant11-7	ant11-9	ACC	ACC	
				CA1378	ant11-7	ant11-7	ACC	ACC	ANT
				CA1379	ant11-7	ant11-9	ACC	ACC	
	Siirt-Kezer river	37.9610	41.8573	CA1380	rid3-8	ant11-7	ACC	ACC	RID/ANT
				CA1383	ant11-9	ant3-5	ACC	HE858238	ANT
				CA1384	ant11-9	ant11-9	ACC	ACC	
				OBMB33257	cil2-1	ant10-1	ACC	ACC	
				OBMB33258	ant5-6	cil1-4	HE858230	HE858225	
	Silifke	36 3130	33 9595	OBMB33264	ant5-6	cil1-4	HE858230	HE858225	CIL/ANT
	Shirke	50.5150	55.7575	OBMB33265	ant5-6	cil1-4	HE858230	HE858225	_
			_	OBMB33270	ant5-6	cil1-4	HE858230	HE858225	
				OBMB33271	ant5-6	ant5-6	HE858230	HE858230	ANT

	Silifke	36.3130	33.9595	OBMB33272	ant5-6	cil1-4	HE858230	HE858225	CIL/ANT
				CA925	ant11-7	ant7-3	ACC	ACC	
	Sinop-Ayancık	41.9483	34.7749	CA926	ant11-7	ant11-7	ACC	ACC	ANT
				CA927	ant11-7	ant11-7	ACC	ACC	
				CA930	ant11-7	ant11-6	ACC	ACC	
	Sinon Corzo	41 9567	25,0000	CA932	ant11-7	ant7-3	ACC	ACC	ANT
	Shiop-Geize	41.8507	33.0999	CA933	ant11-7	ant7-3	ACC	ACC	ANI
				CA934	ant11-7	ant7-3	ACC	ACC	
				CA920	ant11-7	ant7-3	ACC	ACC	
	Sinon Erfelek Cohanlar	41 0543	35.0196	CA921	ant11-7	ant7-3	ACC	ACC	ANT
	Shiop-Effect Çobalilar	41.9545		CA922	ant11-7	ant7-3	ACC	ACC	ANI
Turkey				CA923	ant11-7	ant7-3	ACC	ACC	
	Sinop-Kabalı village	41.8468	35.0503	CA936	ant11-7	ant11-6	ACC	ACC	ANT
Turkey		41.0400	33.0303	CA938	ant11-7	ant11-7	ACC	ACC	AN
				CA1090	ant11-7	ant3-5	ACC	HE858238	
				CA1091	ant11-7	ant3-5	ACC	HE858238	
	Sivas-Hafik lake	39 8727	37 3825	CA1092	ant11-7	ant11-7	ACC	ACC	ANT
	Sivas-Halik lake	39.0727	57.5625	CA1093	ant11-7	ant3-5	ACC	HE858238	AU
				CA1094	ant11-7	ant11-7	ACC	ACC	
				CA1095	ant11-7	ant11-7	ACC	ACC	
				CA1100	ant11-7	ant11-7	ACC	ACC	
	Sivas-Kızılırmak	39.7154	37.0117	CA1101	ant11-7	ant11-7	ACC	ACC	ANT
				CA1102	ant11-7	ant11-7	ACC	ACC	
				CA1085	ant11-7	ant11-7	ACC	ACC	ANT
	Sivas-Tecer river	39.6922	37.0034	CA1087	ant11-7	ant11-7	ACC	ACC	
				CA1088	ant3-5	ant3-5	HE858238	HE858238	

				CA1097	ant11-7	ant11-7	ACC	ACC	
	Sivas-Tödürge lake	39.8713	37.6068	CA1098	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1099	ant11-7	ant11-7	ACC	ACC	
				CBCA6444	ant11-7	ant11-7	ACC	ACC	
	Sorgun Dam	38.6555	29.3386	CBCA6445	ant7-3	ant10-1	ACC	ACC	ANT
				CBCA6446	ant7-3	ant7-3	ACC	ACC	
				CA1300	rid3-8	rid3-7	ACC	ACC	RID
	Sanluurfa Bozova	37 3588	38 5281	CA1302	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Şainturia-Bozova	57.5588	36.5264	CA1303	rid3-8	rid3-8	ACC	ACC	RID
				CA1304	cil1-4	cil1-4	HE858225	HE858225	CIL
		37.1488		CA1294	rid3-8	ant11-7	ACC	ACC	RID/ANT
	Şanlıurfa-Harran road		38 0681	CA1295	rid3-8	rid3-8	ACC	ACC	RID
Turkey			38.9081	CA1296	rid3-8	ant3-5	ACC	HE858238	RID/ANT
				CA1298	rid3-8	rid3-7	ACC	ACC	RID
				CA1306	rid3-8	ant3-5	ACC	HE858238	RID/ANT
				CA1307	rid3-8	ant3-5	ACC	HE858238	KID/AIVI
	Şanlıurfa-Mardin road 30. km	37.1467	38.1144	CA1308	ant3-5	ant3-5	ACC	HE858238	ANT
				CA1309	rid3-8	ant3-5	ACC	HE858238	RID/ANT
				CA1310	ant11-7	ant3-5	ACC	HE858238	ANT
				YECA15144	ant2-1	ant10-1	ACC	ACC	
				YECA15145	ant4-3	ant10-1	ACC	ACC	
				YECA15146	ant2-1	ant4-3	ACC	ACC	
	Tefenni	37.2333	29.7113	KYECA15147	ant2-1	ant4-3	ACC	ACC	ANT
				KYECA15149	ant3-5	ant10-5	HE858238	ACC	_
				MAHACA15151	ant3-5	ant3-5	HE858238	HE858238	
				MAHACA15152	ant3-5	ant2-1	HE858238	ACC	

				CA1508	ant11-7	rid2-1	ACC	ACC	PID/ANT
				CA1509	ant11-7	rid1-2	ACC	ACC	KID/AN1
	Tekirdağ-Bıyıkali lake	41.0114	27.3930	CA1510	ant7-2	ant11-7	ACC	ACC	
				CA1511	ant11-7	ant11-7	ACC	ACC	ANT
				CA1512	ant11-7	ant10-4	ACC	ACC	
				CA1513	ant11-7	ant11-7	ACC	ACC	ANT
				CA1514	ant11-7	rid1-2	ACC	ACC	
	Tekirdağ Kenenekli dam	41 1121	27 5510	CA1515	ant7-3	rid1-3	ACC	ACC	RID/ANT
	Teknuag-Kepenekii uani	41.1121	27.3319	CA1516	ant11-7	rid1-3	ACC	ACC	
				CA1517	ant11-7	ant11-7	ACC	ACC	ANT
				CA1518	ant7-3	ant11-7	ACC	ACC	
Turkey				CA1503	ant7-3	ant7-3	ACC	ACC	ANT
	Tekirdağ-Yazır lake		27.398	CA1504	ant7-3	rid3-4	ACC	ACC	
Turkey		40.9330		CA1505	ant11-7	rid1-2	ACC	ACC	RID/ANT
				CA1506	ant11-7	rid1-7	ACC	ACC	
				CA1507	ant11-7	ant11-7	ACC	ACC	ANT
				CA1070	ant11-7	ant3-2	ACC	ACC	ANT
	Tokat-Avlunlar stream	40.5114	36.7342	CA1071	rid2-1	ant3-2	ACC	ACC	
				CA1072	rid2-1	ant3-2	ACC	ACC	KID/ANT
				CA1079	ant11-7	ant11-7	ACC	ACC	
	Tokat-Cördük plataeu	40.2207	36 5622	CA1080	ant11-7	ant3-5	ACC	HE858238	ANT
	Tokai-Çorduk platacu	40.2207	30.3022	CA1081	ant11-7	ant3-2	ACC	ACC	ANT
				CA1082	ant11-7	ant11-7	ACC	ACC	
	Tokat-Kömec Vesilirmak	40 3396	36 4605	CA1055	ant11-7	ant11-7	ACC	ACC	ANT
	Tokat-Kömeç Yeşilırmak	+0.3370	36.4605	CA1059	ant11-6	ant3-2	ACC	ACC	
	Tokat-Yalancı stream	40.4106	36.7153	CA1062	ant11-7	ant7-3	ACC	ACC	ANT

				CA1063	ant11-7	rid2-1	ACC	ACC	RID/ANT
	Tokat-Yalancı stream	40.4106	36.7153	CA1064	ant11-7	ant7-3	ACC	ACC	ANT
				CA1065	ant11-7	ant7-3	ACC	ACC	ANI
				KO60318	ant11-6	rid2-1	ACC	ACC	RID/ANT
	Tokat Yusufoğlan	40.3421	36.9325	KO60319	ant11-7	ant11-6	ACC	ACC	ANT
				KO60320	ant3-5	ant7-3	HE858238	ACC	AN
				CA792	ant11-6	ant6-2	ACC	ACC	
	Trabzon-Arsin	40.9515	39.9009	CA793	ant11-6	ant6-2	ACC	ACC	ANT
				CA795	ant6-2	ant6-2	ACC	ACC	
			_	CA782	ant11-7	ant6-2	ACC	ACC	
	Trabzon-Darica	41.0718	39.5113	CA783	ant11-7	ant11-7	ACC	ACC	ANT
Turkey				CA787	ant11-7	ant7-3	ACC	ACC	
	Trabzon Derecik	41.0558	39.3900	RYDCA61256	ant7-3	ant6-2	ACC	ACC	ANT
Turkey	Trabzon-Söğütlü	41 0064	39.6310	CA790	ant11-7	ant7-3	ACC	ACC	ANT
		41.0004	59.0510	CA791	ant11-7	ant11-2	ACC	ACC	
				OBMBCA46373	cil1-4	bed1-5	HE858225	ACC	CIL/BED1
				OBMBCA46374	cil1-4	cil2-1	HE858225	ACC	СП
	Türkoğlu	37.3808	36.8764	OBMBCA46376	cil1-4	cil2-1	HE858225	ACC	CIL
				OBMBCA46377	cil1-4	ant3-5	HE858225	ACC	CIL/ANT
				OBMBCA46378	cil1-4	cil1-4	HE858225	HE858225	CIL
				CA1562	ant7-2	ant7-3	ACC	ACC	
	Valova-Altinova stream	40 7084	29 4730	CA1563	ant7-2	ant7-3	ACC	ACC	ANT
	Talova-Attilova sticali	40.7004	29.4750	CA1565	ant9-1	ant7-2	ACC	ACC	ANT
				CA1567	ant7-2	ant7-3	ACC	ACC	
	Yalova-Taşköprü	40 6801	29.3907 —	CA1568	ant7-2	ant7-3	ACC	ACC	ANT
		40.6801		CA1570	ant7-3	ant7-3	ACC	ACC	

	Valova Tasköprü	40.6801	29 3907	CA1572	ant7-3	ant11-7	ACC	ACC	ANT
	таюча-таҙкорти	40.0801	29.3907	CA1573	ant7-2	ant7-3	ACC	ACC	ANI
				ISCA1579	ant3-5	ant4-3	HE858238	ACC	
				ISCA1580	ant3-5	ant4-3	HE858238	ACC	
				ISCA1581	ant10-1	ant10-1	ACC	ACC	
				ISCA1582	ant2-1	ant10-1	ACC	ACC	ANT
	Vesilova	37 53/18	29 6473	ISCA1583	ant10-4	ant10-4	ACC	ACC	
	i cșnova	57.5540	27.0475	ISCA1584	ant4-4	ant10-1	ACC	ACC	
				ISCA1585	ant4-3	ant11-7	ACC	ACC	
				ISCA1587	ant2-1	ant8-5	ACC	ACC	
				ISCA1588	ant10-1	ant10-1	ACC	ACC	
Turkey				ISCA1589	ant4-3	ant2-1	ACC	ACC	
	Yozgat-Gedikhasanlı	39.5720		CA1111	rid2-1	ant3-2	ACC	ACC	RID/ANT
Turkey			35.1373	CA1112	ant11-7	ant3-5	ACC	HE858238	ANT
				CA1113	ant11-6	ant7-3	ACC	ACC	
				CA1114	ant11-7	ant11-6	ACC	ACC	
				CA1117	rid2-1	ant3-5	ACC	HE858238	RID/ANT
	Vozgat Mükramin laka	30 8030	35 2215	CA1119	ant11-7	ant11-7	ACC	ACC	ANT
	i ozgat-wuktemin iake	39.8030	55.2215	CA1120	rid2-1	ant11-7	ACC	ACC	
				CA1121	rid2-1	ant11-7	ACC	ACC	KID/ANT
				CA1105	ant11-7	ant11-7	ACC	ACC	
	Yozgat-Sorgun	39.8042	35.2035	CA1106	ant11-7	ant11-6	ACC	ACC	ANT
				CA1107	ant11-7	ant11-6	ACC	ACC	
				CA1124	ant11-7	ant11-7	ACC	ACC	ANT 8
	Yozgat-Yerköy Delice river	39.6117	34.5031	CA1125	ant11-7	ant11-7	ACC	ACC	
	102gat-Terkoy Dence river			CA1126	ant3-2	ant3-5	ACC	HE858238	

	Yozgat-Yerköy Delice river	39.6117	34.5031	CA1127	ant10-4	ant3-5	ACC	HE858238	ANT
				TBCASE0154	cil1-4	cil2-2	HE858225	ACC	
				TBCASE0155	cil1-4	cil1-4	HE858225	HE858225	CII
				TBCASE0157	cil1-4	cil1-2	HE858225	ACC	CIL
				TBCASE0158	cil1-5	cil1-7	ACC	ACC	
	Yumurtalık	36.6925	35.6288	TBCASE0160	cil1-4	rid3-8	HE858225	ACC	CIL/RID
				TBCASE0161	cil1-4	cil1-4	HE858226	HE858226	CIL
				TBCASE0163	cil1-4	cil1-8	HE858225	ACC	
				TBCASE0164	cil1-4	cil2-1	HE858225	ACC	
				TBCASE0165	cil1-4	cil1-5	HE858225	ACC	
		38.3443		CA638	ant11-7	ant3-5	ACC	HE858238	
Turkey	Van-Edremit Çiçekli		43.1881	CA639	ant11-8	ant11-8	ACC	ACC	ANT
				CA641	ant3-5	ant3-5	HE858238	HE858238	
Turkey				CA642	ant11-8	ant3-5	ACC	HE858238	
			42 1020	CA645	ant11-7	ant11-7	ACC	ACC	ANT
	Van Gevas	38 3070		CA647	ant11-8	ant3-5	ACC	HE858238	
	vaii-Oevaş	38.3079	45.1252	CA649	ant11-7	ant11-4	ACC	ACC	
				CA651	ant11-8	ant11-8	ACC	ACC	
				CA652	rid3-8	ant3-5	ACC	HE858238	RID/ANT
	Van-Muradiye Bendimahı River	38.9366	43 6611	CA653	rid3-8	rid3-8	ACC	ACC	RID
	van-Muraurye Bendiniani Kiver	58.7500	45.0011	CA654	rid3-8	ant3-5	ACC	HE858238	RID/ANT
				CA656	ant11-9	ant3-5	ACC	HE858238	ANT
				CA1732	ant10-1	ant10-4	ACC	ACC	ANT
	Usak-Gediz river	38 7878	29 2243	CA1733	ant4-3	ant11-7	ACC	ACC	
		56.7678	27.2273	CA1734	ant11-7	ant10-1	ACC	ACC	
				CA1735	ant11-7	ant11-7	ACC	ACC	

	Uşak-Gediz river	38.7878	29.2243	CA1736	ant11-7	ant10-1	ACC	ACC	ANT
				CA1742	ant1-4	ant3-2	ACC	ACC	
				CA1743	ant7-3	nt11-7ant10-1ACCACCunt1-4ant3-2ACCACCunt7-3ant10-1ACCACCnt11-7ant10-4ACCACCint3-2ant11-7ACCACCint3-2ant11-7ACCACCint7-3ant11-7ACCACCint1-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt12-1rid1-7ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant10-1ACCACCnt11-7ant1-6ACCACCnt11-7ant1-6ACCACCant7-3ant7-3ACCACCnt11-6ant1-7ACCACCnt11-7ant12-1ACCACCnt11-6ant12-1ACCACCnt11-6ant12-1ACCACCant6-2ant6-2ACCACCant6-2ant6-2ACCACC			
	Usak Göğan laka	28 7207	20 5562	CA1744	ant11-7	ant10-4	ACC	ACC	ANT
	Uşak-Oogen lake	38.7207	29.3303	CA1745	ant3-2	ant11-7	ACC	ACCANTACCACCACCACCACCANTACCANTACCACCACCANT	
				CA1746	ant7-3	ant11-7	ACC	ACC	
				CA1747	ant11-7	ant10-1	ACC	ACC	
				CA1737	ant10-4	ant10-4	ACC	ACC ACC ACC ACC	
	Usak Karakoo laka	38 6542	20 3336	CA1738	ant11-7	ant10-1	ACC	ACC	ANT
	Oşak-Kalakoç lake	38,0342	29,3330	CA1740	ant11-7	ant10-1	ACC	ACC	ANI
				CA1741	ant10-1	ant10-1	ACC	ACC	
Turkey	Zonguldak	41 4271	31 7268	FGCA67251	ant3-2	ant3-2	ACC	ACC	ANT
Turkey	Zonguluak	71.7271	51.7200	FGCA67252	ant12-1	rid1-7	ACC	ACC	RID/ANT
	Zonguldak-Alaplı			CA1013	ant11-7	ant10-1	ACC	ACC	
		41.1515	31.3521	CA1014	ant11-7	ant7-3	ACC	ACC	ANT
				CA1016	ant11-6	ant11-6	ACC	ACC	
				CA1003	ant11-7	ant11-6	ACC	ACC	
	Zonguldak-Kilimli	41.4005	31.6837	CA1004	ant7-3	ant7-2	ACC	ACC	ANT
				CA1006	ant7-3	ant7-3	ACC	ACC	
				CA996	ant11-6	ant7-3	ACC	ACC	ANT
				CA997	ant11-7	ant12-1	ACC	ACC	AN
	Zonguldak-Şirinköy	41.5065	31.9726	CA998	ant7-3	rid1-7	ACC	ACC	RID/ANT
				CA1000	ant11-6	ant12-1	ACC	ACC	ANT
				CA1001	ant11-3	ant11-6	ACC	ACC	
Ukraine	Crimea-Theodosia town	45 0458	35.3716 -	42-1	ant6-2	ant6-2	ACC	ACC	ANT
Ukraine	Crimea-Theodosia town	45.0458		42-2	ant6-1	rid2-5	ACC	ACC	RID/ANT

				42-3	ant7-3	rid2-5	ACC	ACC	
				42-4	ant6-2	ant6-1	ACC	ACC	ANT
	Crimea-Theodosia town	45.0458	35.3716	42-5	ant6-2	ant6-1	ACC	ACC	ANI
				42-7	ant6-1	ant6-1	ACC	ACC	
				42-8	ant6-1	rid2-5	ACC	ACC	RID/ANT
	Kharkiv-Chepel	49.2666	36.9166	57-1	ant8-2	rid1-5	ACC	ACC	RID/ANT
				44-01	rid1-5	rid1-5	ACC	ACC	RID
				44-02	ant8-2	rid1-5	ACC	ACC	
				44-3	ant7-3	rid2-5	ACC	ACC	RID/ANT
				44-8	ant7-3	rid1-5	ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC ACC		
	Kharkiv-Gaidary village	49.6205	36.3286	44-9	ant7-3	ant7-3	ACC	ACC	ANT
				44-10	ant7-3	rid1-5	ACC	ACC	
Ukraine				44-12	ant8-2	rid1-5	ACC	ACC	RID/ANT
Okraine				44-13	ant7-3	rid1-5	ACC	ACC	ANT RID/ANT RID/ANT RID/ANT ANT ANT ANT RID/ANT RID/ANT RID/ANT RID/ANT RID/ANT ANT RID/ANT ANT RID/ANT ANT RID/ANT ANT RID/ANT ANT
				44-25	ant7-3	ant7-3	ACC	ACC	ANT
				46-01	ant7-3	ant7-3	ACC	ACC	
				46-2	ant7-3	ant7-3	ACC	ACC	ANT
	Kharkiy Krasnokutek	50.0527	35 1494	46-4	ant7-1	ant7-1	ACC	ACC	
	Kilaikiv-Kiasilokutsk	50.0527	55.1494	46-6	ant11-7	ant7-3	ACC	ACC	
				46-7	rid2-5	ant7-3	ACC	ACC	RID/ANT
				46-8	ant7-3	ant7-3	ACC	ACC	ANT
				54-1	ant7-3	rid1-5	ACC	ACC	RID/ANT
	Kharkiv-Kolomak	49.8166	35.2833	54-2	ant8-2	ant8-2	ACC	ACC	ANT
				54-3	ant7-3	ant7-3	ACC	ACC	AINT
	Kharkiy-Lintsi village	50 2291	36.3955	45-01	ant7-3	ant7-3	ACC	ACC	ANT
	Kharkiv-Liptsi village	50.2291		45-02	ant8-2	rid1-5	ACC	ACC	RID/ANT

				45-3	ant8-2	rid1-5	ACC	ACC	RID/ANT
				45-7	ant7-3	ant7-3	ACC	ACC	ANT
LUracino	Wheelvin Lintsi village	50 2201	26 2055	45-8	ant7-3	rid1-5	ACC	ACC	
Okrailie	Kharkiv-Liptsi vinage	50.2291	30.3933	45-9	ant8-2	rid1-5	ACC	ACC	
				45-12	ant8-2	rid1-5	ACC	ACC	KID/AN1
				45-14	ant8-2	rid1-5	ACC	ACC	
	Andijan-near Khanabad village	40.7999	72.9666	4-01	sp nov-3	sp nov-3	ACC	ACC	SP NOV
	Karakalpakistan-Takhtakupyr town	43.0333	60.25	14-12	ter-4	ter-4	ACC	ACC	TER
TT-h-1-i-t	Kashkadarya-near Dehkanabad village	38.3300	66.5000	10-10	ter-1	ter-1	ACC	ACC	TER
Uzbekistan	Consordour d	20 (922	(7.0666	7-8	sp nov-2	sp nov-2	ACC	ACC	SP NOV
	Samarkand	39.0833	07.0000	7-12	ter-2	ter-2	ACC	ACC	TER
	Tashkent-Novyi Chinaz village	40.9166	68.7166	6-29	sp nov-4	sp nov-4	ACC	ACC	SP NOV
Outgroups									
Create	Demati/Crete	35.0333	25.2833	Hotz19856/ZMB56960	cre-2	cre-3	FN432374	FN432375	
Greece	Skinias/Crete	35.0632	25.3083	Hotz19852/ZMB56959	cre-1	cre-1	FN432376	FN432376	CRE
Create	Igoumenitsa	39.5000	20.2666	Hotz19421	epe-1	epe-1	FN432369	FN432369	EDE
Greece	Lechena/Peloponnese	37.9174	21.2667	Hotz19552	epe-2	epe-2	FN432370	FN432370	EPE
Itoly	Metaponto	40.3737	16.8028	No.6311	ber-1	ber-1	FN432381	FN432381	DED
italy	Tarsia	39.6166	16.2666	Hotz19586	ber-2	ber-2	FN432382	FN432382	DEK
Italy	Carbonare	45.9333	11.2166	-	les-1	les-1	FN432384	FN432384	
Cormony	Parlin Staahlin			3466	les-1	les-1	FN432384	FN432384	LES
Germany	Bernn-Stechnin			3467	les-1	les-1	FN432384	FN432384	

				16863	shq-1	shq-1	ACC	ACC		
Montenegro	Lake Skutari	42.2410	19.1061	16861	shq-2	shq-2	ACC	ACC	SHQ	
				16914	shq-1	shq-1	ACC	ACC		
France	Ligagnaeu	43.5333	4.7500	Lib11	per-1	per-1	FN432377	FN432377	PER	
Algoria	Abadla	31.0197	-2.7360	Hotz16704	sah-1	sah-1	FN432380	FN432380	SAD	
Algeria	El Golea	30.5732	2.8920	Hotz16737	sah-2	sah-2	FN432379	FN432379	SAR	

Table 3.4. Calculation of genetic distance among mitohondrial subgroups, main haplogroups, and outgroup species. Arithmetic means and standart deviations of Tamura-Nei distances (Tamura and Nei, 1993) within (blue) and between (black) main haplogroups (MHG1-8), outgroups (gray coloured) and subgroups of MHG6 and MHG2, estimated with a gamma distributed shaped parameter (G = 0.325) and rate heterogeneity among groups in the program MEGA 5.10 (Tamura et al., 2011). Subgroups of MHG6 and MHG2 did not compare with each other because only the amount of genetic distance within MHGs was tried to estimate.

	MHG6a	MHG6b	MHG6c	MHG6d	MHG2a	MHG2b
MHG6a	0.008 ±0.001					
MHG6b	0.025 ±0.004	0.003 ±0.001				
MHG6c	0.019 ±0.003	0.017 ±0.003	0.007 ±0.001			
MHG6d	0.024 ±0.004	0.023 ±0.004	0.017 ±0.003	0.006 ±0.001		
MHG2a	-	-	-	-	0.009 ±0.002	
MHG2b	-	-	-	-	0.022 ±0.003	0.008 ±0.002

1. MHG1	0.008 ±0.001															
2. MHG2	0.079 ±0.010	0.014 ±0.002														
3. MHG3	0.099 ±0.012	0.093 ±0.011	0.005 ±0.001													
4. MHG4	0.085 ±0.011	0.080 ±0.010	0.081 ±0.011	0.003 ±0.001												
5. MHG5	0.092 ±0.011	0.088 ±0.011	0.087 ±0.010	0.041 ±0.006	0.007 ±0.001											
6. MHG6	0.080 ±0.010	0.079 ±0.009	0.076 ±0.010	0,046 ±0.007	0.046 ±0.006	0.014 ±0.002										
7. MHG7	0.094 ±0.011	0.086 ±0.010	0.086 ±0.011	0,054 ±0.008	0.053 ±0.007	0.032 ±0.005	0.009 ±0.001									
8. MHG8	0.102 ±0.012	0.095 ±0.011	0.096 ±0.012	0.063 ±0.008	0.057 ±0.008	0.052 ±0.007	0.058 ±0.008	0.001 ±0.001								
9. P.cretensis	0.183 ±0.021	0.159 ±0.019	0.178 ±0.019	0.167 ±0.020	0.179 ±0.021	0.176 ±0.021	0.181 ±0.021	0.197 ±0.022								
10. P.epeiroticus	0.253 ±0.030	0.269 ±0.032	0.279 ±0.036	0.294 ±0.036	0.259 ±0.031	0.260 ±0.032	0.284 ±0.034	0.267 ±0.034	0.315 ±0.036							
11. P.lessonae	0.283 ±0.035	0.269 ±0.032	0.255 ±0.029	0.265 ±0.030	0.227 ±0.024	0.255 ±0.027	0.274 ±0.030	0.264 ±0.030	0.321 ±0.037	0.343 ±0.041						
12. P.bergeri	0.302 ±0.037	0.279 ±0.033	0.253 ±0.030	0.257 ±0.029	0.236 ±0.026	0.244 ±0.027	0.257 ±0.029	0.245 ±0.028	0.330 ±0.041	0.349 ±0.043	$0.050 \\ \pm 0.008$					
13. P.shqipericus	0.300 ±0.038	0.267 ±0.032	0.288 ±0.034	0.286 ±0.035	0.262 ±0.033	0.278 ±0.032	0.292 ±0.035	0.273 ±0.033	0.313 ±0.037	0.336 ±0.043	0.120 ±0.016	0.109 ±0.015				
14. P.perezi	0.367 ±0.044	0.333 ±0.039	0.357 ±0.043	0.349 ±0.041	0.348 ±0.040	0.354 ±0.042	0.370 ±0.043	0.357 ±0.044	0.398 ±0.045	0.453 ±0.055	0.397 ±0.050	0.387 ±0.047	0.374 ±0.046			
15. P.saharicus	0.457 ±0.054	0.406 ±0.046	0.475 ±0.054	0.435 ±0.049	0.402 ±0.046	0.433 ±0.050	0.439 ±0.049	0.424 ±0.048	0.449 ±0.048	0.457 ±0.053	0.462 ±0.054	0.440 ±0.052	0.447 ±0.049	0.342 ±0.039		
16. P.nigromaculatus	0.520 ±0.062	0.490 ±0.057	0.526 ±0.061	0.501 ±0.061	0.516 ±0.064	0.506 ±0.058	0.510 ±0.061	0.530 ±0.061	0.523 ±0.058	0.593 ±0.068	0.525 ±0.063	0.513 ±0.061	0.490 ±0.059	0.436 ±0.050	0.463 ±0.046	

Table 3.5. Calculation of genetic distance among nuclear subgroups, main allele groups and outgroup species. Arithmetic means and standart deviations of the Tamura-3-parameter (T92) model (Tamura, 1992) within (blue) and between (black) main allele groups (Anatolia, Asia and Ridibundus), *P.bedriagae 2, P.cypriensis*, outgroups (gray coloured) and subgroups of Anatolia MAG and allele groups of Asia and Ridibundus MAGs estimated with a gamma distributed shaped parameter (G = 0.622) in the program MEGA 5.10 (Tamura et al., 2011).

ANATOLIA MAG									
	MAGa	MAGb	MAGc	MAGd	MAGe				
MAGa	0.003 ±0.001								
MAGb	0.013 ±0.003	0.003 ±0.001							
MAGc	0.013 ±0.003	0.006 ±0.002							
MAGd	0.014 ±0.003	0.007 ±0.002	0.005 ±0.002	0.004 ±0.001					
MAGe	0.018 ±0.004	0.010 ±0.002	0.009 ±0.002	0.010 ±0.002	0.003 ±0.001				

	ASIA MAG										
	CILI	CIL3	TER	SP NOV	BED1						
CILI	0.002 ±0.001										
CIL3	0.004 ±0.001	0.001 ±0.001									
TER	0.005 ±0.002	0.005 ±0.002	0.001 ±0.001								
SP NOV	0.010 ±0.003	0.007 ±0.002	0.011 ±0.003	0.001 ±0.001							
BED1	0.007 ±0.002	0.005 ±0.002	0.008 ±0.003	0.011 ±0.003	0.003 ±0.001						

RIDIBUNDUS MAG											
	RID1 RID2 RID3 KUR										
RID1	0.004 RID1 ±0.001										
RID2	0.006 ±0.002	0.002 ±0.001									
RID3	0.010 ±0.003	0.005 ±0.002	0.001 ±0.001								
KUR	0.011 ±0.003	0.002 ±0.001									

	1	2	3	4	5	6	7	8	9	10	11	12	13
1. Anatolia MAG	0.010 ±0.002												
2. Asia MAG	0.040 ±0.006	0.006 ±0.001											
3. Ridibundus MAG	0.039 ±0.006	0.013 ±0.003	0.007 ±0.001										
4. Levant MAG (bed2)	0.050 ±0.007	0.043 ±0.007	0.046 ±0.007	0.001 ±0.001									
5. Cyprus MAG	0.039 ±0.006	0.034 ±0.006	0.038 ±0.006	0.041 ±0.007	0.002 ±0.001								
6. P.cretensis	0.035 ±0.006	0.033 ±0.006	0.036 ±0.006	0.035 ±0.006	0.021 ±0.005								
7. P.epeiroticus	0.025 ±0.005	0.025 ±0.005	0.025 ±0.005	0.042 ±0.007	0.041 ±0.007	0.037 ±0.006							
8. P.lessonae	0.038 ±0.010	0.025 ±0.008	0.033 ±0.009	0.029 ±0.009	0.031 ±0.009	0.030 ±0.008	0.029 ±0.009						
9. P.bergeri	0.041 ±0.010	0.028 ±0.008	0.036 ±0.010	0.032 ±0.009	0.034 ±0.009	0.033 ±0.009	0.032 ±0.010	0.006 ±0.004					
10. P.shqipericus	0.054 ±0.008	0.044 ±0.007	0.042 ±0.007	0.041 ±0.007	0.054 ±0.008	0.045 ±0.007	0.044 ±0.007	0.044 ±0.011	0.050 ±0.012				
11. P.perezi	0.070 ±0.009	0.056 ± 0.008	0.058 ±0.008	0.070 ±0.010	0.064 ±0.009	0.062 ±0.009	0.061 ±0.009	0.042 ±0.010	0.045 ±0.010	0.077 ±0.011			
12. P.saharicus	0.072 ±0.009	0.061 ±0.009	0.063 ±0.009	0.076 ±0.011	0.067 ±0.010	0.068 ±0.010	0.064 ±0.009	0.053 ±0.011	0.056 ±0.012	0.080 ±0.011	0.031 ±0.006		
13. P.nigromaculatus	0.130 ±0.020	0.113 ±0.018	0.120 ±0.018	0.113 ±0.019	0.116 ±0.020	0.111 ±0.019	0.125 ±0.020	0.127 ±0.023	0.126 ±0.023	0.146 ±0.025	0.108 ±0.018	0.125 ±0.020	

(Groups	Unique molecular variations		Allele	Groups	
		-	ANT1	ANT2	ANT3	-
		C: 122	A or C: 434	A:	434	
		G: 137	T: 524	C:	524	
		C: 155	A: 1053	C: 1053	A: 1053	
		C: 162	T: 1069	C:1069	T:1069	
	MAGa	C: 184				
		G: 450				
		T: 732				
		4G: 514-517				
		5CA: 738-747				
		6T: 1135-1140				
		-	ANT4	ANT5	-	-
		G: 541	T: 1057	C: 621		
	MAGh	G: 641				
	11100	5G: 514-518				
		5CA: 738-747				
		6T: 1135-1140				
		C: 605				
IAG		A: 641				
A M	MAGc	4G: 514-517				
0L1		5CA: 738-747				
VAT		6T: 1135-1140				
AI		-	ANT6	ANT7	ANT8	-
			A:9	T: 780		
			T: 23	A: 851		
			C: 68	C: 905		
	MAGd	A: 300	C: 255	8 bp deletion:		
		5CA: 738-747	C: 410	938-947		
			A: 514	A: 970		
			A: 580			
			4G: 514-517	4G: 514-517	3-4G: 514-517	
			6T: 1135-1140	5T: 1135-1139	5-6T: 1135-1140	
		-	ANT9	ANT10	ANT11	ANT12
		G: 319	A: 471	C: 384		T: 759
		T: 527				
		C: 532				
	MAGe	A: 533				
		C: 584				
		3G : 514-516				
		5CA : 738-747				
		6T: 1135-1140				

 Table 3.6. Group specific unique molecular variations for nuclear SAI-1+RanaCR1 gene.

Groups	Unique molecular variations			Allele Groups		
	-	CILI	CIL3	TER	SP NOV	BED1
		A or G: 762	C or T: 697	A: 469	C: 174	A or C: 384
ŋ				A: 483	C: 379	T or C: 411
MA	3G: 514-516				C: 390	A: 413
ASIA	5T: 1135-1139				G: 438	G: 419
					T: 478	
					T: 506	
		6CA: 738-749	6CA: 738-749	6CA: 738-749	6CA: 738-749	4-6CA: 738-749
	-	RID1	RID2	RID3	KUR	-
75		G or T: 316	G or A: 538	G: 63	T: 186	
4AC		A: 413	A or G: 569	T or C: 446	A: 202	
JS N	3G: 514-516	G: 419	A or G: 733	A or G: 489	A: 415	
IQN		A or C: 899	T or C: 740	T: 759	124 bp deletion	
BUI				A: 1014	: 758 - 890	
RIDII					C: 1057	
		5-6CA: 738-749	6-7CA: 738-751	8-10CA: 738-757	6CA: 738-749	
		5T: 1135-1139	5-6T: 1135-1140	6T: 1135-1140	6T: 1135-1140	

CHAPTER 4

CONCLUSIONS

In the first manuscript;

 \checkmark In the 5' region of the water frog serum albumin gene, several potential regulatory elements were identified. These were TATA box, hepatocyte-specific promoter element, CAT box, amphibian downstream element, ADEI and ADEII, which could be involved in tissue specific expression.

✓ In the 3' region of the water frog serum albumin gene, several conserved cis-regulatory elements were found: poly(A)-limiting elements a and b (PLEa, PLEb), U-rich upstream element 1 (USE1), poly(A) signal (PAS), U-rich upstream element 2 (USE2), polyadenylation site (PAS), and GU-rich downstream element (GU-rich DSE).

 \checkmark In contrast to strong length conservation in protein coding exons among distinct vertebrate species, intron lengths in the albumin gene of *P. lessonae* showed a considerable variation, and were quite longer than those in other vertebrate species especially 3,4, 5, 8, 10, 11 and 13.

✓ Retroelements (belonging distinct classes of LTR, non-LTR elements, DNA transposons, and simple repeats) found upstream of the 5' UTR, downstream of the 3' UTR and within individual albumin introns could be one of the important drivers of the albumin gene evolution.

 \checkmark Particularly Tc1 like elements (TLE) has 11 truncated copies in the serum albumin gene representing seven distinct intact TLEs, which could have an important impact on not only serum albumin gene but also genome evolution.

 \checkmark Derived consensus elements of these seven TLE obtained from several truncated copies in the *P. lessonae* genome indicated that some of them could be recently transferred horizontally to the *P. lessonae* genome since some of them had several long copies and a few active copies in the genome such as PL Tc1-1Ory and PL Tc1-DR1 elements.

 \checkmark In contrast, others could be potentially very old elements since they have several short truncated copies, very long insertion and deletions in both left and right ITRs or the transcriptional regulatory region, and none of copies has an intact ORF such as PL Tc1-10Xt, PL Tc1-RT, PL Tc1-PP and PL Tc1-FR3.

✓ All these had ITRs varying in size from 29 bp to 221 bp which contained DRs in their extremities. The size of the transcriptional regulatory region changed from 72 bp (PL Tc1-FR3) to 187 bp (PL Tc1-RT). Except PL Tc1-DR1 and PL Tc1-FR3, they all contained a TA duplication at the end of ITRs.

 \checkmark In addition to TLEs, six truncated hAT-like elements (HLE), representing three distinct HLE were identified in the serum albumin gene. These were named as PL hAT-2-TS, hAT-2-ET, hAT-2-OG elements that all could be ancient elements because they had indels in both ITRs, the transcriptional regulatory region or ORF.

 \checkmark Phylogenetic analysis of nucleotide and aa sequences of albumin gene exons gave similar topological results except the position of *P. epeiroticus* which formed a sister group to *P. ridibundus*.

In the manuscript 2,
\checkmark mtDNA and nuDNA markers revealed discordances in the tree topology, the level of genetic differentiation, the number of genetic stocks, the times of divergence and the patterns of geographic distribution.

✓ mtDNA tree represented several well diverged monophyletic groups while nuDNA tree contained both weakly supported and some well supported groups.

✓ In the mtDNA tree, *P. bedriagae* (MHG2) indicated an ingroup species, clustering with *P. ridibundus* (MHG1) as a sister group, whereas in the nuDNA tree it (Levant MAG5) formed a clade within the outgroup species *P. shqipericus/P. lessonae/ P. bergeri*.

 \checkmark *P. cretensis* showed a divergence earlier than *P. cypriensis* in the mtDNA tree, but these two formed sister species in the nuDNA tree. The estimated time of divergence for mtDNA indicated divergence of P. cretensis before 9 Ma while the estimated time of divergence for nuDNA indicated the same divergence time both for P. cypriensis and P. cretensis about 6-5 Ma.

 \checkmark *P. epeiroticus* split off before *P. cretensis/P. cypriensis* in the mtDNA tree, but in the nuDNA tree was more closely related with the Anatolia and Asia/Rid MAGs.

 \checkmark *P. ridibundus* in the mtDNA tree formed a sister group with *P. bedriagae* from the Levant whereas in the nuDNA tree it clustered with the Asia MAG2, including groups from Central Asia, central-southern Turkey and the Levant (BED1) which indicated very recent divergence (2.0-1.7 Ma).

 \checkmark In contrast to the well diverged Cilician (MHG4 and 5) and Central Asia (MHG7 and 8) monophyletic groups in the mtDNA tree about 3.5-4.5 Ma, their relationships within the Asian MAG2 were not fully resolved. nuDNA findings indicated a very shallow divergence among these groups about 1.5-0.5 Ma.

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EDUCATION

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FOREIGN LANGUAGES

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PUBLICATIONS

Theses:

M. Sc. Thesis: Detection of species boundaries in the *Rana ridibunda* complex of Southwestern Turkey using mitochondrial ND3 marker.

Research Articles:

1. Plötner, J., Baier, F., Akın, Ç., Mazepa, G., Schreiber, R., Beerli, P., Litvinchuk, S.N., Bilgin, C.C., Borkin, L., Uzzell, T. (2012) Genetic data reveal

that water frogs of Cyprus (genus Pelophylax) are an endemic species of Messinian origin. *Zoosystematics and . Evolution.* 88: 261-283.

2. Akın, Ç., Bilgin, C.C., Beerli, P., Westaway, R., Ohst, T., Litvinchuk, S.N., Uzzell, T., Bilgin, M., Hotz, H., Guex, G.-D., Plötner, J. (2010b). Phylogeographic patterns of genetic diversity in eastern Mediterranean water frogs were determined by geological processes and climate change in the Late Cenozoic. *Journal of Biogeography* 37: 2111-2124.

3. Akın, Ç., Bilgin, M. & Bilgin C.C. (2010) Discordance between ventral colour and mtDNA haplotype in the water frog *Rana* (*ridibunda*) *caralitana*, 1988 Arıkan. *Amphibia-Reptilia* 31: 9-20.

International Congress Presentations:

Oral presentations:

1. Akın, Ç., Bilgin, C.C., Hotz, H., Beerli, P., Westaway, R., Ohst, T., Guex, G.-D., Litvinchuk, S.N., Uzzell, T., Bilgin, M., Plötner, J. (2009). Use of genetic divergence in water frogs to constrain geodynamics and landscape development in the eastern Mediterranean Region. Abstract: International Workshop on Active Tectonic Studies and Earthquake Hazard Assessment in Syria and Neighboring Countries, Arab School of Science and Technology – Damascus – Syria, 17-19 November, pp, 92-93.

2. Çilingir, F.G., Akın Pekşen, Ç., Ambarlı, H., Beerli, P., Bilgin, C.C. (2014) Brown bears from Turkey show exceptionally high maternal lineage diversity. Ecology and Evolutionary Biology Symposium (12-13 July, İstanbul), pp: 13-14.

3. Akın Pekşen, Ç, Bilgin, C.C., Beerli, P., Westaway, R., Schreiber, R., Mazepa, G., Uzzell, T., Plötner, J. (2014) Do discordant mitochondrial and nuclear distribution patterns indicate introgression following secondary contact between Anatolian water frog lineages (Pelophylax cf. bedriagae) and European water frog lineage (Pelophylax ridibundus) in Anatolia? Ecology and Evolutionary Biology Symposium (12-13 July, İstanbul), pp: 19.

Poster presentation:

1. Akın, Ç., Bilgin, M., Kaya, B., Bilgin, C.C. (2007) Discordance between ventral color and mtDNA haplotype in the water frog *Rana (ridibunda) caralitana*. Abstract (poster presentation): Origin and Evolution of Biota in Mediterranean Climate Zones – Zurich – Switzerland, 14-15 July, pp, 14.

National Congress Presentations:

Oral presentations:

1. Akın, Ç., Bilgin, M., Bilgin, C.C. (2006) Rana (*Ridibunda*) caralitana Arıkan 1988 (Amphibia: Ranidae) taksonunun yayılış sınırlarının moleküler yöntemlerle saptanması. 18. Ulusal Biyoloji Kongresi (26-30 Haziran, Aydın). Bildiri Kitabı Sayfa: 75.

2. Akın, Ç., Bilgin, C.C., Hotz, H., Beerli, P., Westaway, R., Ohst, T., Guex, G.-D., Litvinchuk, S.N., Uzzell, T., Bilgin, M., Plötner, J. (2010). Anadolu su kurbağalarının (Pelophylax (Rana)) mitokondriyal DNA genetik çeşitliliğinin geçmiş jeolojik olaylarla şekillenmesi. 20. Ulusal Biyoloji Kongresi (21-25 Haziran, Denizli). Bildiri Kitabı Sayfa: 165-166.

3. Akın, Ç. Kılıç, A., Kürüm, V., Plötner, J., Bilgin, C.C. (2011) Anadolu su kurbağaları (Ranidae: Pelophylax) ticaretinin ekolojik etkileri. 10. Ulusal Ekoloji ve Çevre Kongresi (3-7 Ekim, Çanakkale). Bildiri Kitabı Sayfa: 24.

4. Kunduz, E., Akın, Ç., Bilgin, C.C. (2011) Moleküler yöntemlerle kuşlarda eşey tayini. 10. Ulusal Ekoloji ve Çevre Kongresi (3-7 Ekim, Çanakkale). Bildiri Kitabı Sayfa: 25.

5. Çilingir, F.G., Akın, Ç., Ambarlı, H., Bilgin, C.C. (2012) Zor yakalanan hayvan gruplarından boz ayıların (Ursus arctos) dışkı, kıl, ve postlarından girişimsiz genetik örnekleme başarısı. 21. Ulusal Biyoloji Kongresi (3-7 Eylül, İzmir).

Poster presentation:

1. Kunduz, E., Akın, Ç., Bilgin, C.C. (2012) Kuşlarda moleküler yöntemlerle eşey belirlenmesinde kullanılan üç primer sisteminin sonuçlarının kıyaslanması. 21. Ulusal Biyoloji Kongresi (3-7 Eylül, İzmir).

Book Chapter:

1. Plötner, J., Uzzell, T., Beerli, P., Akın, Ç., Bilgin, C.C., Haefeli, C., Ohst, T., Köhler, F., Schreiber, R., Guex, G.-D., Litvinchuk, S.N., Westaway, R., Reyer, H.-U., Pruvost, N., Hotz, H (2010). Genetic divergence and evolution of reproductive isolation in eastern Mediterranean water frogs. In: Glaubrecht M (Ed). Evolution in Action. Case studies in Adaptive Radiation, Speciation and the Origin of Biodiversity. Special volume originating from contributions to the Priority Programme SPP 1127 "Radiations: Origins of Biological Diversity" of the DFG. Springer, Heidelberg, Berlin, pp. 373-403.

PROJECTS:

1. Investigation of population biology and the effect commercial trade on eastern Mediterranean water frog populations (*Pelophylax* spp). TUBITAK TBAG 112T913.

2. Detection of genetic diversity in the eastern, southeastern Anatolia, Black Sea and east of Central Anatolian water frog populations. TUBİTAK TBAG 111T584.

3. Genetic divergence and evolution of reproductive isolation in Eastern Mediterranean water frogs. Complementary action among German, Switzerland and Turkey, financed by DFG (Germany) (2002-2009).

4. Detection of species boundaries in the *Rana ridibunda* complex of Southwestern Turkey using mitochondrial ND3 marker. (BAP-08-11-DPT.2002K120510-DK-14). OYP-DPT Project.

RESEARCH ACTIVITY:

1. Studies on molecular markers at Museum für Naturkunde, Berlin, Germany (supervised by Dr. Jörg Plötner)

i. Application of ITS2 markers on the southwestern Anatolian water frog populations for three months in 2006

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AWARDS/GRANTS:

1. First Honor in B. Sc., Gazi University University, June 2002

2. Training course grant for Conservation Genetics: Assessing populations structure and dynamics through the use of molecular markers and novel computational models", Château Liblice, Czech Republic, 1-7 September 2008.

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