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Molecular Systematics of *Cercocarpus* H.B.K. (Rosaceae)

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Molecular Systematics of *Cercocarpus* H.B.K. (Rosaceae)

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For Traci, the love of my life, without which there is nothing

For Mom and Dad

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Molecular Systematics of *Cercocarpus* H.B.K. (Rosaceae)

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Brian David Vanden Heuvel, Ph.D.

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Supervisor: C. Randal Linder

Cercocarpus H.B.K. (Rosaceae) is a New-World genus comprised of montane xerophytic shrubs and trees found in deserts, chaparral, and montane regions of western North America. *Cercocarpus* has been shown to be infected by *Frankia*, a nitrogen-fixing soil actinomycete. *Frankia* strains provide their hosts with a source of fixed nitrogen, a nutrient that often controls plant growth. This dissertation set out to create molecular phylogenies for *Cercocarpus* for later studies of coevolutionary relationships between *Cercocarpus* and *Frankia*. Only by knowing the evolutionary histories of both lineages can an in-depth study of coevolution proceed. *Cercocarpus* lacks a consensus classification or phylogeny. Past confusion about the number and circumscription of species within *Cercocarpus* arose from a lack of clear morphological discontinuities between taxa. Variations in leaf morphology show a continuum within and between taxa, making delineation of species difficult. Due to the variation in morphology,

extensive sampling was conducted of all described taxa of *Cercocarpus*. The external transcribed spacer (ETS) of the nuclear ribosomal DNA was developed in *Cercocarpus* for the purpose of phylogenetic reconstruction. We document unexpected, deep coalescence of paralogous ETS types in *Cercocarpus*. This is the first record of the maintenance of ETS paralogues that were not produced by recent hybridization and lack of concerted evolution or pseudogene formation. We produced three phylogenies for *Cercocarpus* taxa, one based on ETS sequence data, another on AFLP presence/absence data, and a third based on chloroplast intergenic spacer sequences. Areas of congruence between the three trees supports clear taxonomic conclusions. Most notably, *C. montanus* is distinct from *C. betuloides*, the two newly described species of *Cercocarpus* have affinities to *C. fothersgilloides*, and the endangered *C. traskiae* is more closely related to *C. betuloides* than *C. fothersgilloides*, as previously thought. Taxon sampling had a great impact on tree topology, highlighting the need to do thorough sampling for closely related plant groups. We also discovered all three independent phylogenies were significantly incongruent, suggesting a complex, reticulating evolutionary history.

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Chapter 1: Introduction

GENERAL INTRODUCTION

Cercocarpus H.B.K. (Rosaceae) is a New World genus composed of montane xerophytic shrubs and trees found in deserts, chaparral, and montane regions of western North America. Considerable attention has been paid to the ecology and management of *Cercocarpus* because the evergreen leaves found on most members of the genus have high levels of protein and are important winter wildlife and livestock forage in the western United States and Mexico (Blauer et al. 1975, Davis and Brotherson 1991). The high level of protein found in leaves of *Cercocarpus* is due to an association with *Frankia*, a nitrogen-fixing soil actinomycete. *Frankia* strains provide their hosts with a source of fixed nitrogen, a nutrient that, when absent, limits plant growth. The host plant, in turn, provides fixed carbon to the *Frankia* strain (Baker and Schwintzer 1990). Much like legumes associated with *Rhizobia*, plants that associate with *Frankia* have a distinct advantage over other plants because their growth is not limited by the absence of nitrogen and can inhabit nitrogen poor soils. Therefore, *Cercocarpus* can play an important role in ecosystem development by securing unstable surfaces and promoting growth of other plants by adding nitrogen into the soil (Klemmedson 1979).

My current and future research interests are to study the coevolutionary history of the mutualistic association between *Cercocarpus* and *Frankia*. *Cercocarpus*, the largest genus of actinorhizal plants in the Rosaceae, is a group of reasonable size (approx. 10 species in 4 species complexes) and geographic distribution (western North America), making it ideally suited for the study of an actinorhizal symbiosis. One of the requirements of a coevolutionary study is to create phylogenetic hypotheses for both members in the mutualistic association. This dissertation addresses the construction of phylogenetic hypotheses for *Cercocarpus*, and will be crucial for a study of the coevolutionary relationships between *Cercocarpus* and *Frankia*.

The systematics of *Cercocarpus* are largely unknown; the genus lacks a consensus classification or phylogeny. Since it was first described, *Cercocarpus* has undergone three revisions (Schneider 1905b, Rydberg 1913, Martin 1950), each varying widely in the ranks assigned to taxa and the number of species. Confusion about the number and circumscription of species within *Cercocarpus* arises primarily from the lack of clear morphological delimitations between taxa (Schneider 1905b, Rydberg 1913, Martin 1950). Variations in leaf morphology show a continuum within and between taxa, making it difficult to define boundaries among species (Brayton and Mooney 1966, Mortenson 1973). In recent years, molecular data have proved useful for resolving relationships among

morphologically similar species and for understanding the diversity within species (Soltis and Soltis 1998 and references therein).

This dissertation is composed of three independent chapters, each addressing a different aspect of the evolution and systematics of the genus *Cercocarpus*, using molecular characters. To facilitate discussion of evolutionary relationships in *Cercocarpus*, the second part of this introductory chapter will give a background overview of the taxonomic and systematic work on *Cercocarpus* that has been completed to date.

The remainder of the dissertation (Chapters 2-4) is organized as independent, interrelated papers. Each chapter will be submitted for publication as an independent paper, complete with introduction, materials and methods, results, and discussion.

The second chapter investigates the molecular evolution of the external transcribed spacer (ETS) of the nuclear ribosomal DNA (nrDNA) within *Cercocarpus*. The ETS region was initially developed for use as a phylogenetic marker in *Cercocarpus*. In order to create a phylogenetic tree for a recently diverged lineage, a DNA region must be identified that contains sufficient phylogenetically informative variation (Soltis and Soltis 1998). Preliminary studies found little informative variation in the internal transcribed spacer (ITS) region of the nuclear ribosomal DNA (nrDNA) repeat, a common nuclear marker used for phylogenetic reconstruction at low taxonomic levels in plants (Baldwin

et al. 1995). We, therefore, developed the external transcribed spacer (ETS) of the nrDNA repeat for phylogenetic reconstruction in *Cercocarpus* to search for the variation we needed, as well as take advantage of the benefits (concerted evolution and easy amplification) that the ribosomal repeat displays (Nickrent and Soltis 1998).

Chapter two documents four main ETS paralogues that exhibit a deep coalescence in the genus. This was an unexpected result because concerted evolution should be homogenizing the rDNA copies. The observation of multiple rDNA copies is not unprecedented (Dvorak 1990, Suh et al. 1993, Buckler and Holtsford 1996, O’Kane et al. 1996, Waters and Schaal 1996, Wendel et al. 1995, Sang et al. 1995, and Campbell et al. 1997). Instances of multiple rDNA copies have been observed in many plant groups, and explained using hypotheses of hybridization and pseudogene formation (Buckler et al. 1997). Multiple ETS types have also been found in previous studies, yet the multiple ETS types coalesced within species, having little impact on the phylogenetic relationships reconstructed between species (Linder et al. 2000). This was not the case in *Cercocarpus*. We were unable to explain convincingly the multiple types found in *Cercocarpus* using recent hybridization or pseudogene formation. We discuss other possible reasons for the multiple paralogous ETS types with deep coalescence in *Cercocarpus*, the implications that multiple ETS types have for

concerted evolution in the rDNA repeat, and hypothesize about the sequence of events that led to the current ETS diversity.

In the third chapter, we report two nuclear phylogenetic hypotheses for *Cercocarpus*. To address correctly previous hypotheses about species and varietal designations, we conducted dense taxon sampling across all recognized taxa (see second half of this introductory chapter). The first phylogenetic hypothesis used amplified fragment length polymorphisms (AFLP) as characters for reconstruction. The second hypothesis was obtained using one of the four ETS types identified in Chapter 2. We found the two phylogenies were significantly incongruent, even though we sampled the same individuals. We address the implications and possible reasons for the incongruency. We also test taxonomic and evolutionary hypotheses made by previous authors for *Cercocarpus* using both the ETS and AFLP trees. Finally, we found that the taxonomic sampling we had done was critical to a correct understanding of the evolution of *Cercocarpus*. Reduced datasets for both AFLP and ETS phylogenies had well supported incongruent tree topologies. This result is important beyond *Cercocarpus*, highlighting the need for dense taxon sampling in other phylogenetic studies in recently evolved plant lineages.

The fourth chapter presents a phylogeny based on chloroplast DNA sequence data. We did not find enough phylogenetically informative variation for the complete resolution of all *Cercocarpus* taxa. The chloroplast phylogeny was

significantly incongruent to both the nuclear ETS and amplified fragment length polymorphisms (AFLP) phylogenies, and areas of congruence and incongruence are discussed.

Overall, this dissertation is a model for current limits and difficulties when attempting to reconstruct phylogenies for closely related groups of plants. Figures and Tables will be included in their respective chapters; however, references have been compiled into a single bibliography at the end of the dissertation, along with relevant appendices.

SYSTEMATIC BACKGROUND OF *CERCOCARPUS H.B.K.*

Cercocarpus was first collected near Mexico City by Humboldt and Bonpland and was described and illustrated by Kunth (1823). It was identified as a part of the Rosaceae based on the presence of a tubular calyx and a single one-celled ovary that produced an achene. Because of the presence of one, solitary ovule in the ovary and a lack of petals, it was also recognized as an undescribed genus. The first collection was named *Cercocarpus fothergilloides* H.B.K.

During the next 70 years, many more collections were made and new species and varieties described. These included *Cercocarpus montanus* described by Rafinesque in 1832 and individuals currently named *C. betuloides*, collected in California by Nuttall in the 1830's.

There have been three formal revisions of *Cercocarpus*. A comparison of all three taxonomic treatments is illustrated in Figure 1.1. C. K. Schneider

(1905b) conducted the first formal revision of *Cercocarpus* using borrowed material from the Missouri Botanical Garden. Schneider recognized 10 species and 12 varieties (Figure 1.1). P. A. Rydberg re-evaluated *Cercocarpus* in 1913 in his treatment for the *Flora of North America*. Rydberg recognized 21 species and no varieties. Due to the criticism of Rydberg's revision of *Cercocarpus* by later authors, who considered many of the species described by Rydberg not worthy recognition, F. L. Martin, a Masters Student at Pomona College, followed with the third and last revision in 1950. Martin was able to view much more material that had been collected in the 37 years following Rydberg's revision. Martin recognized only six species and 11 varieties (Figure 1.1). Such a drastic reduction in recognized species left many botanists perplexed, preferring instead an intermediate number of species between the extremes of Rydberg's (1913) and Martin's (1950) revisions (Little 1953). Although Martin viewed more material in his revision, his classification has not been widely accepted as evidenced by its absence from current western states floras (Kearny and Peebles 1951, Munz 1959, Hitchcock and Cronquist 1961, Lis 1993).

Much of the disagreement between taxonomic treatments arises from a lack of characters and intergrading morphological variation between taxa. Martin (1950) addressed this difficulty when he noted on page 91:

All the members of this genus are related very closely; intergradations of all distinguishing characteristics is the rule rather than the exception, and hybridization apparently occurs even between those entities sufficiently distinct to merit specific rank. A feature which seems characteristic of one entity often appears sporadically in other entities (91).

Martin's admission (1950) of a lack of characters and intergradations of forms was then used to justify his drastic reduction in the number of species in *Cercocarpus*. The key, then, for further studies in this group is to find new characters to use to describe taxa.

In order to help clarify the species delineations, Richard Lis (1992) approached *Cercocarpus* using leaf architecture characters. Based on this work, he proposed a re-organized classification that included four species complexes; a *ledifolius* species complex, a *betuloides* species complex, a *montanus* species complex, and a *fothergilloides* (Mexican) species complex. Lis (1993) followed this study by resurrecting the *C. betuloides* name and re-classifying the California taxa for the Jepson manual to reflect his view of the diversity

A list of the recognized *Cercocarpus* taxa used in this study can be found in Table 1.1 and Figure 1.1. The recognized taxa are organized following the species complexes hypothesized by Lis (1992). The following is a description of the diversity, past and current taxonomic entities, and geographic distribution of the four species complexes.

Table 1.1: A list of the recognized *Cercocarpus* taxa for this study. Included is the publication citation.

taxon	Place of publication
<i>montanus</i> complex	
<i>Cercocarpus montanus</i> Raf.	<i>Atl. J.</i> pg. 146. 1832.
<i>C. montanus</i> Raf. var. <i>montanus</i> F. L. Martin	<i>Brittonia</i> 7(2). pg. 101. 1950.
<i>C. montanus</i> Raf. var. <i>glaber</i> (S. Wats.)F.L. Martin	<i>Brittonia</i> 7(2). pg. 101. 1950.
<i>C. montanus</i> Raf. var. <i>paucidentatus</i> (S. Wats.)F.L. Martin	<i>Brittonia</i> 7(2). pg. 104. 1950.
<i>C. montanus</i> Raf. var. <i>argenteus</i> (Ryd.)F.L. Martin	<i>Brittonia</i> 7(2). pg. 104. 1950.
<i>betuloides</i> complex	
<i>Cercocarpus betuloides</i> Nutt.	Torr. and Gray, <i>Flora of North America.</i> pg. 427. 1840.
<i>C. betuloides</i> Nutt. var. <i>betuloides</i>	<i>Jepson Manual,</i> pg 950. 1993.
<i>C. betuloides</i> Nutt. var. <i>blancheae</i> (C.K.Schneid.)R.F. Thorne	<i>Aliso</i> 9(2). pg. 193. 1978.
<i>Cercocarpus minutiflorus</i> Abrams	<i>Bull. Torr. Bot. Club</i> 37. pg. 149. 1910.
<i>fothergilloides</i> complex	
<i>Cercocarpus fothergilloides</i> H.B.K	<i>Nov. Gen. And Sp.</i> 6. pg 232. 1832.
<i>C. fothergilloides</i> H.B.K. var. <i>fothergilloides</i> Henrickson	ined.
<i>C. fothergilloides</i> H.B.K. var. <i>medranoanus</i> Henrickson	ined.
<i>C. fothergilloides</i> H.B.K. var. <i>mojadensis</i> (C.K. Schneid.)Henrickson	ined.
<i>Cercocarpus pringlei</i> (C.K. Schneid.)Ryd.	Rydberg. <i>North American Flora</i> 22. pg 420. 1913
<i>Cercocarpus macrophyllus</i> C. K. Schneid.	<i>Ill. Handb. Laubholz.</i> 1. pg 530. 1905a
<i>Cercocarpus traskiae</i> Eastw.	<i>Proc. Cal. Acad.</i> 3. pg. 136. 1898.
<i>ledifolius</i> complex	
<i>Cercocarpus ledifolius</i> Nutt.	Torr. and Gray, <i>Flora of North America.</i> pg. 427. 1840.
<i>C. ledifolius</i> Nutt. var. <i>intermontanus</i> N. Holmgren	<i>Brittonia</i> 39(4). pg 424. 1987.

<i>C. ledifolius</i> Nutt. var. <i>ledifolius</i>	<i>Brittonia</i> 39(4). pg 426. 1987.
<i>Cercocarpus intricatus</i> S. Wats.	<i>Proc. Amer. Acad. Arts</i> 10. pg 346. 1875
New taxa	
<i>Cercocarpus mexicanus</i> Henrickson	<i>Sys. Bot.</i> 12. pg. 293. 1987.
<i>Cercocarpus rzedowskii</i> Henrickson	<i>Sys. Bot.</i> 12. pg. 296. 1987.

Taxonomic background of the *Cercocarpus ledifolius* complex

Members of the *Cercocarpus ledifolius* complex are found distributed in the Great Basin (Figure 1.2), and can be easily identified by their leaves with entire margins (Figure 1.3). The type specimen for *Cercocarpus ledifolius* was collected in Caribou Co., Idaho, near the Bear River of the Timpanagos near the celebrated “Beer Springs” by Nuttall in July, 1834 (see Table 1.1). In 1875, Sereno Watson described *Cercocarpus intricatus* from a collection he made near the mouth of the American Fork Canyon, near Saint George, Utah. In his description he mentioned that *Cercocarpus intricatus* was probably closely allied to *Cercocarpus ledifolius*, but differed by its low, intricately branched growth and by its smaller, tightly revolute leaves. Sixteen years later, Marcus E. Jones (1891b) subsumed *Cercocarpus intricatus* to a variety of *C. ledifolius* based on his observation that at the type locality, he found a complete transition in morphology between *C. ledifolius* and *C. intricatus* as he decreased in altitude. Also in 1891a, Jones described *C. arizonicus* from a collection in Coconino Co.,

Arizona, near Willow Springs. Jones recognized that *C. arizonicus* was very similar in morphology to *C. ledifolius* var. *intricatus*, but differed in flower size, length of persistent style on the achene, and the vestiture of the leaves.

In the first formal revision of the genus in 1905b, Schneider recognized the *Cercocarpus ledifolius* group, which included *Cercocarpus ledifolius* Nutt. and *C. intricatus* S.Wats.. Schneider (1905b) further described *Cercocarpus ledifolius* Nutt. var. *intercedens* C.K. Schneid. as a taxon with smaller, more revolute leaves that were intermediate between *C. ledifolius* and *C. intricatus*, and *Cercocarpus intricatus* S.Wats. var. *villosus* C.K.Schneid. that was similar to *C. intricatus*, yet had a much denser vestiture. *Cercocarpus arizonicus* was not recognized, but was used as a synonym of *C. intricatus* var. *villosus*.

Rydberg (1913) recognized *Cercocarpus ledifolius*, *C. intricatus*, and *C. arizonicus*. He further described *Cercocarpus hypoleucus* Ryd. from a collection near Melrose Montana, to replace *C. ledifolius* var. *intercedens*.

Martin (1950), in the last formal revision, recognized the *Cercocarpus ledifolius* complex as containing three varieties based mostly on leaf size. Individuals with the largest leaves were given the name *Cercocarpus ledifolius* Nutt. var. *typicus* C.K. Schneid. Individuals with intermediate sized leaves were described as *C. ledifolius* var. *intercedens* C.K. Schneid., and *C. ledifolius* var. *intricatus* referred to the small leafed individuals. Holmgren (1987), while preparing a treatment for the *Intermountain Flora*, recognized that the actual type

for *Cercocarpus ledifolius* Nutt. was actually *C. ledifolius* var. *intercedens* from Schneider (1905b). This meant that the large-leafed taxon was not named at any rank. Holmgren therefore re-classified the *Cercocarpus ledifolius* complex to include two varieties: *C. ledifolius* var. *intermontanus* and *C. ledifolius* var. *ledifolius*, with variety *intermontanus* being the more common, large-leafed taxon. He did not include *Cercocarpus intricatus* as a variety of *ledifolius*, recognizing it as a distinct species. For this study, we will use the taxonomy from Holmgren's treatment, but have tested if there is indeed molecular support for his classification (Chapter 3 and 4). Holmgren's re-classification is illustrated in Figure 1.1.

Much of the leaf diversity (Figure 1.3) and intergradations in leaf morphology between the two *Cercocarpus ledifolius* varieties (variety *intermontanus* and variety *ledifolius*) and *C. intricatus* (Figure 1.4) has been attributed to hybridization. In the original description of the *Cercocarpus ledifolius* complex, Schneider (1905b) considered his variety *intercedens* to be a hybrid taxon between variety *intricatus* and variety *typicus*. Holmgren (1987) agreed with this view, recognizing that in zones of contact, intermediate forms could be observed. Brayton and Mooney (1966) conducted a detailed study of fifteen populations of *Cercocarpus ledifolus* that displayed leaf variation. They found that individuals of variety *intermontanus* were at higher altitudes, and individuals classified as *C. intricatus* were found at much lower, drier sites.

Although individuals with intermediate leaf morphologies (*C. ledifolius* var. *ledifolius*) occupied the intermediate elevations between the two “parental populations,” leaf variation indicative of all three taxa was also found in the intermediate zone. Brayton and Mooney (1966) attributed the elevated degree of leaf variation to hybridization coupled with different selective conditions year-to-year, resulting in the establishment of different leaf types. We have also tested this assumption using molecular data (Chapter 3 and 4).

Taxonomic background of the *Cercocarpus montanus* complex

Rafinesque described *Cercocarpus montanus* in the 1832 from a collection by E. P. James who accompanied Long on his first expedition to Colorado during the summer of 1820. Nuttall later collected a member of the same taxon on the Platte River in Colorado and named it *Cercocarpus parvifolius* in 1839. Other early collections and species descriptions came from Asa Gray (1853), who named *Cercocarpus breviflorus* from a collection by Wright near Frontera, New Mexico, which may be Fronteras, Sonora, Mexico as commented by Rydberg (1913). *Cercocarpus breviflorus* was recognized as distinct from *C. montanus* and *C. parvifolius* by its spiny, short habit and smaller leaves. Watson (1882) described a similar taxon to *Cercocarpus breviflorus*, who named it *Cercocarpus parvifolius* Nutt. var. *paucidentatus* from a collection in San Luis Potosi in Mexico. These collections had much smaller leaves than the original *C. parvifolius* collections. Jones (1891b) recognized the commonalities between

Cercocarpus parvifolius and *C. brevifolius*, and therefore subsumed *C. breviflorus* as a variety of *parvifolius* under the name *C. parvifolius* Nutt. var. *breviflorus* Colville ex. M.E. Jones.

Schneider (1905b), in the first formal revision, recognized both *C. parvifolius* Nutt. and *C. breviflorus* Gray. He further described two new varieties of *C. parviflorus* Nutt.: variety *hirsutissimus* C.K.Schneid. from a collection by Hayden in the Laramie Mountains, Wyoming, and variety *minimus* C.K.Schneid. from an Englemann collection from Utah. Rydberg (1913) later found that *C. parvifolius* var. *minimus* was actually a specimen of *Holodiscus microphyllus*. Schneider also described *C. breviflorus* Gray var. *eximus* C.K. Schneid. from a collection by Earle from the El Captain mountains, Lincoln Co., New Mexico. Schneider further described *Cercocarpus treleasei* from a collection in San Luis Potosi, Mexico. He did not recognize *C. parvifolius* var. *paucidentatus* S.Wats.

Rydberg (1913) recognized that *C. montanus* Raf. and *C. parvifolius* Nutt. were the same taxon, and used *C. montanus* in his Flora of North America. He also recognized that *C. treleasei* and *C. parvifolius* var. *paucidentatus* S.Wats. were the same taxon, elevating Watson's variety *paucidentatus* to specific rank. *C. breviflorus* Gray was recognized as being distinct, based on glabrate adaxial leaf surfaces, less tomentum on the abaxial leaf surface, and a more northerly distribution. Rydberg elevated *C. breviflorus* var. *eximus* to specific rank, and

further described two new species: *Cercocarpus flabellifolius* Ryd. from a collection in southern Utah, and *Cercocarpus argenteus* from Randall Co., Texas.

Martin (1950) subsumed all of the afore-mentioned species and varieties under three varieties of *C. montanus*: *C. montanus* var. *montanus*, *C. montanus* var. *paucidentatus*, and *C. montanus* var. *argenteus* (Figure 1.1). Lis's (1992) view of the *Cercocarpus montanus* complex also included the three varieties as described by Martin (1950), yet he maintained that *Cercocarpus montanus* var. *paucidentatus* displayed sufficient differences in leaf morphology to be recognized at the species level. For this dissertation, we will refer to this taxon at the varietal rank. It differs from variety *montanus* and variety *argenteus* by having much smaller leaves with fewer marginal teeth. It is broadly distributed from Arizona across New Mexico into Texas and south into Mexico. Within this complex, Lis (1992) found *Cercocarpus montanus* var. *montanus* and *C. montanus* var. *argenteus* were very closely related. Both are winter deciduous shrubs, while all other members of *Cercocarpus* are evergreen. *C. montanus* var. *montanus* is broadly distributed from western South Dakota to northern New Mexico and from western Oklahoma to western Utah (Figure 1.6). *C. montanus* var. *argenteus* is found primarily in Palo Duro Canyon in the Texas panhandle, eastern New Mexico, and in the Guadalupe Mountains of west Texas (Figure 1.6).

A fourth taxon recognized as part of the *Cercocarpus montanus* complex for this study is *C. montanus* var. *glaber*, described from a collection from California. Martin (1950) recognized this taxon as a widespread Californian taxon. McVaugh (1952) first misused *Cercocarpus montanus* var. *glaber* (S. Wats)F.L. Martin and later by Correll and Johnston (1970) in the *Manual of the Vascular Plants of Texas*, to describe broad-leafed individuals with approximately the same geographic distribution as variety *paucidentatus*. Since Lis (1992) reclassified the Californian diversity under the name *C. betuloides*, there is no opportunity for confusion. For the sake of convenience, we will continue to use *C. montanus* var. *glaber* to refer to the taxon as McVaugh (1952) and Correll and Johnston (1970) intended and not use it to refer to a Californian taxon.

Because of the lack of discontinuities in leaf morphology between varieties in the *Cercocarpus montanus* complex, which is the primary source of characters for classification, the naturalness of the varieties are in question (Figure 1.7) (Chapter 2 and 3).

Taxonomic background of the *Cercocarpus betuloides* complex

Nuttall collected specimens of *Cercocarpus* from the mountains of Santa Barbara, California during the 1830's. Nuttall sent specimens both to Torrey and Gray, who named it *C. betuloides* Nutt. (*Flora of North America* 1840), and to Hooker, who named it *C. betulaefolius* Nutt. ex Hook., (*Icones plantarum* pl. 322 1840). Watson (1876) did not recognize *Cercocarpus betuloides* Nutt. or *C.*

betulaefolius Nutt. as displaying sufficient morphological differences to warrant specific status, and described *C. parvifolius* Nutt. var. *glaber* S.Wats.

Schneider (1905b) subsumed both these California taxa under one name, *C. betulaefolius*, with three varieties: variety *typicus* C.K. Schneid., variety *blancheae* C.K. Schneid., and variety *minor* C.K. Schneid. (Figure 1.1). Rydberg (1913) noted that *C. betuloides* had been published before *C. betulaefolius*, and recognized only *C. betuloides*. Rydberg did not recognize the varieties of Schneider, but instead described four new species: *C. alnifolius* Ryd. from a collection by Trask near Avalon, Santa Catalina Island; *C. douglasii* Ryd. from a collection by Douglas in California; *C. rotundifolius* Ryd. from a collection by Grant near Los Angeles; and *C. macrourus* Ryd. from a collection by Baker in Modoc Co., California. Additionally, Rydberg recognized *C. minutiflorus*, a species described by Abrams (1910) from San Diego (Figure 1.1).

In the last formal revision of *Cercocarpus*, Martin (1950) grouped all the described, broad-leaved Californian diversity into three varieties of *C. montanus* (Figure 1.1). This taxonomic move was rejected by many of the floras of the Pacific states, including the *Arizona Flora* (Kearny and Peebles 1951) and the *Flora of Southern California* (Munz 1979), all of which retained the name *C. betuloides*. After his study of leaf architecture characters, Lis (1992) recognized a close relationship between Martin's varieties *glaber*, *minutiflorus*, and *macrourus*. Because these three varieties are all located in California (Figure 1.8), Lis (1993)

used the name *C. betuloides* and continued the tradition in Pacific states of recognizing *C. betuloides* as a separate entity than *C. montanus*.

For this dissertation, we will recognize the taxonomy of Lis (1993) for the California diversity of *Cercocarpus*. Lis delimited two species, *C. betuloides* with three varieties and *C. minutiflorus* (Figure 1.1, Figure 1.9).

Taxonomic background of the *Cercocarpus fothergilloides* complex

Schneider (1905a,b) described three new taxa similar to *Cercocarpus fothergilloides*: *C. mojadensis* C.K. Schneid., *C. mojadensis* var. *pringlei* C.K. Schneid., and *C. macrophyllus* C.K. Schneid. Rydberg (1913) raised the variety *pringlei* to specific rank, but made no more changes to the taxonomy of this group. Because of a lack of material, Martin (1950) did not reclassify this species complex.

After his study of leaf architectural characters, Lis (1992) reorganized the *fothergilloides* complex to contain *C. fothergilloides*, *C. pringlei*, *C. mojadensis*, *C. macrophyllus* and *C. montanus* var. *traskiae*. Martin (1950) had placed *C. traskiae* as a variety of *C. montanus* based on overlapping traits between *C. traskiae* and *C. montanus* var. *glaber* and variety *blancheae*. The close association of *C. montanus* var. *traskiae* with the other Mexican taxa based on leaf architecture led Lis (1993) to raise *C. traskiae* to the specific level in his treatment in the Jepson Manual.

James Henrickson (in prep.) is currently revising the *Cercocarpus fothergilloides* complex and his treatment can be seen in Figure 1.1. Henrickson recognizes four species, *C. fothergilloides*, *C. pringlei*, *C. macrophyllus*, and *C. traskiae*. *Cercocarpus fothergilloides* is described containing three varieties, including the variety *mojadensis*, a taxon that Schneider (1905b) described at the rank of species. Henrickson also recognizes *C. pringlei* with two varieties, but notes that there are strong overlaps in many characteristics with *C. fothergilloides*. *Cercocarpus macrophyllus* is recognized as a distinct species by a large number of characteristics, but particularly by distinctively large leaf blades. Henrickson continues to recognize *Cercocarpus traskiae* as a distinct species, separate from *C. montanus*. The geographic ranges for these taxa are found in Figures 1.10.

Cercocarpus rzedowskii* and *C. mexicanus

Recently, two new species of *Cercocarpus*, *C. rzedowskii* and *C. mexicanus*, were recently described from northern Mexico (Figure 1.11) because of their very distinct leaf morphology (Henrickson 1987). Although Lis (1992) was unable to include these species into his analysis of *Cercocarpus*, Henrickson (1987) recognized similarity between *C. mexicanus* and the *montanus* complex, and noted *C. rzedowskii* had similarities to the *ledifolius* complex.

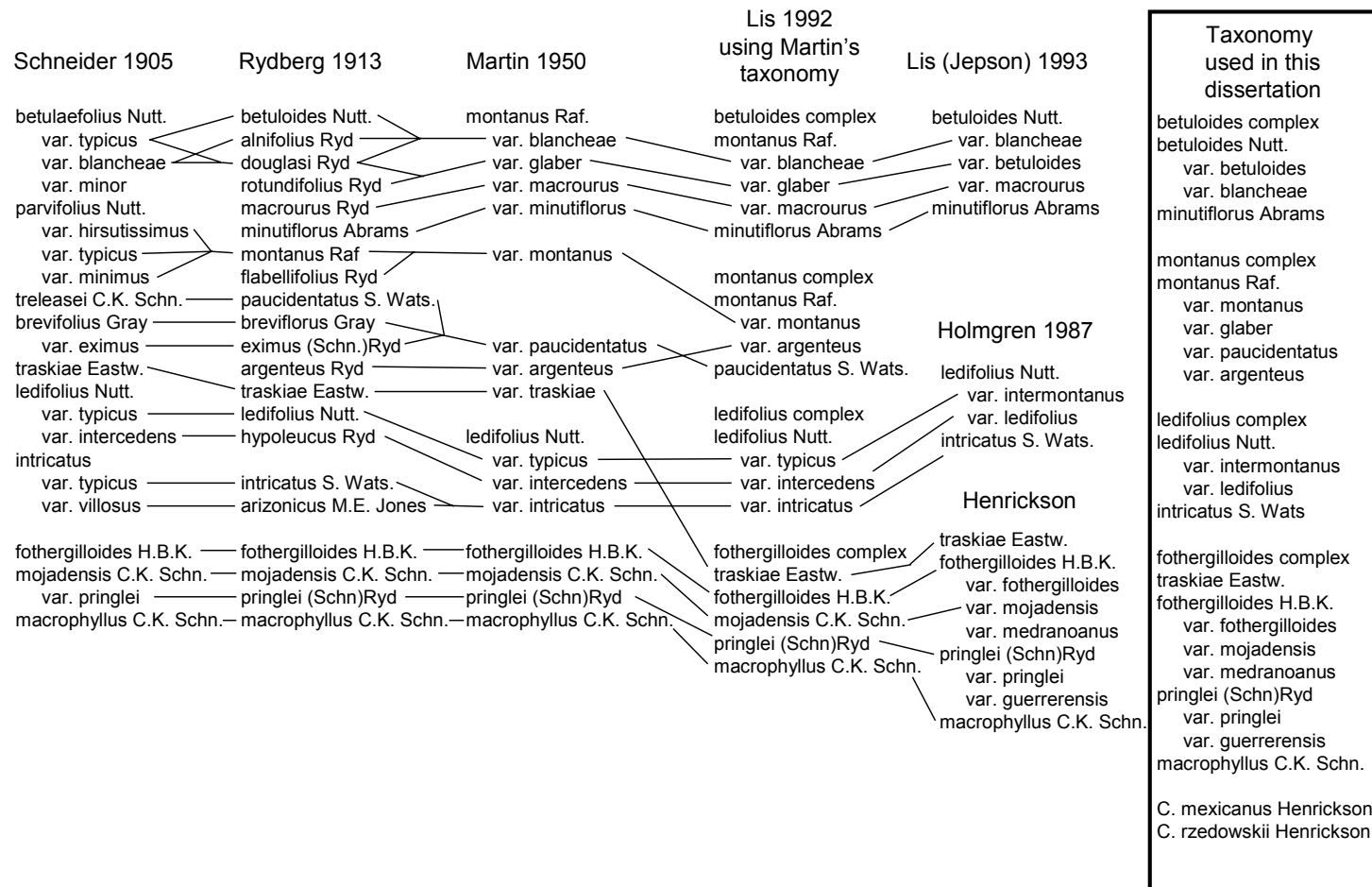


Figure 1.1. Representation of the different treatments of *Cercocarpus* H.B.K. (Rosaceae) and the taxonomy used in this dissertation, including the three complete revisions (Schneider 1905b, Rydberg 1913, and Martin 1950) and infrageneric revisions (Holmgren 1987, Lis 1993, and Henrickson in manuscript).

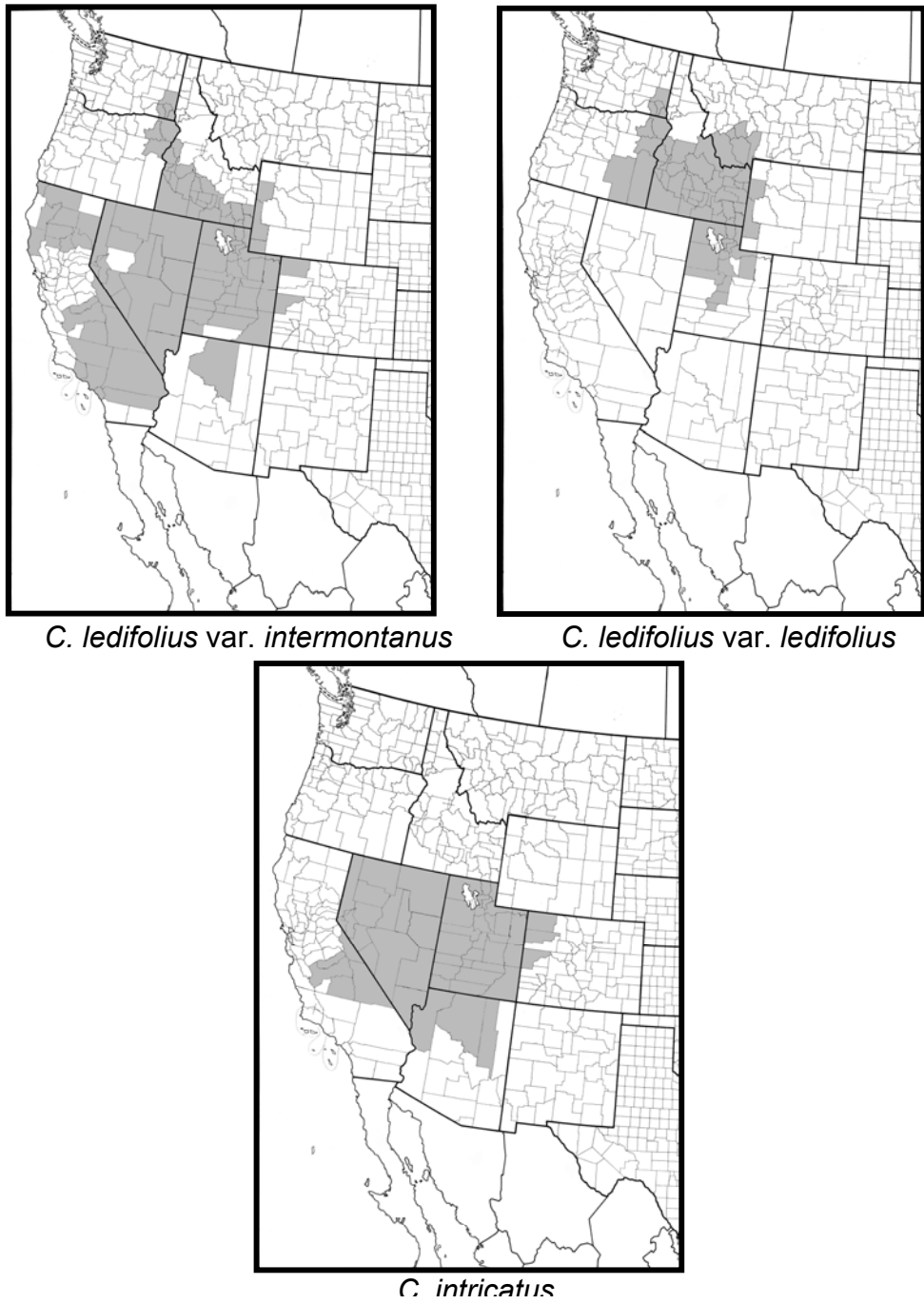


Figure 1.2. Geographic distribution of the *Cercocarpus ledifolius* complex

**The *ledifolius* complex:
Varieties of *C. ledifolius*
and *C. intricatus***



***C. ledifolius* var. *C. ledifolius* *C. intricatus*
intermontanus var. *ledifolius***

Figure 1.3. A picture of representative leaves for the taxa in the *Cercocarpus ledifolius* complex as delimited by Holmgren (1987). *C. ledifolius* var. *intermontanus* is found on the left, *C. ledifolius* var. *ledifolius* is found in the middle, and *C. intricatus* is found on the right. The black and white squares found on the bottom and to the left of the picture are square centimeters for scale.

Leaf diversity from sympatric populations



C. ledifolius var. *intermontanus* → *C. ledifolius* var. *ledifolius* → *C. intricatus*

Figure 1.4. A picture showing mature leaf variation found in a single population of *Cercocarpus ledifolius* from White Pine County, Nevada. The individuals displaying leaves on the right would be identified as *C. ledifolius* var. *intermontanus*. Individuals displaying leaves in the middle would be identified as *C. ledifolius* var. *ledifolius*, and individuals displaying leaves on the left would be identified as *C. intricatus*. The black and white squares found on the bottom and to the left of the picture are square centimeters for scale.

The *montanus* complex: Varieties of *C. montanus*

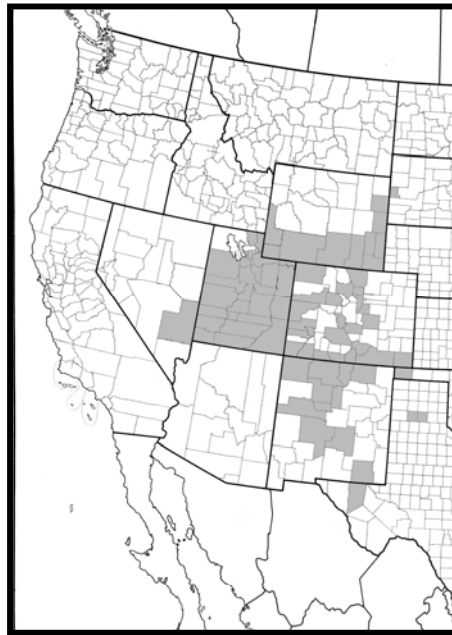


C. montanus var.
montanus

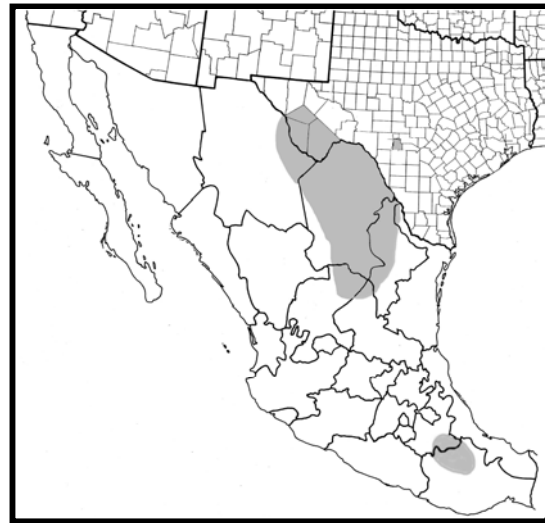
C. montanus
var. *glaber*

C. montanus var.
paucidentatus

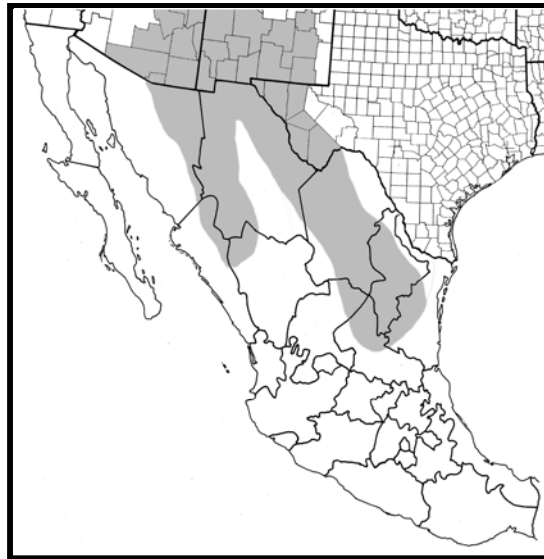
Figure 1.5. A picture of representative leaves for three of the four varieties in the *Cercocarpus montanus* complex as delimited by Martin (1950). *C. montanus* var. *montanus* is found on the left, *C. montanus* var. *glaber* is found in the middle, and *C. montanus* var. *paucidentatus* is found on the right. The black and white squares found on the bottom and to the left of the picture are square centimeters for scale.



C. montanus var. *montanus*



C. montanus var. *glaber*



C. montanus var. *paucidentatus*



C. montanus var. *argenteus*

Figure 1.6. Geographic distribution of the *Cercocarpus montanus* complex.

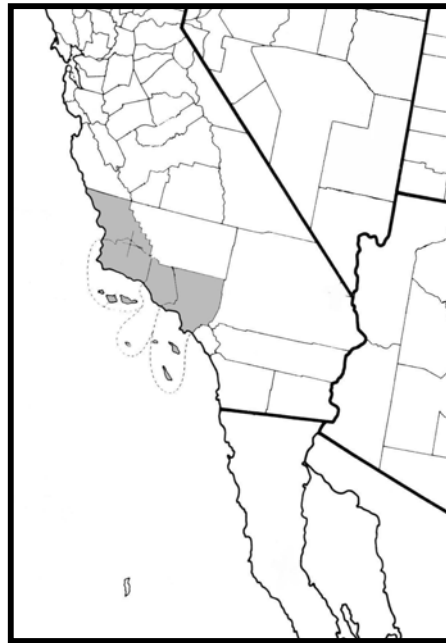
Leaf diversity from population 30
variety *glaber* → variety *paucidentatus*



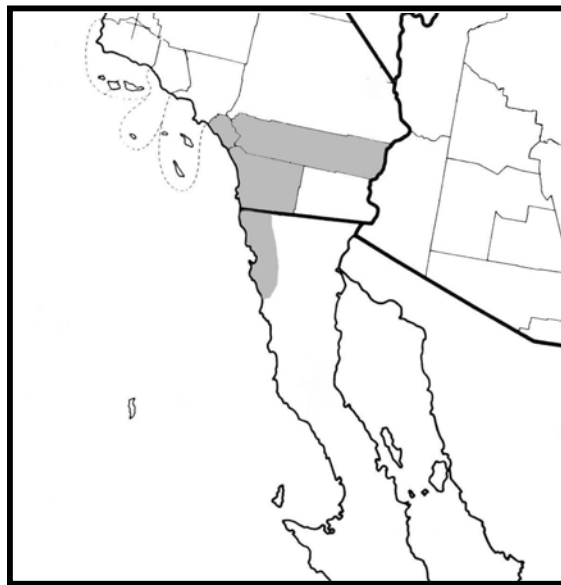
Figure 1.7. A picture showing mature leaf variation found in a single population of *Cercocarpus montanus* var. *paucidentatus* from Catron County, New Mexico. The black and white squares found on the bottom and to the right of the picture are square centimeters for scale.



C. betuloides var. *betuloides*



C. betuloides var. *blancheae*



C. minutiflorus

Figure 1.8. Geographic distribution of the *Cercocarpus betuloides* complex.

The *betuloides* complex: Varieties of *C. betuloides* and *C. minutiflorus*



C. betuloides var. *betuloides*

C. minutiflorus

C. betuloides var. *blancheae*

Figure 1.9. A picture of representative leaves for the taxa in the *Cercocarpus betuloides* complex as delimited by Lis (1993). *C. betuloides* var. *betuloides* is found on the left, *C. betuloides* var. *blancheae* is found in the middle, and *C. minutiflorus* is found on the right. The black and white squares found on the bottom and to the right of the picture are square centimeters for scale.

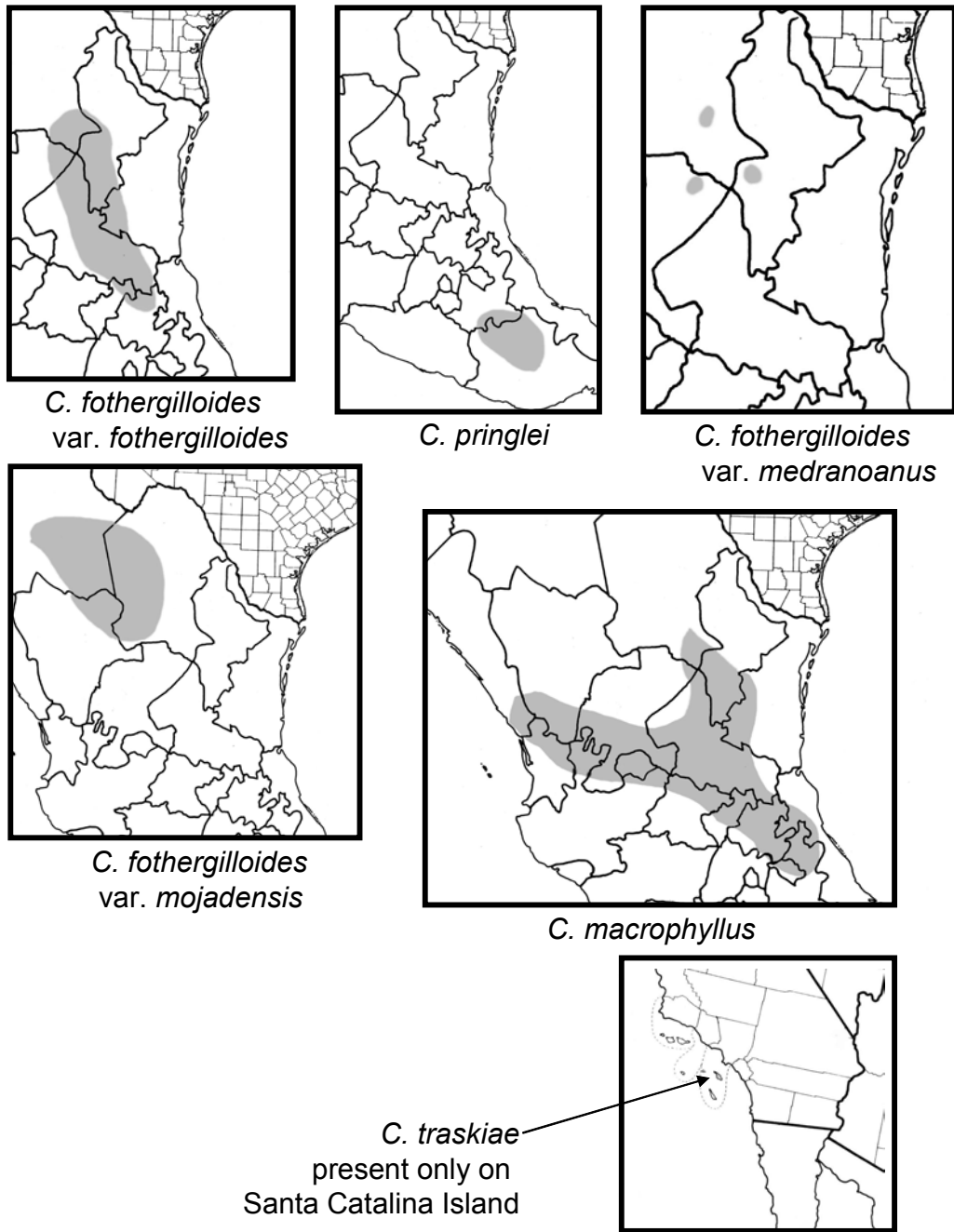


Figure 1.10. Geographic ranges for *Cercocarpus fothergilloides* complex.

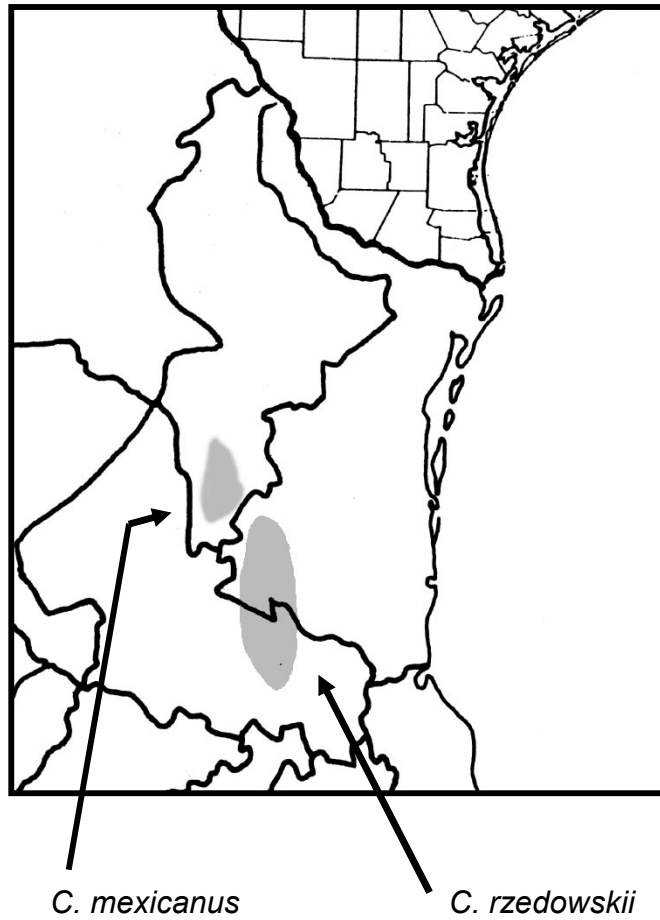


Figure 1.11. Geographic ranges for *Cercocarpus mexicanus* and *C. rzedowskii* (Henrickson 1987).

Chapter 2: Deep Coalescence of Multiple Paralogous Nuclear Ribosomal DNA Copies within Individuals of the Genus *Cercocarpus*

INTRODUCTION

Nuclear ribosomal DNA (nrDNA) is arranged in tandem arrays consisting of 100's of copies, that are generally located in the nucleolar organizing region (NOR), and can be found from one to several regions within the genome (Appels and Honeycutt 1986, Thompson and Flavell 1988). Within a single genome, the sequence similarity of each rDNA repeat is usually extremely high. This production of homogeneity between rDNA repeats is referred to as concerted evolution (Arnheim et al. 1980, Zimmer et al. 1980). Mechanisms which might drive concerted evolution include: gene conversion, slippage, transposition, or unequal crossing over (Arnheim et al. 1980, Dover 1982, Arnheim 1983).

Most attempts at phylogenetic reconstruction in plants using nuclear gene sequences have turned to nuclear rDNA. The reasons for using rDNA are well reviewed (Appels and Honeycutt 1986, Mindell and Honeycutt 1990, Hillis and Dixon 1992, Hamby and Zimmer 1992, Sanderson and Doyle 1992, Nickrent and Soltis 1995). These reasons include: 1) the ease of amplification due to the extremely high copy number and the tandem repeat structure coupled with the availability of universal primers; 2) the heterogeneity of rate of sequence evolution between different regions, which allows study at almost any taxonomic

level; and 3) the observation that the rDNA copies undergo rapid concerted evolution, which allow direct sequencing of PCR products, despite the extremely high copy number.

However, if concerted evolution does not eliminate all rDNA variation between rDNA repeats within or between arrays before a taxon speciates (i.e., if the rate of concerted evolution is slower than the rate of speciation) or, if two divergent species come together and form a hybrid, then descendent species will possess multiple rDNA repeats of divergent paralogues until concerted evolution homogenizes the repeat. In recent years the number of reports of paralogous rDNA repeats in certain plant species has increased (Dvorak 1990, Suh et al. 1993, Buckler and Holtsford 1996, O’Kane et al. 1996, Waters and Schaal 1996, Wendel et al. 1995, Sang et al. 1995, and Campbell et al. 1997). Paralogous rDNA repeats in an individual plant have usually been attributed to one of three causes; hybridization, allopolyploidization, and pseudogene formation. With hybridization and allopolyploidization, the paralogous rDNA repeats’ origins (i.e., the parental species’ rDNA types) can be identified and often lend insight into the diversity and evolutionary history of the taxon being studied. In the case of pseudogene formation, the functional rDNA repeats are usually identified by secondary structure stability analysis and base pair composition (Buckler et al. 1997). In this chapter, we present the first evidence for multiple, paralogous rDNA repeats with a deep coalescence that do not appear to be the product of

hybridization, allopoloidization, or pseudogene formation. Using the external transcribed spacer sequences (ETS) from the nuclear rDNA present in the taxa of the genus *Cercocarpus* (Rosaceae), we document at least three deep-coalescing copies of the ETS. We also document a complex evolutionary history for the ETS region in *Cercocarpus*.

MATERIALS AND METHODS

Collection of Plant Material and DNA Extraction

The systematics of *Cercocarpus* are largely unknown; the genus lacks a consensus classification or phylogeny. For this report, we followed the suggested taxonomy of Lis (1992) and Holmgren (1987) for the *Cercocarpus* taxa found in the United States. For the Californian taxa of *Cercocarpus*, we used the taxonomy found in the Jepson manual (Lis 1993). For the Mexican taxa of *Cercocarpus*, we used an unpublished treatment by J. Henrickson (Madroño ined.). We collected leaf material for 289 individuals, from 96 populations representing 16 taxa of *Cercocarpus* (Table 2.1). Leaves from five herbarium sheets are included in this total. The remainder of the samples consisted of field-collected leaf material, dried using silica gel for transport back to the lab. For each leaf sample, DNA was extracted from the collections using a modified CTAB procedure (Doyle and Doyle 1987). Genomic DNA extractions were

cleaned using the Elu-quick DNA purification kit (Schleicher and Schuell) prior to PCR amplification.

Developing the ETS in *Cercocarpus*

To develop the ETS region in *Cercocarpus* for routine PCR amplification, we amplified the entire intergenic spacer (IGS) of *C. montanus* var. *montanus* (population #5) and *C. ledifolius* var. *intricatus* (population #25) (Table 2.2) using the primers *1M* and *18S-2L* (Linder et al. 2000) (Table 2.3) located in conserved regions of the 26S and 18S genes, respectively (Figure 2.1). A negative control was also used that lacked template DNA. Fifty microliter PCRs were performed using 30mM Tricine, pH 8.4, 2mM MgCl₂, 50 mM KCl, 5% acetamide, 100 uM of each dNTP, 1.0 unit of Taq polymerase, and 30 nM of each amplification primer. Twenty nanograms of template DNA were added, and the reactions were overlaid with mineral oil and run on a MJ Research Thermal Cycler programmed for a hot start (95° C, 5 min; 74° C, 7 min) and 30 cycles of 94° C for 30 sec, 50° C for 45 sec, and 72° C for 2 min. A final extension of 72° C for 7 min was also included. Three microliters of the reaction products were run on a 1.5 % agarose gel in 0.5X TBE buffer. PCR products were visualized with ethidium bromide under UV light. The IGS PCR products from *C. montanus* var. *montanus* (population #5) and *C. ledifolius* var. *intricatus* (population #25) were purified using the QIAquick PCR purification kit (QIAGEN) and were sequenced from their 3' end on an ABI Prism 377 automated sequencer using standard

reagents. Because the ETS region was too long to sequence with a single primer, we developed an internal primer *Cerc-ETS1r* (Table 2.3) (Figure 2.1) to sequence through the transcription initiation site (TIS). We identified the TIS based on similarity with the TISs in other plant groups (Linder et al. 2000). As a result of its role in transcription of the rDNA repeat, the TIS is under strong selection to stay conserved, which makes it a good choice for priming throughout *Cercocarpus*. This approach has been successful in other studies (Linder et al. 2000). Both sequences were proofread and then aligned using Clustal W in MegAlign (DNASTar, 1998), followed by the usual adjustment. We designed *Cerc-ETS1f* (Table 2.3) to include the TIS and to have high sequence similarity to both taxa (Figure 2.2).

Table 2.1: Taxa and Populations used to study ETS type variation in *Cercocarpus*.

Taxon	Populations sampled	Individuals sampled	Individuals w/ successful amplifications	ETS types and number of individuals
<i>C. montanus</i> var. <i>montanus</i>	9	26	26	1(26)
<i>C. montanus</i> var. <i>paucidentatus</i>	20	57	56	1(50) 3(3) 1 and 3(3)
<i>C. montanus</i> var. <i>glaber</i>	4	17	16	1(14) 1 and 2(1) 1 and 3(1)
<i>C. montanus</i> var. <i>argenteus</i>	1	1	1	1(1)
<i>C. betuloides</i> var. <i>betuloides</i>	14	44	42	1(28) 2(9) 4(2) 1 and 2(3)
<i>C. betuloides</i> var. <i>blancheae</i>	5	15	12	1(2) 2(7) 3 and 4 (1) 4(2)
<i>C. minutiflorus</i>	3	7	6	1 and 2 (5) 2(1)
<i>C. traskiae</i>	2	19	19	1(1) 1 and 2(3) 2(12) 3(1) 2 and 4(1) 4(1)
<i>C. fothergilloides</i> var. <i>fothergilloides</i>	8	27	23	1(3) 3(18) 3 and 4(2)
<i>C. fothergilloides</i> var. <i>mojadensis</i>	1	1	1	2(1)
<i>C. fothergilloides</i> var. <i>medranoanus</i>	2	6	6	1 and 3(2) 3(4)
<i>C. macrophyllus</i>	3	4	4	1(1) 1 and 2(1)

				1 and 3(2)
<i>C. rzedowskii</i>	3	12	11	1 and 4(5) 4(6)
<i>C. ledifolius</i> var. <i>ledifolius</i>	3	5	4	1(4)
<i>C. ledifolius</i> var. <i>intermontanus</i>	13	35	34	1(34)
<i>C. intricatus</i>	5	15	15	1(15)
Total	96	289	275	1(179) 2(29) 3(26) 4(11) 1 and 2(13) 1 and 3(8) 1 and 4(5) 2 and 4(1) 3 and 4(3)

We successfully amplified 9 samples (populations 8, 12, 14, 15, 17, 18, 19, 20, and 21) (Table 2.2) using the *Cerc-ETS1f* and the *18S-2L* primers. After sequencing the 9 amplifications using the procedures outlined above, we aligned all 9 sequences plus *C. montanus* var. *montanus* (population #5) and *C. intricatus* (population #25) and noticed polymorphisms and small indels in the sequences near the *Cerc-ETS1f* priming site (Figure 2.2). We, therefore, designed another forward primer, *Cerc-ETS2f* (Table 2.3) (Figure 2.2), 25 bp downstream of *Cerc-ETS1f* in a region that was completely conserved in all sequences. We used *Cerc-ETS2f* for all further ETS amplifications. We attempted amplification using *Cerc-ETS2f* on all 289 individuals (Table 2.1). Amplification was as described above for the IGS except for a 1 min. 30 sec. extension at 72° C during the 30 cycles.

Three microliters of the reaction mixture were run on a gel and visualized as described above. Of the 289 samples, 57 failed to amplify, so we attempted the PCR again using the same primers with the FailSafe PCR system (Epicentre Technologies). Twenty five microliter reactions were performed using 12.5 uL of premix A through L, 30 nM of each primer, 1 unit of Taq, and 20 ng of template DNA. Each reaction was overlaid with 30 uL of mineral oil. Amplification was the same as detailed above.

Table 2.2: *Cercocarpus* individuals whose ETS amplification was sequenced. Individuals with * denote the seven individuals used in the probing experiment. † denotes the 14 individuals who were included in the reduced dataset. BVH = Brian Vanden Heuvel, JH = James Henrickson, BC = Bonnie Crozier, Sand = Sanderson

<i>Taxon</i>	Population # and individual	Location and collection number	ETS types
<i>C. montanus</i> var. <i>montanus</i>	5	Douglas Co., CO BVH 5	1
	8	Platte Co., WY BVH 8	1
	10	Custer Co., SD BVH 10	1
	28A	Otero Co., NM BVH 28	1
<i>C. montanus</i> var. <i>paucidentatus</i>	†1	Tamaulipas, MX JH 19218	1
	27C	Eddy Co., NM BVH 27	1
	30A	Grant Co., NM BVH 30	1
	*86B	Bernalillo Co., NM BVH 86	1,3
	87A	Sierra Co., NM BVH 87	4
	87B	Sierra Co., NM BVH 87	3
	90D	Catron Co., NM BVH 90	1,3
	<i>C. montanus</i> var. <i>glaber</i>	3	Coahuila, MX JH 19218
12		Brewster Co., TX JH 20663	1
14		Brewster Co., TX JH 20665	1
15		Brewster Co., TX JH 20666	1
17		Coahuila, MX JH 20637	1
58B		Real Co., TX JH 22275	1
68A		Coahuila, MX BVH and JH 22414	1
75A		Tamaulipas, MX BVH and JH 22429	1
80A	Tamaulipas, MX BVH and JH	1	

22457			
	85A	Coahuila, MX JH 22587	1
<i>C. montanus</i> var. <i>argenteus</i>	131	Culberson Co., TX BVH 131	1
<i>C. betuloides</i> var. <i>betuloides</i>	19	Los Angeles Co., CA BVH 123a	2
	20	Los Angeles Co., CA BVH 123b	2
	†21	Los Angeles Co., CA BVH 123c	2
	62D	Tulare Co., CA BVH 127a	2
	†63B	Tulare Co., CA BVH 127a	4
	†65C	Tulare Co., CA BVH 127b	2
	65D	Tulare Co., CA BVH 127c	2
	*93C	Gila Co., AZ BVH 93	1,2
	118A	Monterey Co., CA BVH 118	2,4
	*120C	San Luis Obispo Co. CA BVH 120	1
<i>C. betuloides</i> var. <i>blancheae</i>	121B	Santa Barbara Co., CA BVH 121	3,4
	121C	Santa Barbara Co., CA BVH 121	2
	122A	Santa Barbara Co., CA BVH 122	2
	123F	Los Angeles Co. CA, BVH 123	2
	123G	Los Angeles Co. CA, BVH 123	1
	126.BC1	Los Angeles Co. CA	2
<i>C. minutiflorus</i>	57B	San Diego Co., CA JH 22311	1,2
	61	San Diego Co., CA JH 22313	1
<i>C. traskiae</i>	52B	Los Angeles Co., CA JH 22312	2
	126A	Los Angeles Co.	2
	126E	Los Angeles Co.	1,2
	†126.29	Los Angeles Co.	2,4
	126.WB3	Los Angeles Co.	4
<i>C. fothergilloides</i> var. <i>fothergilloides</i>	66C	Coahuila, MX BVH and JH 22411	3
	66D	Coahuila, MX BVH and JH 22412	3
	67A	Coahuila, MX BVH and JH 22413	3
	*67B	Coahuila, MX BVH and JH 22413	1
	†71A	Coahuila, MX JH and BVH 22423	3
	†83B	Tamaulipas, MX JH and BVH 22464	3
	*†84A	Nuevo Leon, MX JH and BVH 22482	4
<i>C. fothergilloides</i> var. <i>mojadensis</i>	†18	Coahuila, MX JH 20639	2
<i>C. fothergilloides</i> var. <i>medranoanus</i>	†77B	Tamaulipas, MX BVH and JH 22444	3
	82A	Tamaulipas, MX BVH and JH 22456	1,3
<i>C. macrophyllus</i>	26	Durango, MX BC n/a	1,2
	69A	Coahuila, MX BVH and JH 22415	1
	69B	Coahuila, MX BVH and JH 22415	1,3
<i>C. rzedowskii</i>	76A	Tamaulipas, MX BVH and JH 22434	1,4
	†76E	Tamaulipas, MX BVH and JH	1,4

		22434	
	78A	Tamaulipas, MX BVH and JH 22446	1
	78B	Tamaulipas, MX BVH and JH 22446	4
	*†81A	Tamaulipas, MX BVH and JH 22460	4
<i>C. ledifolius</i> var. <i>ledifolius</i>	41B	Emery Co., UT BVH 41	1
	104	White Pine Co., NV BVH 104	1
<i>C. ledifolius</i> var. <i>intermontanus</i>	45C	Box Elder Co., UT BVH 45	1
	98A	Sierra Co., CA BVH 98	1
	101A	Lander Co., NV BVH 101	1
	102B	Elko Co., NV BVH 102	1
	108C	Clark Co., NV BVH 108	1
	111A	Mono Co., CA BVH 111	1
<i>C. intricatus</i>	*†25	Garfield Co., UT BVH 25	1
	50D	Uintah Co., UT BVH 50	1
	103A	White Pine Co., NV BVH 103	1
<i>Purshia tridentata</i>	†34	La Plata Co., CO BVH 34	Purshia
<i>Purshia plicata</i>	74	Coahuila, MX BVH and JH	Purshia

Sequencing the ETS

We sequenced 102 ETS amplifications representing 76 individuals (Table 2.2), using a sampling strategy to maximize the geographic distribution and ETS length variants for each *Cercocarpus* taxon (Figure 2.3). One of two strategies was employed for sequencing the ETS for each sample. If the ETS amplification produced a single PCR product, the product was sequenced directly as described above. If the ETS amplification produced multiple-length PCR products, we cloned the PCR products using the TOPO-TA cloning kit (Invitrogen). At least 10 clones per amplification were screened for the ETS insert by direct PCR

amplification using the *Cerc-ETS2f* and *18S-2L* primers and the same PCR parameters listed above, except for a 10 min. hot start at 95° C to lyse the cells from the clones. The amplified inserts were visualized on an agarose gel as above. Two to six of these PCR amplifications of cloned material for each ETS length variant were purified (QIAquick PCR purification kit, QIAGEN) and sequenced per sample. Sequencing was performed on an ABI Prism 377 or an ABI Prism 3100 automated sequencer using standard reagents. Sequences were proofread and then aligned using Clustal W in MegAlign (DNASTar, 1998), followed by adjustment by eye.

Table 2.3: Sequences of the primers used in PCR for this study

Primer	Sequence 5'-3'	Direction
<i>1M</i>	TTGCTGCCACCATCCACTGAGAT	Forward
<i>18S-2L</i>	TGACTACTGGCAGGATCAACCAG	Reverse
<i>Cerc-ETS1f</i>	TATAAAGGGGGAGGCCTCATT	Forward
<i>Cerc-ETS2f</i>	TACAAACATATTCTCCCATATCTC	Forward
<i>Cerc-ETS1r</i>	GGCCCAATCTCCTTTCATGTCACT	Reverse
<i>Pur-ETS1r</i>	CGCACGCACAAGGCAGAAGG	Reverse
<i>Pur-ETS2r</i>	CACCGGTGGGCTGACGAAAGG	Reverse
<i>ETS Type 1f</i>	GTTGGTATCAGAGCAGGATCGATATGTC	Forward
<i>ETS Type 1r</i>	GAATCATCGCAGCTCCTCGGGGTGC	Reverse
<i>ETS type 2f</i>	GCCCACCCATGCAATGCACAAAGC	Forward
<i>ETS type 2r</i>	GATCGAGACACTGGCACCCTATCGC	Reverse
<i>ETS type 3f</i>	GATTGGATCATATTGATGAATACCC	Forward
<i>ETS type 3r</i>	GCGGATGGAGATGGTGGTAGGCTGGC	Reverse

After alignment with Clustal W, we observed four length variants and designated them ETS type 1 (1000 bp), ETS type 2 (800 bp), ETS type 3 (750 bp), and ETS type 4 (500 bp). We found large regions (80-600 bp) that were unalignable between ETS length variant types (Figure 2.3). We searched these

putative unalignable regions for conserved alignable regions using pair-wise dot-plot comparisons in MegAlign (minimum window = 20 bp; >70% similarity), and employing representatives for each ETS length variant type in the analyses. We used the ETS sequence from *C. montanus*. var. *paucidentatus* (pop#1) to represent the ETS type 1 sequences, the ETS sequence *C. montanus* var. *blancheae* (pop#21) to represent ETS type 2 sequences, the ETS sequence *C. fothergilloides* var. *mojadensis* (pop#66) to represent ETS type 3 sequences, and the ETS sequence *C. rzedowskii* (pop#81) to represent type 4 sequences (Table 2.2).

The ETS sequences for *Purshia tridentata*, and *Purshia plicata* were obtained to serve as outgroups for this study. *Purshia* has consistently been identified as the sister group to *Cercocarpus* in recent molecular phylogenies (Morgan et al. 1994, Evans et al. 2000). The complete IGS in these species was amplified as described above for *Cercocarpus*. Then we sequenced from the 3' end of the amplified product using *18S-2L* and the reverse internal primers *Pur-ETS1r* (5'-CGCACGCACAAGGCAGAAGG-3') and *Pur-ETS2r* (5'-CACCGGTGGGCTGACGAAAGG-3') until we identified the TIS.

Testing Coalescence of the ETS Types found in *Cercocarpus*

After identifying four ETS types in *Cercocarpus*, we created a gene tree to examine the origin and historical relationships between the ETS types. All alignable regions from the ETS sequences, regardless of type, were aligned using Clustal W, followed by usual adjustment. This included the 5' alignable, 3'

alignable, and the α and β regions (Figure 2.3). For ETS type 3 sequences, only the 5' common region could be included in the aligned sequence matrix. Only the 3' common region of the ETS sequences of *P. tridentata* and *P. plicata* could be included to serve as the outgroups (Figure 2.3). Therefore, ETS type 3 sequences' placement in the phylogeny were based only on their relationship to the other ETS types and not on a direct comparison to the outgroup. All phylogenetic analyses were carried out in PAUP 4.0.0d64 (Swofford 1999). Maximum parsimony (MP) heuristic searches were conducted using the following options: ACCTRAN, gaps treated as missing data, sequences added by stepwise addition, TBR, unordered, and equal weight characters. Support for branches were found using 100 bootstrap replicates. A maximum likelihood (ML) heuristic search was allowed to TBR branch swap for 240 hours, using the HKY85 model, adding sequences by stepwise addition as is and treating gaps as missing data.

Because of the computational time that a ML analysis on a large dataset requires, we created a reduced dataset containing only 14 sequences to examine ETS type evolution fully using the ML optimality criterion. The identity of the 14 sequences can be found in Table 2.2. We conducted two main sets of analyses on this dataset, which differed in the character sets included in the analyses. The first set of analyses included the 5' alignable, 3' alignable, and the α and β regions (511 aligned base pairs). We conducted ML searches using the Jukes-Cantor, Kimura-2, and HKY85 models, adding sequences as-is and using TBR branch

swapping. Then we conducted the likelihood ratio test to find the parameter set that gave us the greatest fit. A bootstrap analysis was also conducted using 100 replicates. Maximum parsimony was also conducted on this reduced dataset as described above. The second set of analyses included just the 5' alignable and 3' alignable regions (382 aligned base pairs). We executed the same ML and MP searches as described above.

Probing for the different ETS types

We tested whether the different ETS length types recovered after amplifications were all part of the rDNA repeat rather than an artifact of non-specific priming outside of the rDNA repeat. Seven individual leaf collections were chosen based on the ETS type they contained (Table 2.2). Total DNA was extracted from these seven samples, using 3-5 grams of silica dried leaf material in a modified CTAB extraction procedure (Doyle and Doyle 1987). The samples were further purified using Cesium chloride/ethidium bromide gradient ultracentrifugation (Jansen and Palmer 1987). Total DNA was digested with EcoRI, which is known to cut infrequently in the rDNA repeat (Jourgensen et al. 1987). Restriction enzyme digestion, electrophoresis, bi-directional blotting to Zetabind nylon (AM CUNO), labeling of recombinant plasmids by nick translation, filter hybridizations, and autoradiography were performed following Palmer (1986) and Jansen and Palmer (1987).

To probe for the 5' end of the 18S gene, we used the 1.3 kb BamHI/HindIII rDNA fragment (Jourgensen et al. 1987) from *Pisum* sp. (kindly provided by Bob Jansen). The ETS type 1 probe was constructed by PCR amplifying 600 bp of the unalignable region of *C. betuloides* var. *betuloides* pop # 93C clone 2. Amplification was accomplished using *ETS Type 1f* and *ETS Type 1r* primers (Table 2.3). This PCR product was then placed into a plasmid using the TOPO TA kit as described above. The ETS type 2 probe was constructed in the same manner using a 350 bp fragment from the type 2 unalignable region. It was amplified using *ETS Type 2f* and *ETS Type 2r* primers (Table 2.3) from *C. betuloides* var. *betuloides* pop # 93C clone 1. The ETS type 3 probe was constructed using a 250 bp fragment from the ETS type 3 unalignable region. It was amplified using *ETS Type 3f* and *ETS Type 3r* primers (Table 2.3) from *C. fothersgilloides* var. *fothersgilloides* pop # 67A clone 3. We did not construct a probe specifically for ETS type 4 because of its sequence-similarity to ETS type 2.

RESULTS

Amplification and Identity of the ETS types in *Cercocarpus*

We successfully amplified the ETS in 275 of the 289 individuals using the *Cerc-ETS2f* primer (Table 2.1) (Table 2.3). We amplified 43 of these samples using premix K from the FailSafe system. Of the 14 samples that failed, 3 were

from herbarium sheets and had very low template DNA concentrations, and the remaining 11 samples that failed contained high levels of polysaccharides even after Elu-quik DNA purification. The 14 samples that failed to amplify using both the Tricine/acetimide and FailSafe systems were not taxon specific, but were distributed fairly evenly among all *Cercocarpus* taxa (Table 2.1). They included one individual of *C. montanus* var. *paucidentatus*, one individual of *C. montanus* var. *glaber*, two individuals of *C. betuloides* var. *betuloides*, three individuals of *C. betuloides* var. *blancheae*, one individual of *C. betuloides* var. *minutiflorus*, four individuals of *C. fothergilloides* var. *fothergilloides*, one individual of *C. rzedowskii*, and one individual of *C. ledifolius* var. *ledifolius*.

The distribution of the four different ETS length types (ETS type 1, type 2, type 3, and type 4) amplified can be seen in Table 2.4. ETS type 1 was the most frequent, followed sequentially by type 2, type 3, and type 4. Twenty nine amplifications contained multiple ETS types. The most frequent ETS type combination was ETS type 1 and 2, followed by ETS type 1 and type 3 and ETS type 1 and type 4. The least frequent combination of ETS types was ETS type 2 and type 4.

Table 2.4: Overall distribution of ETS types in individuals sampled in *Cercocarpus*

ETS types	Number of individuals
Only having ETS type 1	180
Only having ETS type 2	29
Only having ETS type 3	26

Only having ETS type 4	11
ETS type 1 and 2	12
ETS type 1 and 3	8
ETS type 1 and 4	5
ETS type 2 and 4	1
ETS type 3 and 4	3

The taxonomic distribution of the ETS types is illustrated in Table 2.5. Five taxa were found to only contain ETS type 1. These five taxa are all found in the Rocky Mountains and the Great Basin. The Californian taxa of *C. betuloides* var. *blancheae* and *C. traskiae* contained all four ETS types. The taxa that contain ETS type 3 all have Mexico as part of their distribution.

Table 2.5: Taxonomic distribution of ETS types in individuals sampled in *Cercocarpus*

ETS types	<i>Cercocarpus</i> taxa
Only having ETS type 1	<i>C. montanus</i> var. <i>montanus</i> , <i>C. montanus</i> var. <i>argenteus</i> , <i>C. ledifolius</i> var. <i>intermontanus</i> , <i>C. ledifolius</i> var. <i>ledifolius</i> , <i>C. intricatus</i>
Only having ETS type 2	<i>C. fothergilloides</i> var. <i>mojadensis</i>
ETS type 1 and 2	<i>C. minutiflorus</i>
ETS type 1 and 3	<i>C. montanus</i> var. <i>paucidentatus</i> , <i>C. fothergilloides</i> var. <i>medranoanus</i>
ETS type 1 and 4	<i>C. rzedowskii</i>
ETS type 1, 2, and 3	<i>C. macrophyllus</i> , <i>C. montanus</i> var. <i>glaber</i>
ETS type 1, 2, and 4	<i>C. betuloides</i> var. <i>betuloides</i>
ETS type 1, 3, and 4	<i>C. fothergilloides</i> var. <i>fothergilloides</i>
ETS type 1, 2, 3, and 4	<i>C. betuloides</i> var. <i>blancheae</i> , <i>C. traskiae</i>

Structure and Alignment of the ETS types

We found that distinct portions of each ETS type were alignable between ETS types (Figure 2.3). The 5' alignable region was 174 bp long, and the 3' alignable region was 207 bp long. ETS type 1, type 2, and type 4 contained both the 5' and 3' alignable regions. ETS type 3 only contained the 5' alignable region. The *Purshia* ETS type contained only the 3' alignable region.

Dot plot analyses revealed two small alignable regions embedded in the unalignable regions of ETS type 1, ETS type 2, and the *Purshia* ETS type. We designated them as α and β . The α region was 49 bp long in all ETS types which contained it. The β region was 80 bp in ETS type 1 and the *Purshia* ETS type. ETS type 2 had 34 bp of the 3' end of the α region. ETS type 1 contained two copies of the α region while ETS type 2 and the *Purshia* ETS type contained only 1 (Figure 2.3).

The five ETS types also contained different unalignable regions. The unalignable region in ETS type 1 was 499 bp and was upstream of the α and β region (Figure 2.3). The unalignable region in ETS type 2 totaled 314 bp and was found downstream of the α and β region. ETS type 3 contained an unalignable region that totaled 510 bp and stretched from the 5' alignable region all the way to the 18S gene. ETS type 4 contained only 100 bp of the extreme 3' end of the unalignable region found in ETS type 2. The *Purshia* ETS type contained the full ETS type 2 unalignable region and an unalignable region specific to *Purshia* that totaled 729 bp and stretched upstream from the α region to the TIS.

Relationships of the alignable regions of the ETS types

The sequence matrix of 5', 3', and α and β alignable regions totaled 511 aligned base pairs for 102 ETS sequences (Appendix A). The MP analysis resulted in over 10,000 most parsimonious trees of length 530. Figure 2.4 is a

strict consensus based upon those trees. The *Cercocarpus* ETS types form a clade with high bootstrap support when compared to the outgroup sequences' indicating that the different *Cercocarpus* ETS types likely had their origin within the genus. ETS type 1 and ETS type 3 each form weakly supported clades, indicating a single coalescence or origin for both types (arrows in Figure 2.4). ETS types 2 and 4 form a "phylogenetic grade," with ETS type 3 basally nested within, suggesting that ETS type 3 may have had an early origin from one of those two ETS types. The ML tree (Figure 2.5.) is a strict consensus tree of five most-likely trees. The ML tree agrees with the MP tree in that the ETS type 1 and ETS type 3 clades are monophyletic, and the ETS type 2 and type 4 sequences form a grade. The ML tree differs from the MP consensus tree in the placement of some of the type 2 and type 4 sequences. The differences between the two reconstructions occur on branches with very little bootstrap support in the MP tree. Bootstrap support for particular reconstructions in the ML trees based on the full dataset were not calculated due to computational limitations.

The results of the ML and MP searches on the reduced dataset of 14 sequences can be seen in Figure 2.6. For the first analysis, where the α and β regions are included, we found that the MP and ML analysis gave the same tree topology. Both analyses supported the basal position of ETS type 1 sequences with moderate to high support. The analyses do not support the monophyly of ETS type 2 or 4 (Figure 2.6). When the α and β regions were excluded from the

analyses, the MP and ML analyses resulted in more than 1 tree. The strict consensus trees from the two analyses were congruent. Once again, ETS type 2 and 4 are not monophyletic. In this case, the ETS type 1 clades and the clade including *C. fothergilloides* var. *mojadensis* (18) and *C. fothergilloides* var. *mojadensis* (83B) are unresolved at the base of the tree, sister to the rest of the taxa. In both analyses with the reduced dataset, the members of the ETS type 4 clade found in the basal position in the MP and ML consensus trees from the full dataset are found nested within the type 2 and 3 clade with moderate bootstrap support.

Probing

The results from the probing can be seen in Table 2.6. Only one band was produced on our filters after sequentially hybridizing with three probes. The ETS type 3 probe failed during the labeling reaction and produced negative results. The size of the band was identical for each individual between probes. The size of the band varied between individuals from 8.0 kb to 8.7 kb (Table 2.6).

Table 2.6: Results of the ETS type probing experiment. Listed are four ETS probes hybridized to seven individuals. Numbers in columns refer to the size of the band in kilobases.

Individual	18S probe	ETS type 1 probe	ETS type 2 probe	ETS type 3 probe
<i>C. intricatus</i> (25)	8.3	8.3	8.3	n/a
<i>C. bet.</i> var. <i>betuloides</i> (120C)	8.7	8.7	8.7	n/a
<i>C. bet.</i> var. <i>betuloides</i> (93C)	8.3	8.3	8.3	n/a
<i>C. foth.</i> var. <i>fothergilloides</i> (84A)	8.0	8.0	8.0	n/a
<i>C. mot.</i> var. <i>paucidentatus</i> (86B)	8.7	8.7	8.7	n/a
<i>C. foth.</i> var. <i>fothergilloides</i> (67B)	8.3(weak)	8.3(weak)	8.3(weak)	n/a
<i>C. rzedowskii</i> (81A)	8.7	8.7	8.7	n/a

DISCUSSION

ETS variation in *Cercocarpus*

We have demonstrated the occurrence of four ETS types in the taxa of *Cercocarpus* that differ from one another both in length and nucleotide substitution. The occurrence of variation, both nucleotide substitutions and length variation, in the ETS region in vascular plants is nothing new (Bena et al. 1998, Markos and Baldwin 1999, Linder et al. 2000). Many previous studies have documented both types of variation occurring both inter-specifically and intra-specifically. In fact, this variation has been harnessed successfully for

phylogenetic reconstruction (Bena et al. 1998, Markos and Baldwin 1999, Linder et al. 2000). What is unique about the ETS variation reported here in *Cercocarpus* is the nature of the variation.

In sharp contrast to the lack of variation in length of the coding regions and the ITS in the rDNA within an individual or a species, the length of the ETS has been shown to have inter-specific or even intra-specific variation (Linder et al 2000, Markos and Baldwin 1999). To date, both inter-specific and intra-specific variation in ETS length has been attributed to the presence of varying numbers of tandem sub-repeats. In *Cercocarpus*, we have demonstrated length variation in the ETS that is not due to varying numbers of sub-repeats. Instead, the length variation we have observed is caused by the occurrence of unalignable regions of different length present in the center of the ETS. What is perhaps the most interesting about the placement of the unalignable regions is the conservation of the length of the 5' and 3' alignable regions surrounding the unalignable regions in all four ETS types. ETS type 3 only contains the 5' alignable region, yet the length is still conserved to the same base pair as in the other ETS types.

Why the length of the 5' and 3' alignable regions is so conserved between ETS types is still a mystery. One possibility we explored for the conservation of the 5' and 3' alignable regions is that they are a product of a transposable element, short interspersed elements (SINE), or long interspersed elements (LINE). We were unable to find any nucleotide sequence evidence for footprints of known

transposons in the 5' or 3' alignable region or in the unalignable regions. We also conducted BLAST searches in GenBank for all of the different regions in all the ETS types and did not uncover any probable matches to known transposons. Another possibility for the conserved nature of the alignable regions may be that the inherent secondary structure of the ETS is maintaining the length of the alignable regions in a stem, and the unalignable regions are found in a loop and therefore less constrained. To explore this possibility, we submitted ETS sequences of each type to folding programs that use minimum free energy. We were unable to find any evidence for stems or loops that were correlated with the base pair positions of the alignable and unalignable regions.

Are some of the ETS types pseudogenes?

Plant ribosomal pseudogenes have been identified in the Winteraceae (Suh 1993), *Gossypium* (Wendel et al. 1995), *Tripsacum* (Buckler and Holtsford 1996), *Zea* (Buckler and Holtsford 1996), and *Nicotiana* (Buckler et al. 1997). Pseudogenes are identified in rDNA most often using ITS sequences that have low GC content, high substitution rate in conserved motifs, and low thermodynamic stability where certain conserved secondary structures are predicted (Buckler et al. 1997). Paralogues due to pseudogene formation are also diagnosed by large indel events that radically change the secondary structure (Mayol and Rossello 2001). As described above, we were unable to find any obvious evidence for secondary structure features in the ETS sequences in

Cercocarpus. There is no known secondary structure for the ETS to test against or a predicted base composition for the ETS. The only sequence motif that is known to be present in a functional ETS sequence is the TIS, which all four ETS types we recovered do possess.

Based on the probing data and the priming location of the *18S-2L* primer within the 5' end of the 18S gene, we have good evidence that at least three of the four ETS types recovered are located in the same fragment as the 5' end of the 18S gene and are not the product of non-specific priming outside the rDNA. Also, we identified individuals who possess only one of each of the four ETS types. If any one of the ETS types were a pseudogene, we would not expect to find individuals possessing only that ETS type. We would expect those individuals not to survive due to non-functional ribosomes. Therefore, it seems that each ETS type is part of a functional rDNA copy.

Are the multiple ETS types a product of hybridization or allopolyploidization?

In past studies, when multiple paralogous rDNA repeats in individuals were attributed to hybridization or allopolyploidization, each parental rDNA type was identified by comparison to “pure” parental individuals (Rieseberg et al. 1990, Soltis et al. 1991, Sang et al. 1995). These “pure” parental individuals can be identified because concerted evolution has homogenized the rDNA repeats within the individuals of the parental taxon. The occurrence of multiple rDNA

types in the hybrid individual is due to the relatively recent creation of the hybrid, where concerted evolution has not had enough time to fully homogenize the paralogous rDNA repeats. In the data presented here, the different ETS types were not taxon specific, as one would expect in a “pure” taxon, where concerted evolution has homogenized all the rDNA repeats. The ETS types were distributed throughout the majority of the *Cercocarpus* taxa sampled. Although hybridization is known to occur within *Cercocarpus* (Rieseberg 1989, 1994), to explain the distribution of ETS types throughout the *Cercocarpus* taxa using recent hybridization events, we would have to postulate a high level of long-distance hybridization. For example, *C. macrophyllum* (collection 26 from Durango, Mexico) is sister to *C. ledifolius* var. *intermontanus* (collection 45C from Utah) within the ETS type 1 clade, yet *C. macrophyllum* (collection 26) is sister to *C. betuloides* var. *betuloides* (collection 20 from California). The distances in this example exceed 1600 km. Gene flow in plants can occur through the movement of seeds or pollen. *Cercocarpus* displays both seed and pollen characteristics that favor dispersal. *Cercocarpus* achenes have a persistent, plume-like style that is thought to promote dispersal (Russel 1997). In addition, *Cercocarpus* has a floral morphology (apetalous and exerted stamens) and pollen morphology (small with smooth exine) that suggests wind pollination, although no formal studies of the pollination biology of *Cercocarpus* have been completed to date. Yet, when studying the patterns of seed dispersal in *C. ledifolius*, Russel

and Schupp (1998) found that both seed density and seed dispersal were highly influenced by microhabitat conditions, with most if not all the seeds being trapped in the leaf litter near the location of dispersal. Also, although the small, light pollen grains in *Cercocarpus* do disperse readily, the probability of a pollen grain landing on a receptive stigmatic surface decreases sharply as both the density of individuals goes down and distance increases, both of which occur between *Cercocarpus* populations. Therefore, we do not believe that long distance gene flow explains the distribution of ETS types in *Cercocarpus*.

The results we have obtained are also inconsistent with allopolyploidization as the process by which we have multiple, paralogous rDNA repeats. There have been eight chromosome counts spanning the four *Cercocarpus* taxa *C. betuloides* var. *betuloides*, *C. ledifolius* var. *ledifolius*, *C. ledifolius* var. *intermontanus*, and *C. montanus* var. *montanus*, and all have been constant at $X=9$ (Morley 1949, McArthur and Sanderson 1985). The chromosome counts for species in other genera closely related to *Cercocarpus* are also constant at $X=9$. There has been no evidence for recent polyploidization events within *Cercocarpus* or the genera allied with *Cercocarpus*.

Deep Coalescence of multiple ETS types in *Cercocarpus*

Using the alignable regions present in each ETS type, we showed that all of the ETS types found in *Cercocarpus* form a clade with high bootstrap support when compared to the outgroup *Purshia*. The monophyletic nature of the ETS

types in *Cercocarpus* could change with broader sampling in the genus *Purshia* and other genera closely allied with *Cercocarpus* in the Rosaceae. The ETS type 1 sequences are monophyletic, rooted at (Figure 2.5, 2.6A, 2.6B) or near the base of the gene trees (Figure 2.4), suggesting an early origin within the genus. ETS types 2, 3 and 4 also show a deep coalescence and are found sister to ETS type 1 clade (Figure 2.5, 2.6A). ETS type 2 and ETS type 4 sequences occur polyphyletic in both the MP and ML reconstructions in both the complete and reduced dataset, with ETS type 3 being monophyletic and nested within the phylogenetic grade. We believe that these results cannot be explained by simply inter-specific variation. In other studies of rDNA diversity within a species using internal transcribed spacer (ITS) sequences, the divergence between two different types did not exceeded 5% (Baldwin et al. 1995, Vargas et al. 1998, Steane et al. 1999). Yet we found the different ETS types within taxa and from single individuals differed by both length and sequence content due to the presence of long, unalignable regions. Multiple inter-specific ETS sequences from *Argyranthemum* and *Helianthus* coalesced within a species, leading to the conclusion that concerted evolution was working, but not as rapid as in the ITS or coding regions of the rDNA repeat (Linder et al. 2000). On occasions when multiple rDNA types have been discovered, which do not coalesce within a species and are not associated with hybridization or allopolyploidization, one of the rDNA types has been shown to feature pseudogene qualities (Buckler et al.

1997, Mayol and Rossello 2001). Although we cannot rule out that some of the different ETS types are pseudogenes, we have presented evidence above for why we believe the ETS types found in *Cercocarpus* are functional. If the different ETS types are functional, which is likely, it is clear that the different ETS repeats are not being homogenized due to concerted evolution, even when they are present in the same individual.

The unique nature of a deep coalescence of ETS types implies there is a lack of concerted evolution in the ETS region in *Cercocarpus*. The phenomenon of concerted evolution has been documented, yet the mechanism by which it proceeds is still not truly known. If concerted evolution indeed proceeds by unequal crossing over, then large mutations that interfere with successful pairing between strands may break down concerted evolution. It may be that the unalignable regions we found in the ETS types of *Cercocarpus* are examples of this.

Origin of multiple ETS types in *Cercocarpus*

Using the phylogenies we created for the ETS types and the structure and location of different alignable and unalignable regions found in each ETS type, we developed a hypothesis for the steps that might have occurred to create the current ETS diversity (Figure 2.7). To create an ancestral ETS sequence, we must account for the presence of the 3' alignable, the ETS type 2 unalignable, and the α

and β alignable regions in both the extant *Purshia* ETS type and *Cercocarpus* ETS types 1 and 2. Although other evolutionary scenarios with the same number of steps exist, the ancestral ETS sequence must contain these three regions. We propose, as the first step in this scenario, the insertion of the *Purshia* unalignable region into the *Purshia* ETS type and the insertion of the 5' alignable region into an ancestral *Cercocarpus* ETS. Whether or not the type 1 unalignable region was present in the putative ancestral ETS or was inserted along with the 5' alignable region has the same number of steps in this scenario. Based on the gene tree topology and the shared features of ETS type 1 and ETS type 2 (mainly the position of the α and β regions) we propose that ETS type 1 and type 2 were each created by the deletion of different regions from this proposed ancestral *Cercocarpus* ETS type. We do not have any evidence from the gene tree to make inferences about the sequence of the type 1 and type 2 creations. In the MP and ML analyses using the reduced dataset including the α and β regions, ETS type 1 is most basal, yet when we excluded the α and β regions, ETS type 1 and type 2 are unresolved at the base of the tree. The ETS type 3 clade is found nested within the ETS type 2/type 4 grade, suggesting an origin from ETS type 2. Based on the sequence and structural similarity, as well as their placement in the gene tree, we believe ETS type 4 is a derivative of ETS type 2 by way of a deletion event which excises the α and β regions and a portion of the ETS type 2 unalignable region. According to the gene tree, it appears this deletion event has

occurred 3 times. The ETS type 4 sequences take a more basal position to all the ETS type 2 sequences in the MP tree (Figure 2.4). Although this is contradictory to the arrangement one would expect based on our phylogenies (Figure 2.4 and 2.5), our other analyses (Figure 2.6) show at least one individual of ETS type 2 (*C. fothergilloides* var. *mojadensis* 18) is found at the base of the tree. This arrangement is most likely an artifact of the dataset in the MP analysis. The ETS type 2 sequences within the aligned dataset contain more information due to the inclusion of the α and β regions, which the ETS type 4 sequences do not contain. The lack of the α and β regions in ETS type 4 sequences forces them to a more basal position due to the absence of informative variation contained in the α and β regions. Also, there would be a low probability of a ETS type 2 arising from a type 4 multiple times to include all of the alignable regions in common with ETS type 1 and the outgroup *Purshia*.

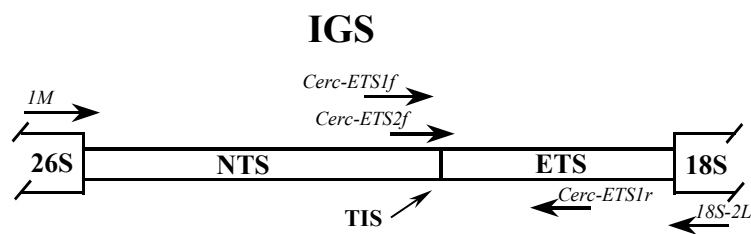


Figure 2.1. A representation of the intergenic spacer (IGS) between the 18S and 26S ribosomal genes in the nuclear genome. The genes 26S and 18S are shown by the large boxes. Transcription of the rDNA repeat begins at the transcription initiation site (TIS). The general location and direction of the main primers used in this study are shown by the arrows and the italicized text. The size of the IGS in *Cercocarpus* is approximately 2.5 kb. The size of the ETS in *Cercocarpus* ranges from 1000 bp to 500 bp.

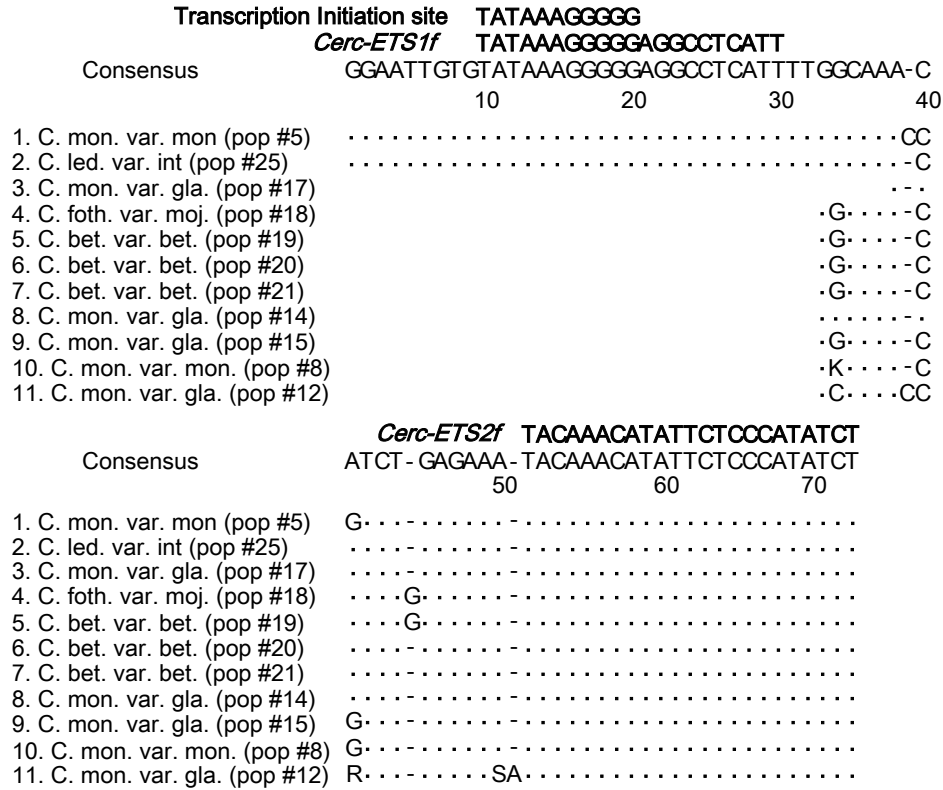


Figure 2.2. Alignment of ETS sequences in the region of the TIS and the placement of the *Cerc-ETS1f* and *Cerc-ETS2f* primers. The consensus sequence is shown on top. The dots refer to the same base as found in the consensus sequence, the dashes refer to gaps. The identity of each sequence is represented by an abbreviation. The abbreviation begins with C, referring to *Cercocarpus*. The three letters after the first period are the first three letters of the species name. var. refers to variety. The last three letters are an abbreviation of the first three letters of the variety name. The number in parentheses refers to the collection number. C. mon. var. mon (5) and C. int (25) were generated using the *Cerc-ETS1r* primer. Also aligned are the 9 samples successfully amplified and sequenced using the *Cerc-ETS1f* and *18S-2L* primers.

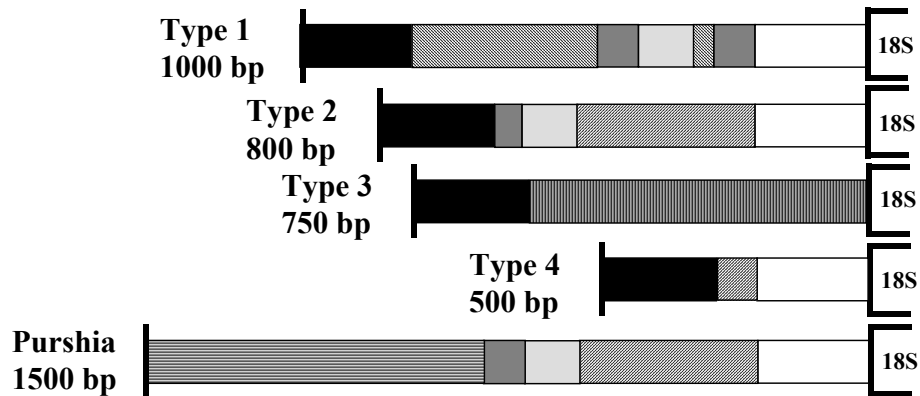
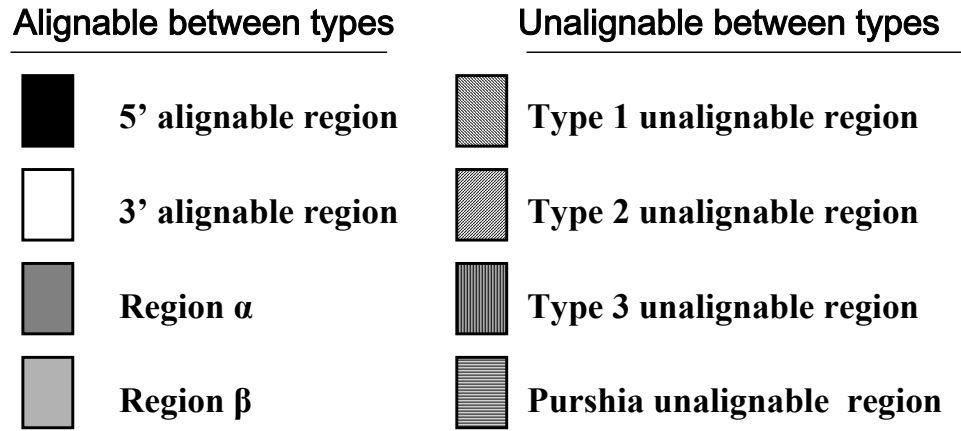


Figure 2.3. Representation of all four ETS types found in *Cercocarpus* and the ETS found in *Purshia*. Figure is not to scale. Representations of the ETS region begin at the transcription initiation site (TIS), shown by the vertical black bar on the left, and continue through to the 18S gene. Alignable and unalignable regions are shown by different hatching patterns, the identities are defined in the key.

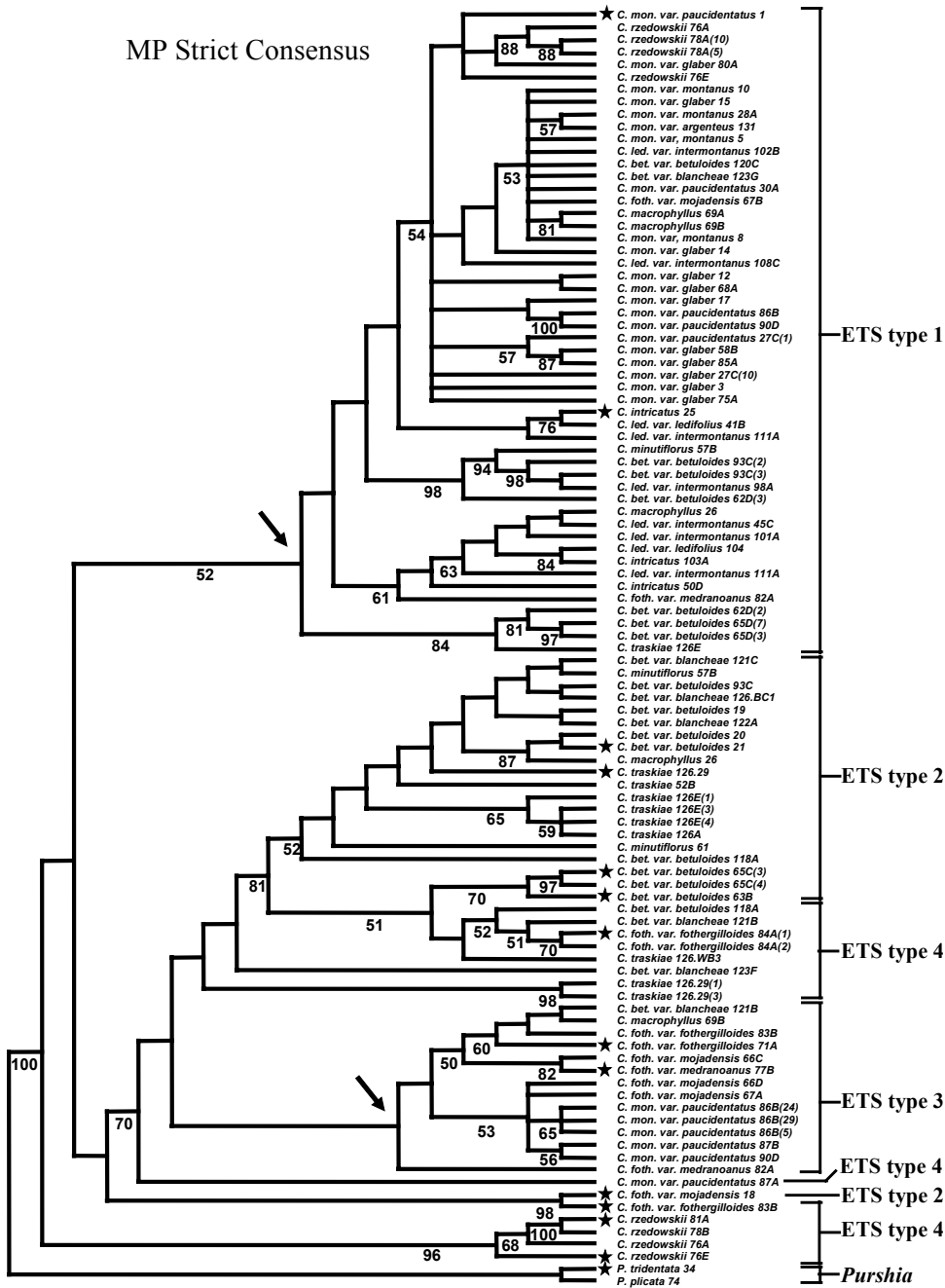


Figure 2.4. A maximum parsimony (MP) strict consensus tree including all ETS types of over 10,000 most parsimonious trees of length 530. Numbers below the branches indicate bootstrap support. The identity of the individual that produced each ETS sequence is abbreviated and shown on the right. The abbreviation begins with either C, referring to *Cercocarpus*, or P referring to *Purshia*. The three letters after the first period are the first three letters of the species name. var. refers to variety. The numbers in parentheses refer to the collection numbers. The ETS type designation of each sequence is shown on the right (Figure 2.3). The stars located to the left of the abbreviated name of the individual show the 14 sequences used in further analysis (Figure 2.6). The large arrows show the monophyletic nature of ETS type 1 and 3.

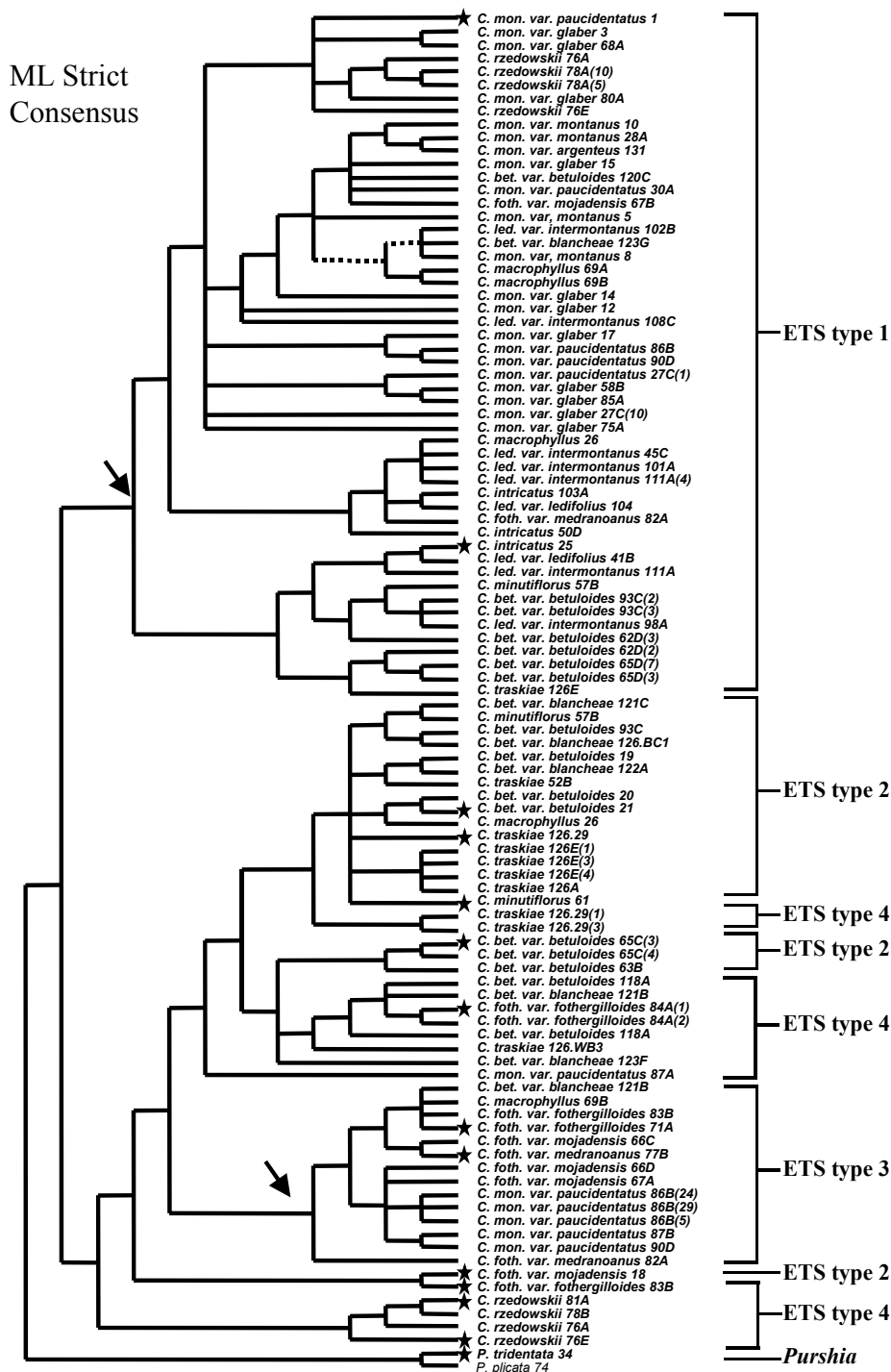
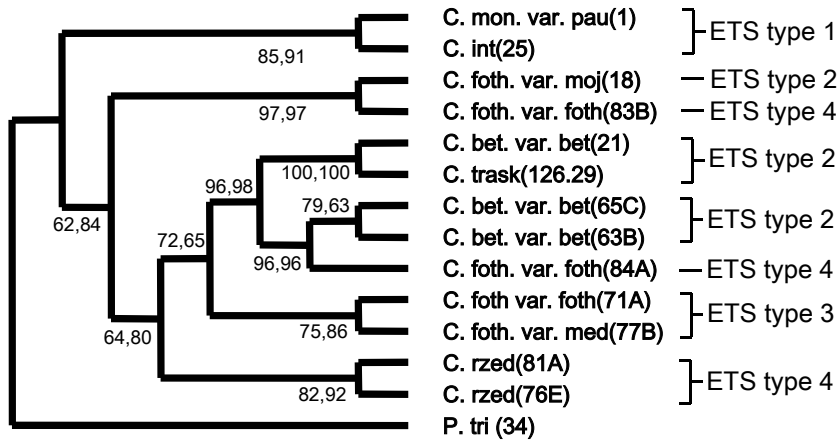


Figure 2.5. A maximum likelihood (ML) cladogram for all ETS types with a score of $-\ln 3851.04666$ produced from an aligned dataset of 512 bp of ETS sequence. This is one of five most likely trees. The dotted lines represent the branches that collapse in the strict consensus. The identity of the source of the ETS sequence and the ETS type designations are as in Figure 2.4. The small stars refer to the 14 ETS sequences used in further analyses. The large arrows highlight the monophyletic nature of ETS type 1 and 3.

A. With α and β



B. Without α and β

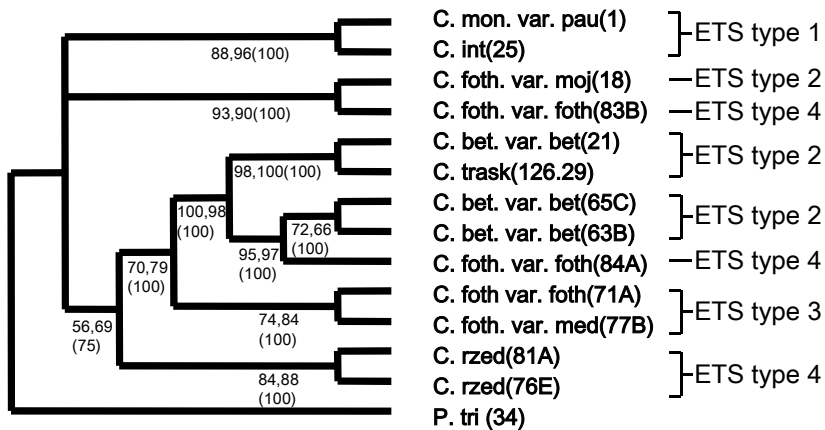


Figure 2.6. Cladograms representing both MP and ML analyses for the two reduced ETS datasets. **A.** A cladogram representing both MP and ML analyses using the 5' alignable, 3' alignable, and α and β regions. The two analyses resulted in the same topology. The MP tree had a length 182. The ML had a likelihood score of $-\ln = 1616.56253$. The numbers below the branches indicate bootstrap support. The numbers before the comma are bootstrap percentages from the MP analysis, the number after the comma bootstrap percentages from the ML analysis. The identities of the sequences are abbreviated as in Figure 2.2. The ETS type is designated as in Figure 2.4 and 2.5. **B.** A strict consensus cladogram representing both MP and ML analyses using just the 5' alignable and 3' alignable regions. The MP analyses produced 4 most parsimonious trees with a length of 141. The ML analysis resulted in 2 most likely trees with a score of $-\ln = 1246.24172$. Numbers below the branches are measures of bootstrap support. The number before the comma is the bootstrap values from the MP analysis; the number after the comma is the bootstrap value from the ML analysis. The numbers in the parentheses are the majority rule consensus values from the MP analysis.

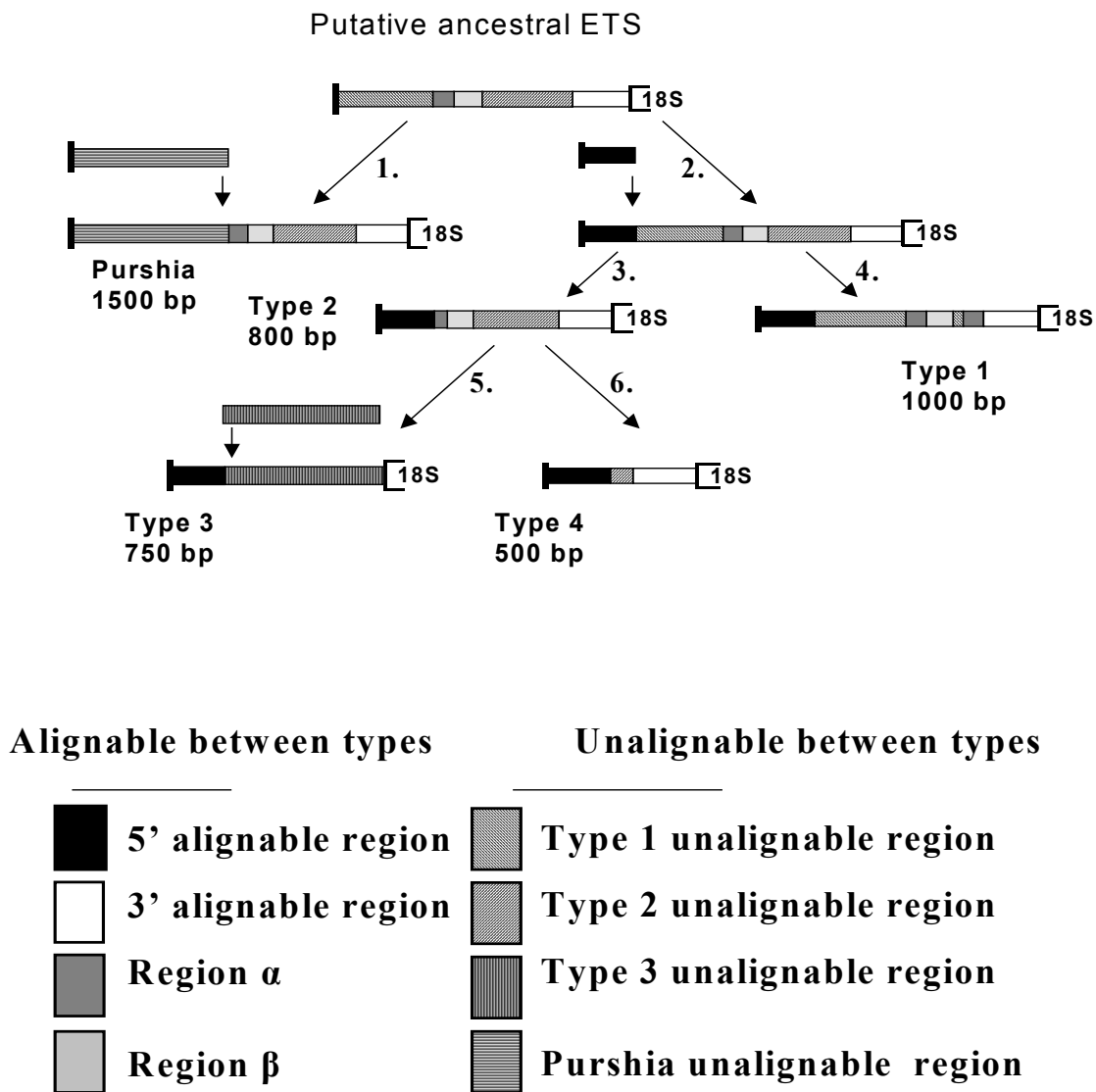


Figure 2.7. A proposed series of steps for the evolution of multiple ETS types in *Cercocarpus* and *Purshia*. Alignable and unalignable regions are illustrated as in Figure 2.3, the identities are defined in the key. 1) From the proposed ancestral sequence, the *Purshia* unalignable region is inserted. 2) From the proposed ancestral sequence, the 5' alignable region is inserted. 3) ETS type 2 is created by the deletion of the type 1 unalignable region. 4) ETS type 1 is created by the deletion of the type 2 unalignable region. 5) ETS type 3 is created by the insertion of the type 3 unalignable region. 6) ETS type 4 is created by the deletion of the central part of the ETS type 2 unalignable region.

Chapter 3: A nuclear phylogeny for *Cercocarpus* H.B.K. (Rosaceae) using AFLP and ETS sequence data

INTRODUCTION

Cercocarpus H.B.K. (Rosaceae) is a small genus made up of shrubs and trees found in chaparral and arid montane regions of western North America. Currently, *Cercocarpus* does not have a consensus classification or phylogeny. Since it was first described, *Cercocarpus* has undergone three revisions (Schneider 1905b, Rydberg 1913, Martin 1950), each varying widely in the ranks assigned to taxa and the number of species. The first revision (Schneider 1905b) described 10 species with 12 varieties. Rydberg (1913), in the second revision, described 21 species. Rydberg never recognized sub-specific taxa. The last revision by Martin (1950) described only six species with 11 varieties. Confusion about the number and circumscription of species within *Cercocarpus* results from the absence of clear morphological delimitations between taxa and, possibly, the lack of molecular phylogenetic treatments. Historically, the taxonomy in *Cercocarpus* has used leaf characteristics as the major source of characters for definition. Variations in leaf morphology show a continuum within and between taxa making it difficult to define boundaries among species (Brayton and Mooney 1966, Mortenson 1973), and making a phylogeny based on these characters misleading due to severe homoplasy (see Figure 1.4). It is clear that new characters, not based on gross morphology, are needed to resolve evolutionary

relationships among species and the diversity within species, which have otherwise remained elusive.

The predominant hypothesis (Schneider 1905b, Brayton and Mooney 1966) for the origin of intergradations in leaf characters between putative distinct taxa of *Cercocarpus* has been hybridization and introgression. Individuals displaying intermediate leaf forms are thought to be F1 hybrids, and the intergradations of leaf form between the hybrids and each parental taxon arises from varying amounts of introgression of leaf traits into both parental taxa at the zone of contact. A second hypothesis, mentioned briefly by Brayton and Mooney (1966) describes leaf variation in *Cercocarpus* populations as a plastic response to varying habitats, or a mixture of different ecotypes, where similar genotypes have been selected to have different leaf shapes, based on microhabitat selection.

Molecular phylogenetic treatments may be useful to test these hypotheses for the origin of leaf-based variation. If leaf variation is indeed due to hybridization, we might expect the two “parental taxa“ to unite in two, distinct monophyletic groups. The placement of the individuals with intermediate leaf morphology may differ, uniting with either “parental taxon” depending on whether the marker was inherited uni-parentally (chloroplast markers) or if concerted evolution has converted all copies of the marker in the putative hybrid to one “parental taxon”. Individuals with intermediate morphology may also be

found in an unresolved, basal position if there has been recombination between the two parental marker types in the hybrid individuals.

If leaf variation in populations of *Cercocarpus* is a product of plastic characters responding to different conditions or ecotypal variation, we might expect different tree topologies than those presented above based on hybridization. In this case, we would expect a single lineage for all leaf shapes with no clear clades representing the two “parental taxa.” We might expect instead to see clades corresponding to geographic/ecological position and not leaf shape.

A phylogeny for *Cercocarpus* can also help test three specific hypotheses about the evolution and taxonomic affinities of particular *Cercocarpus* taxa. These hypotheses are listed below. 1) Stutz (1974) claimed that *Cercocarpus ledifolius* is the ancestral species from which both *C. montanus* and *C. intricatus* originated by peripheral isolation. 2) Lis (1992) claimed that there are similarities between *Cercocarpus traskiae*, found only on Santa Catalina Island off the coast of southern California and *C. fothergilloides* in Mexico based on leaf architectural characters. This is a surprising hypothetical relationship due to the historical view that *Cercocarpus traskiae* is closely related to *C. betuloides* of California (Martin 1950, Rieseberg 1989, 1994). 3) We can also test if *Cercocarpus rzedowskii* and *C. mexicanus*, two recently described species, are related to *C. ledifolius* and *C. montanus*, respectively (Henrickson 1987).

In this study, we developed the first phylogenetic hypotheses for *Cercocarpus* using nuclear molecular characters. We constructed phylogenies for *Cercocarpus* using amplified fragment length polymorphisms (AFLP) markers (Vos et al. 1995) and sequence data from the external transcribed spacer (ETS) of the nuclear ribosomal DNA (nrDNA). AFLPs (Vos et al. 1995) have been used extensively to examine relationships among closely related groups of plants, including cultivars, varieties, and species (Mueller and Wolfenbarger 1999, Zhang et al. 2001, Koopman et al. 2001, Hedren et al. 2001). In this study, we used AFLPs to construct phylogenies for *Cercocarpus*, concentrating on the *C. montanus* and *C. ledifolius* species complexes, to elucidate species and variety delimitations and relationships among *Cercocarpus* taxa. Initially, we explored the internal transcribed spacer (ITS) region of the nrDNA as a potential marker for phylogenetic reconstruction within *Cercocarpus*. The ITS region has been used with success for phylogenetic reconstruction at the generic and specific level in other plant groups (Baldwin et al. 1995). After finding the ITS region lacking in informative variation, we developed the ETS region as a DNA sequence marker for phylogenetic reconstruction. The details of the ETS's development for phylogenetic reconstruction and its molecular evolution are reported in Chapter 2. The ETS has been shown to be phylogenetically informative in very closely related groups, including island taxa (Linder et al. 2000, Baldwin and Markos

1998). Here, we present the results of the phylogenetic reconstruction using the ETS and their implications for the relationships within *Cercocarpus*.

MATERIALS AND METHODS

Collection and classification of Plant Material

We followed the proposed taxonomy of Lis (1992) with the exception of the California taxa - for which we used the taxonomy in the Jepson manual (Lis 1993) - and the Mexican taxa, for which we used an unpublished treatment by J. Henrickson (personal communication). We used 153 accessions representing 16 taxa of *Cercocarpus* (Table 3.1). Leaves from five herbarium sheets were included in this total. The remaining 148 accessions were field-collected leaf material, dried using silica gel for transport to the lab. The morphological characters used to distinguish species and sub-specific categories are listed in Table 3.2. Members of the genus *Purshia* were used as the outgroup for this study. *Purshia* has been shown to be sister to *Cercocarpus* in recent molecular phylogenies based on both chloroplast (Morgan et al. 1994) and nuclear genes (Rodgers et al. 1998).

Table 3.1: *Cercocarpus* taxa included in this study, including codes used in this paper, voucher numbers, collectors, and geographic position. BVH = Brian Vanden Heuvel, JH = James Henrickson. * denote individuals sampled in the ETS dataset. Taxa have been grouped by species complex according to Lis (1992).

taxon	code	location	Collector and Voucher
<i>montanus</i> complex			
<i>C. montanus</i> var. <i>montanus</i>	*5	Castle Rock, Douglas Co., CO USA	BVH 5
	*8	Hartville, Platte Co., WY USA	BVH 8
	*10	Dewey, Custer Co., SD USA	BVH 10
	*28A	Bailey Canyon, Otero Co., NM USA	BVH 28
<i>C. montanus</i> var. <i>glaber</i>	*3	Jame, Coahuila, MX	JH 20439B
	*12	Chisos Mts., Brewster Co., TX USA	BVH and JH 20662
	13	Chisos Mts., Brewster Co., TX USA	BVH and JH 20663
	*14	Chisos Mts., Brewster Co., TX USA	BVH and JH 20664
	*16	Chisos Mts., Brewster Co., TX USA	BVH and JH 20665
	*17	Sierra de la Modera, Coahuila, MX	JH 20637
	*58B	Leaky, Real Co., TX USA	JH 22275
	58D	Leaky, Real Co., TX USA	JH 22275
S.7	Guadalupe Mts., Culberson Co., TX USA	Correll 18482	
<i>C. montanus</i> var. <i>paucidentatus</i>	*1	La Perdida, Tamaulipas, MX	JH 19218
	27A	Guadalupe Mts., Eddy Co., NM USA	BVH 27
	27B	Guadalupe Mts., Eddy Co., NM USA	BVH 27
	*27C	Guadalupe Mts., Eddy Co., NM USA	BVH 27
	*30A	Pinos Altos, Grant Co., NM USA	BVH 30A
	30D	Pinos Altos, Grant Co., NM USA	BVH 30D
	30E	Pinos Altos, Grant Co., NM USA	BVH 30E
	30F	Pinos Altos, Grant Co., NM USA	BVH 30F
	31	Gila Mts., Grant Co., NM USA	BVH 31
	32B	Gila Mts., Grant Co., NM USA	BVH 32
	68A	Jame, Coahuila, MX	BVH and JH 22414
	68B	Jame, Coahuila, MX	BVH and JH 22414
	68C	Jame, Coahuila, MX	BVH and JH 22414
	70A	Jame, Coahuila, MX	BVH and JH 22416
	70B	Jame, Coahuila, MX	BVH and JH 22417
	72A	Cerro Potosi, Coahuila, MX	BVH and JH 22424
	72B	Cerro Potosi, Coahuila, MX	BVH and JH 22424
	*75A	Inturbide, Coahuila, MX	BVH and JH 22429
	*80A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22457
	80C	Valle Hermosa, Tamaulipas, MX	BVH and JH 22457
	80D	Valle Hermosa, Tamaulipas, MX	BVH and JH 22457
	*85A	Sierra de la Encantada, Coahuila, MX	BVH and JH 22587
	85B	Sierra de la Encantada, Coahuila, MX	BVH and JH 22587
	85D	Sierra de la Encantada, Coahuila, MX	BVH and JH 22587
	85E	Sierra de la Encantada, Coahuila, MX	BVH and JH 22587
	86A	Tunnel Springs, Bernalillo Co., NM USA	BVH 86
	*86B	Tunnel Springs, Bernalillo Co., NM USA	BVH 86
	87A	Kingston, Sierra Co., NM USA	BVH 87
	89A	Mogollon Mts., Catron Co., NM USA	BVH 89
	89B	Mogollon Mts., Catron Co., NM USA	BVH 89
	90A	San Francisco Mts., Catron Co., NM USA	BVH 90
	90B	San Francisco Mts., Catron Co., NM USA	BVH 90
*90C	San Francisco Mts., Catron Co., NM USA	BVH 90	
92A	Rose Peak, Greenlee Co., AZ USA	BVH 92	

	92B	Rose Peak, Greenlee Co., AZ USA	BVH 92
	95A	Tonto State Park, Gila Co., AZ USA	BVH 95
	*96A	Strawberry, Gila Co., AZ USA	BVH 96
	96B	Strawberry, Gila Co., AZ USA	BVH 96
	HEN1	San Antonio Pena Nevada, Nuevo Leon, MX	JH HEN1
	S.3	Guadalupe Mts., Culberson Co., TX USA	Correll 24804
<i>C. montanus</i> var. <i>argenteus</i>	*131	Guadalupe Mts., Culberson Co., TX USA	McVaugh 7424
<i>betuloides</i> complex			
<i>C. betuloides</i> var. <i>betuloides</i>	19	La Conada, Los Angeles Co., CA USA	JH 20731
	20	La Conada Los Angeles Co., CA USA	JH 20732
	*62D	Porterville, Tulare Co., CA USA	BVH 127a
	63B	Lake Hume, Tulare Co., CA USA	BVH 127b
	65C	Camp Nelson, Tulare Co., CA USA	BVH 127c
	*65D	Camp Nelson, Tulare Co., CA USA	BVH 127c
	*93C	Miami, Gila Co., AZ USA	BVH 93
	112	Yreka, Siskiyou Co., CA USA	BVH 112
	113A	Callahan, Siskiyou Co., CA USA	BVH 113
	116	Weaverville, Trinity Co., CA USA	BVH 116
	118A	Tularcitos Ridge, Monterey Co., CA USA	BVH 118
	*120C	American Canyon, San Luis Obispo Co., CA USA	BVH 120
	127A	Camp Wishone, Tulare Co., CA USA	BVH 127
	*128A	Oakhurst, Madera Co., CA USA	BVH 128
<i>C. betuloides</i> var. <i>blancheae</i>	130A	Hathaway Pines, Calaveras Co., CA USA	BVH 130
	52B	Rancho Santa Ana Botanic Garden, Los Angeles Co., CA USA	JH 22312
	121A	Parisima Hills, Santa Barbara Co., CA USA	BVH 121
	122A	Solvang, Santa Barbara Co., CA USA	BVH 122
	123F	Santa Monica Mts., Los Angeles Co., CA USA	BVH 123
	*123G	Santa Monica Mts., Los Angeles Co., CA USA	BVH 123
	125A	Santa Catalina Island, Los Angeles Co., CA USA	BVH 125
<i>C. minutiflorus</i>	*57B	Del Mar, San Diego Co., CA USA	JH 22311
	61	Fallbrook, San Diego Co., CA USA	BVH 127a
<i>fothergilloides</i> complex			
<i>C. traskiae</i>	126A	Santa Catalina Island, Los Angeles Co., CA USA	Thorne 36737
	*126E	Santa Catalina Island, Los Angeles Co., CA USA	Thorne 36737
<i>C. fothergilloides</i> var. <i>fothergilloides</i>	66C	Los Lirios, Coahuila, MX	BVH and JH 22411
	66D	Los Lirios, Coahuila, MX	BVH and JH 22412
	67A	Los Lirios, Coahuila, MX	BVH and JH 22413
	*67B	Los Lirios, Coahuila, MX	BVH and JH 22413
	71A	Cerro Potosi, Coahuila, MX	BVH and JH 22423
	83A	Tula, Tamaulipas, MX	BVH and JH 22464
	84G	San Roberto Junction, Nuevo Leon, MX	BVH and JH 22482
<i>C. fothergilloides</i> var. <i>medranoanus</i>	77B	Valle Hermosa, Tamaulipas, MX	BVH and JH 22444
	*82A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22456b
<i>C. fothergilloides</i> var. <i>mojadensis</i>	18	Sierra Encantada, Coahuila, MX	JH 20639
<i>C. rzedowskii</i>	*76A	Miquihuana, Tamaulipas, MX	BVH and JH 22434
	*76E	Miquihuana, Tamaulipas, MX	BVH and JH 22434
	*78A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22446
	81A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22460

<i>C. mexicanus</i>	134A	Coahuila, MX	JH 19295
<i>C. macrophyllus</i>	*26	Durango, MX	Crozier n/a
	*69A	Sierra de la Vega, Coahuila, MX	BVH and JH 22415
	*69B	Sierra de la Vega, Coahuila, MX	BVH and JH 22415
	79	Valle Hermosa, Tamaulipas, MX	BVH and JH 22458
 <i>ledifolius</i> complex			
<i>C. ledifolius</i> var. <i>intermontanus</i>	37A	Casto Canyon, Garfield Co., UT USA	BVH 37
	37B	Casto Canyon, Garfield Co., UT USA	BVH 37
	37C	Casto Canyon, Garfield Co., UT USA	BVH 37
	39	Gooseberry, Sevier Co., UT USA	BVH 39
	45A	Mantua, Box Elder Co., UT USA	BVH 45
	45B	Mantua, Box Elder Co., UT USA	BVH 45
	*45C	Mantua, Box Elder Co., UT USA	BVH 45
	46A	Preston, Franklin Co., ID USA	BVH 46
	46B	Preston, Franklin Co., ID USA	BVH 46
	46C	Preston, Franklin Co., ID USA	BVH 46
	*98A	Verni, Sierra Co., CA USA	BVH 98
	98B	Verni, Sierra Co., CA USA	BVH 98
	98C	Verni, Sierra Co., CA USA	BVH 98
	99A	Santa Rose Mts., Humboldt Co., NV USA	BVH 99
	99B	Santa Rose Mts., Humboldt Co., NV USA	BVH 99
	99C	Santa Rose Mts., Humboldt Co., NV USA	BVH 99
	*101A	Toiyabe Range, Lander Co., NV USA	BVH 101
	102A	Ruby Mts., Elko Co., NV USA	BVH 102
	*102B	Ruby Mts., Elko Co., NV USA	BVH 102
	102C	Ruby Mts., Elko Co., NV USA	BVH 102
	105	Ely, White Pine Co., NV USA	BVH 105
	108A	Spring Mts., Clark Co., NV USA	BVH 108
	108B	Spring Mts., Clark Co., NV USA	BVH 108
	*108C	Spring Mts., Clark Co., NV USA	BVH 108
	110	Whits Mts., Inyo Co., CA USA	BVH 110
	*111A	West Walker river, Mono Co., CA USA	BVH 111
	111B	West Walker river, Mono Co., CA USA	BVH 111
	111C	West Walker river, Mono Co., CA USA	BVH 111
<i>C. ledifolius</i> var. <i>ledifolius</i>	41A	Emery, Emery Co., UT USA	BVH 41
	41B	Emery, Emery Co., UT USA	BVH 41
	*41C	Emery, Emery Co., UT USA	BVH 41
	*104	Ely, White Pine Co., NV USA	BVH 104
	106A	Cannons pass, White Pine Co., NV USA	BVH 106
	106B	Cannons pass, White Pine Co., NV USA	BVH 106
	106C	Cannons pass, White Pine Co., NV USA	BVH 106
	114	Noyes valley, Siskiyou Co., CA USA	BVH 114
<i>C. intricatus</i>	*25	Escalante, Garfield Co., UT USA	BVH 25
	43	Emery, Emery Co., UT USA	BVH 43
	48A	Flaming Gorge, Dagget Co., UT USA	BVH 48
	48B	Flaming Gorge, Dagget Co., UT USA	BVH 48
	48C	Flaming Gorge, Dagget Co., UT USA	BVH 48
	48D	Flaming Gorge, Dagget Co., UT USA	BVH 48
	50A	Vernal, Uintah Co., UT USA	BVH 50
	50B	Vernal, Uintah Co., UT USA	BVH 50
	50C	Vernal, Uintah Co., UT USA	BVH 50
	*50D	Vernal, Uintah Co., UT USA	BVH 50
	*103A	Ely, White Pine Co., NV USA	BVH 103
	103B	Ely, White Pine Co., NV USA	BVH 103
	103C	Ely, White Pine Co., NV USA	BVH 103

<i>Purshia tridentata</i>	*36B	Casto Canyon, Garfield Co., UT USA	BVH 36
	100B	Santa Rose Mts., Humboldt Co., NV USA	BVH 100
<i>Purshia stansburiana</i>	54A	Coliseum Canyon, San Bernardino Co., CA USA	JH 22307-5
<i>Purshia glandulosa</i>	55A	Cima Rd., San Bernardino Co., CA USA	JH 22303-5
<i>Purshia plicata</i>	*74D	Interbide, Coahuila, MX	Nesom 6198

Table 3.2. Eight morphological characters used to define taxa in *Cercocarpus*. The characters listed for *C. montanus* var. *montanus*, *C. montanus* var. *argenteus*, *C. montanus* var. *paucidentatus*, and *C. montanus* var. *glaber* are from Martin (1950). The characters of *C. betuloides* var. *betuloides*, *C. betuloides* var. *blancheae*, *C. minutiflorus*, and *C. traskiae* are from Lis (1993). The characters defining *C. rzedowskii* and *C. mexicanus* are from Henrickson (1987). *C. fothergilloides* var. *fothergilloides*, *C. fothergilloides* var. *mojadensis*, *C. fothergilloides* var. *medranoanus*, and *C. macrophyllus* are from an unpublished treatment by Henrickson (personal communication).

taxon	leaf size (cm)	leaf thickness	leaf shape	abaxial leaf pubescence	# of lateral veins	leaf margin	length of persistent style (cm)	geography
<i>C. montanus</i> var. <i>montanus</i>	1.5 - 3 long 0.4-2 wide	thin	obovate, rotund, flabelliform	short villous, appressed silky	3 - 6	coarse ovate to triangular teeth present on distal half of blade	5 - 6	Great Basin and Rocky Mountains
<i>C. montanus</i> var. <i>glaber</i>	1 - 3 long 0.6 - 2 wide	thin	obovate, elliptical, rotund	glabrate	4 - 6	short, broad triangular apiculate tipped teeth present on distal half of blade	5 - 9	Chihuahuan Desert
<i>C. montanus</i> var. <i>paucidentatus</i>	0.7 - 2 long 0.3 - 1 wide	thin	lanceolate, oblanceolate, narrow obovate	short villous, appressed silky	3 - 5	entire to 3 - 5 short teeth at apex	1.5 - 5	Chihuahuan Desert
<i>C. montanus</i> var. <i>argenteus</i>	2 - 4 long 0.9 - 1.7 wide	thin	oblanceolate, obovate, rotund	appressed silky to wholly	5 - 6	short, broad triangular apiculate tipped teeth present on distal half of blade	5 - 6	TX and NM
<i>C. betuloides</i> var. <i>betuloides</i>	1 - 4 long 0.6 - 2 wide	thin	obovate, rotund	sparsely silky	4 - 7	finely toothed to serrate	5 - 9	CA and AZ
<i>C. betuloides</i> var. <i>blancheae</i>	2.5 - 6 long 2 - 3.5 wide	coriaceous	widely elliptic, ovate	sparsely silky	6 - 10	finely toothed to serrate, margins revolute	3 - 6	CA
<i>C. minutiflorus</i>	1 - 2.5 long 0.7 - 1.5 wide	thin	ovate, rotund	glabrous	3 - 5	entire to short, broad triangular apiculate tipped teeth present on distal half of blade	3 - 7	CA and Baja California

<i>C. traskiae</i>	3 - 6 long 2 - 4 wide	coriaceous	rotund, broadly obovate	woolly	6 - 8	revolute with short, broad triangular apiculate tipped teeth present on distal half of blade	4.5 - 6	Santa Catalina Island, CA
<i>C. macrophyllus</i>	4 - 11 long 2.5 - 7.8 wide	coriaceous to thin	oblong-ovate, oblong-elliptical	glabrous	12 - 18	teeth ovate to acute	5 - 8.5	Mexico
<i>C. fothersilloides</i> var. <i>fothersilloides</i>	2.5 - 4.5 long 1.5 - 3 wide	coriaceous	ovate, broadly ovate	sparsely to moderately villous	5 - 8	serrate to crenate on distal two-thirds of blade	4 - 8.5	Mexico
<i>C. fothersilloides</i> var. <i>mojadensis</i>	1.6 - 2.8 long 1.2 - 2.2 wide	coriaceous	broadly ovate to orbicular	dense, white tomentose	4 - 6	crenate on distal two-thirds of blade	4 - 6.5	Mexico
<i>C. fothersilloides</i> var. <i>medranoanus</i>	1.5 - 2.5 long 1.1 - 1.6 wide	coriaceous	ovate to orbicular	uniform, tight coiled hairs	4 - 6	crenate on distal two-thirds of blade	3 - 4	Mexico
<i>C. rzedowskii</i>	0.4 - 1.5 long 0.4 - 0.6 wide	coriaceous	elliptical	thick, white tomentose	5 - 6	revolute, entire or with 2 - 4 spine tipped teeth on distal one-third	4.5 - 5	Mexico
<i>C. mexicanus</i>	1.7 - 3.1 long 1 - 1.5 wide	coriaceous	narrowly ovate	sparingly silky with large stomatal crypts	4 - 7	serrated on distal two-thirds	4.5 - 5	Mexico
<i>C. ledifolius</i> var. <i>intermontanus</i>	2.5 - 6 long 0.5 - 0.8	coriaceous	elliptical-lanceolate	glabrate	n/a	entire, revolute	4 - 10	Great Basin

	wide							
<i>C. ledifolius</i> var. <i>ledifolius</i>	1 - 2.5 long 0.1 - 0.4 wide	coriaceous	narrowly elliptical- lanceolate	densely- woolly	n/a	entire, revolute	3 - 6	Great Basin
<i>C. intricatus</i>	0.3 - 1 long 0.1 - 0.2 wide	coriaceous	linear	densely- woolly	n/a	entire, revolute to midrib	1 - 2	Great Basin

DNA Extraction

For all leaf samples, DNA was extracted using a modified CTAB procedure (Doyle and Doyle 1987). Genomic DNA extractions were cleaned using the Elu-quik DNA purification kit (Schleicher and Schuell), prior to further work.

AFLP methods

We followed the general protocol of Vos et al. (1995) to produce AFLP fragments. Briefly, sample DNA was restriction digested using the endonucleases EcoRI and MseI and ligated to the appropriate adapters. Two rounds of PCR amplification followed. The first preselective amplification used primers with a 1 base pair (bp) extension, and the second selective amplification used primers with 3 bp extensions. We used three different selective primer combinations having extensions -CCC/-AGC, -CCC/-ATA, and -CGG/-AGC, where the extensions refer to the EcoRI/MseI adapters, respectively. The -CCC EcoRI selective primers were labeled with the flouochrome FAM and the -CGG EcoRI selective primer was labeled with the flouochrome TAMRA (IDT industries).

The amplified fragments were separated and visualized on a MJ BaseStation (MJ Research), and sized using the size standard GS500 (ABI). Fragment scoring and lane tracking were done automatically within the software package Cartographer (MJ Research), and checked and adjusted by eye.

Fragments between 35 and 500 bp were scored. The three primer combinations were combined into one dataset containing 376 presence/absence characters (Appendix B and C).

The data were analyzed in two steps. First, analyses were performed on the complete dataset, containing all 153 accessions. The complete AFLP dataset was analyzed using both distance and parsimony criteria. The distance analysis consisted of NJ clustering and a heuristic search using minimum evolution (ME) distance estimates as implemented in PAUP* 4.09b (Swofford 1999). The ME method searches for tree topologies where the absolute difference between observed distances and path length distances are minimized. The parsimony analysis was performed in PAUP* 4.09b (Swofford 1999). A heuristic search strategy was employed with tree bisection/reconnection (TBR) branch swapping and applied the saving multiple parsimonious trees at each step (MULPARS) in effect. In order to explore a large amount of tree space without running out of memory trying to swap all possible trees to completion, 10,000 addition replicates were performed, using equally weighted characters and saving only 10 trees per replicate. A histogram of tree lengths for all 100,000 trees saved was obtained which allows the identification of multiple, equally parsimonious trees. The validity of using maximum parsimony as an optimality criterion in analyses using AFLP data has been challenged recently (see Koopman et al. 2001 for discussion). The dispute centers the assumption that equal-sized AFLP markers

are homologous and the homoplasies that may be introduced into parsimony analysis due to this assumption. Therefore, we have compared the MP and distance trees to look for incongruence and assessed the strength of support for particular branches with 100 bootstrap replicates using the same parameters as described above.

The second set of analyses used a presence/absence AFLP data subset that matched the same accessions used for the ETS analysis (see below). This reduced dataset contained 49 individuals (Table 3.1) and was used to compare directly to the ETS dataset. Parsimony analyses were performed on this smaller dataset as described above, except 1000 bootstrap replicates were performed for a more accurate indication of branch support.

ETS amplification and sequencing

The methods and details of the development for amplifying the ETS in *Cercocarpus* are found in the preceding chapter (Chapter 2). Briefly, following the methods outlined in Linder et al. (2000), we developed a forward primer located near the transcription initiation site (TIS) that reliably amplified the ETS in individuals of *Cercocarpus*. We discovered five different ETS paralogues (types) in our survey, four within *Cercocarpus* and one type in the outgroup *Purshia*. The four ETS types in *Cercocarpus* have deep coalescences within the genus and none of the ETS types was taxon specific. Only two of the four ETS types (type 1 and type 3) were monophyletic in analyses of all the ETS types, and

ETS type 1 had a deeper coalescence point than ETS type 3 in our analysis. Also, of the 275 successful ETS amplifications, ETS type 1 was found in 180 of these (Chapter 2). Due to the broad distribution, monophyletic nature, and deep coalescence of ETS type 1, we used ETS type 1 sequences for phylogeny reconstruction in *Cercocarpus*.

The total data set included 55 ETS sequences with an aligned length of 977 bp. We included all 53 ETS type 1 sequences, generated from 47 individuals representing 16 taxa of *Cercocarpus*, and two ETS sequences from *Purshia plicata* and *Purshia tridentata* served as outgroups for this analysis (Table 3.1). Six *Cercocarpus* individuals (populations 27C, 62D, 65D, 78A, 93C, and 111A) had two cloned sequences each in the data matrix. .

ETS type 1 products were amplified using the forward primer *Cerc-ETS1f*, located at the transcription initiation site (TIS), or the *Cerc-ETS2f* forward primer, located 14 bp downstream of *Cerc-ETS1f*, and the reverse primer *18S-2L* located in the 5' end of the 18S gene (Chapter 2). Fifty microliter PCRs were performed, as described in Chapter 2, using twenty nanograms of template DNA. The PCRs were overlaid with mineral oil and run on a MJ Research Thermal Cycler. Three microliters of the reaction products were run on a 1.5 % agarose gel in 0.5X TBE buffer. PCR products were visualized with ethidium bromide under UV light and purified using the QIAquick PCR purification kit (QIAGEN). DNA sequencing was performed on an ABI Prism 377 or an ABI Prism 3100 automated sequencer

using standard reagents. Sequences were proofread and then aligned using Clustal W in MegAlign (DNASTar, 1998), followed by adjustment by eye.

Phylogenetic analysis of the ETS

Maximum parsimony and maximum likelihood analyses were performed using the software package PAUP* version 4.0b8 (Swofford 1999). For our maximum parsimony analysis, we employed a heuristic search strategy using 10 random replicates, TBR branch-swapping and the MULTREES optimization. We set MAXTREES to 15,000. Support for branches was evaluated using bootstrap analysis (Felsenstein 1985) and random sequence addition for 100 replicates, using the same parameters.

For our maximum likelihood analysis, we used the HKY85 model with two substitution types, a transition-transversion ratio of 2, use of empirical base frequencies derived from the alignment, and no *a priori* site-specific variation. We used a heuristic search strategy using 10 random replicates and TBR branch-swapping.

Tests for incongruence

To assess the incongruence between the ETS and AFLP datasets, we conducted the "compare-2" test (Faith et al. 1991), the Templeton test (Templeton 1983), and the Kishino-Hasegawa test as implemented in PAUP* version 4.0b8 (Swofford 1999). To compare the AFLP dataset to the ETS dataset, we had to

account for the two cases of paraphyletic ETS type 1 clones from the individuals *C. ledifolius* var. *intermontanus* 111A (clone 4 and 5) and *C. betuloides* var. *betuloides* 62DA (clones 2 and 3) (see Results below). Therefore, we constructed 4 ETS datasets, which differed in the clones that represented the individuals 111A and 62D in all possible combinations (Table 3.3). The compare-2 test examines the number of steps required by a dataset on two different constraint trees. For this study, the first constraint tree was the MP bootstrap consensus tree containing only branches with greater than 50% support from the dataset being tested. The second constraint tree, the test constraint tree, was the rival bootstrap consensus tree using the other dataset.

Table 3.3: A description of the clones included in each of the ETS datasets tested against the cpDNA dataset. 111A refers to the individual *C. ledifolius* var. *intermontanus* collection 111A (Table 3.1). 62D refers to *C. betuloides* var. *betuloides* collection 62D (Table 3.1).

	111A clone 4	111A clone 5	62D clone 2	62D clone 3
ETS dataset 1	included	not included	included	not included
ETS dataset 2	included	not included	not included	included
ETS dataset 3	not included	included	included	not included
ETS dataset 4	not included	included	not included	included

The Templeton test compares the number of steps each individual character needs to fit on two rival tree topologies, and then uses a Wilcoxon sign ranks (WSR) test to ascertain significance. The Kishino-Hasegawa is a parametric test comparing the difference in tree lengths of the dataset on the two rival tree topologies to the variance in the tree length difference over many sites. Significance is determined by comparing the original length difference to a distribution of tree length differences. The distribution is created from examining all the sites where the tree length differs between the two constraint trees, subtracting a mean difference and squaring. If the dataset supports both topologies equally, we would expect tree length differences to be distributed normally with a mean of zero. The trees used in the Templeton and the Kishino-Hasegawa tests are the same as the constraint trees used with the compare-2 test.

Testing the effect of different combinations of individuals representing taxa on tree topology

We constructed MP trees based on reduced AFLP and ETS datasets to determine if reduced sampling within taxa would have significantly altered our conclusions about tree topology. For the AFLP data, we created three reduced datasets (A, B, and C), each including 6 individuals, one for each of the taxa (Table 3.4). For the MP analyses on the three reduced AFLP datasets, we conducted exhaustive searches in PAUP* version 4.0b8 based on the 376 presence/absence characters. We also measured branch support with a bootstrap analysis using 1000 replicates.

Table 3.4: The identity of the individuals used in the reduced AFLP datasets A, B, and C. The numbers in parentheses are the collection number. The collection location and voucher number can be found in Table 3.1.

Species	AFLP Dataset A	AFLP Dataset B	AFLP Dataset C
<i>C. montanus</i>	<i>C. montanus</i> var. <i>montanus</i> (5)	<i>C. montanus</i> var. <i>montanus</i> (5)	<i>C. montanus</i> var. <i>glaber</i> (17)
<i>C. betuloides</i>	<i>C. betuloides</i> var. <i>blancheae</i> (123G)	<i>C. betuloides</i> var. <i>betuloides</i> (65D)	<i>C. betuloides</i> var. <i>blancheae</i> (123G)
<i>C. ledifolius</i>	<i>C. ledifolius</i> var. <i>intermontanus</i> (102B)	<i>C. ledifolius</i> var. <i>intermontanus</i> (102B)	<i>C. ledifolius</i> var. <i>intermontanus</i> (102B)
<i>C. macrophyllus</i>	<i>C. macrophyllus</i> (26)	<i>C. macrophyllus</i> (26)	<i>C. macrophyllus</i> (26)
<i>C. fothergilloides</i>	<i>C. fothergilloides</i> var. <i>medranoanus</i> (82A)	<i>C. fothergilloides</i> var. <i>medranoanus</i> (82A)	<i>C. fothergilloides</i> var. <i>medranoanus</i> (82A)
<i>Purshia tridentata</i>	<i>P. tridentata</i> (34)	<i>P. tridentata</i> (34)	<i>P. tridentata</i> (34)

For the ETS data, we created two reduced datasets (D and E), each including 15 individuals, one for each of the taxa (Table 3.5). Five of the 15 individuals (individuals 131, 123G, 57B, 126E, and 34) were the same in each dataset because we had no other ETS sequences to represent that taxon. We conducted a MP heuristic search analysis on the two ETS reduced datasets based on 977 aligned bases. Bootstrap support was determined using 100 replicates.

Table 3.5: The identity of the individuals used in the reduced ETS datasets D and E. The numbers in the columns are the collection numbers. The collection location and voucher number can be found in Table 3.1.

taxon	ETS reduced dataset D	ETS reduced dataset E
<i>C. montanus</i> var. <i>montanus</i>	28A	28A
<i>C. montanus</i> var. <i>paucidentatus</i>	1	30A
<i>C. montanus</i> var. <i>glaber</i>	3	14
<i>C. montanus</i> var. <i>argenteus</i>	131	131
<i>C. betuloides</i> var. <i>betuloides</i>	93C	128C
<i>C. betuloides</i> var. <i>blancheae</i>	123G	123G
<i>C. minutiflorus</i>	57B	57B
<i>C. traskiae</i>	126E	126E
<i>C. fothergilloides</i>	var. <i>fothergilloides</i> 67B	var. <i>medranoanus</i> 82A
<i>C. ledifolius</i> var. <i>intermontanus</i>	102B	101A
<i>C. ledifolius</i> var. <i>ledifolius</i>	104	41B
<i>C. intricatus</i>	25	25
<i>Purshia tridentata</i>	34	34

RESULTS

Relationships in *Cercocarpus* based on AFLP characters

NJ analysis

The NJ analysis produced a phenogram that contained four clusters, corresponding with some exceptions to the four species complexes proposed by Lis (1992) (Figure 3.1). Cluster 1 consisted of members identified as belonging to the *Cercocarpus montanus* complex, with the exception of the inclusion of five individuals of the *C. ledifolius* complex highlighted by arrows in Figure 3.1. There was no support for the unity of any of the *Cercocarpus montanus* varieties described by Martin (1950), although there was support for the clustering of three of the four *C. montanus* var. *montanus* accessions (5, 8, and 10). Cluster 2 contained individuals identified as belonging to the *Cercocarpus betuloides* complex, with the inclusion of one individual of *C. fothergilloides* var. *fothergilloides* (18), one individual of *C. mexicanus* (134A), two individuals of *C. montanus* (variety *glaber* 17 and variety *paucidentatus* 75A), and two individuals of *C. traskiae* (126A and E) all identified by arrows in the Figure 3.1. The two individuals of *Cercocarpus minitiflorus* clustered together with moderate bootstrap support.

Cluster 3 consisted exclusively of individuals identified as belonging to the *Cercocarpus ledifolius* species complex. There was no support for the clustering of *Cercocarpus intricatus* or the varieties of *C. ledifolius*. Cluster 4

consisted exclusively of individuals representing the *Cercocarpus fothergilloides* complex and the newly described species *C. rzedowskii*.

The only cluster in the distance analysis that had bootstrap support above 50 was cluster 4 and the outgroup *Purshia* (Figure 3.1). The *Cercocarpus montanus* cluster 1 was found in the most derived position, sister to the *C. betuloides* cluster 2. The *Cercocarpus ledifolius* cluster 3 was found sister to both cluster 1 and 2. Cluster 4 sits sister to cluster 1, 2, and 3.

ME analysis

The minimum evolution distance analysis also yielded four clusters that corresponded to the four species complexes, although the relationship between the clusters was different (Figure 3.2). Cluster 1 contained individuals identified exclusively as belonging to the *Cercocarpus montanus* complex, although I did not find any support for the varieties as described by Martin (1950). Cluster 2, found sister to cluster 1, contained individuals of the *Cercocarpus ledifolius* complex and one individual identified as *C. montanus* var. *paucidentatus* (HEN1) identified by an arrow in Figure 3.2. As in the NJ analysis, we found no support for the varieties of *Cercocarpus ledifolius* (Holmgren 1987) or *C. intricatus*. Cluster 3, which was found sister to clusters 1 and 2, contained individuals of the *Cercocarpus betuloides* complex, with the inclusion of one individual of *C. fothergilloides* var. *fothergilloides* (18), one individual of *C. mexicanus* (134A), two individuals of *C. montanus* (variety *glaber* 17 and variety *paucidentatus*

75A), and two individuals of *C. traskiae* (126A and E) all identified by arrows in the Figure 3.2. The two individuals of *Cercocarpus minutiflorus* once again group together. The final cluster, cluster 4, contained individuals of the *Cercocarpus fothergilloides* complex. We found *Cercocarpus rzedowskii* grouped in this cluster with high bootstrap support.

MP analysis

The parsimony analysis resulted in one most-parsimonious tree of length 2643, consistency index of 0.134, and retention index of 0.484 (Figure 3.3). The parsimony tree contained four main groups. Clade 1 contained almost all members of *Cercocarpus montanus*, but did not support the monophyly of any of the varieties. Two individuals of *Cercocarpus montanus* (var. *paucidentatus* 75A and var. *glaber* 17) were found nested within the *C. betuloides* grade (Figure 3.3). Clade 1 also contained two individuals identified as *Cercocarpus ledifolius* var. *intermontanus* (individuals 37C and 98C), highlighted by arrows in Figure 3.3. Members of the *Cercocarpus betuloides* complex were found in a phylogenetic grade basal to clade 1. The two individuals of *Cercocarpus traskiae* were found in this grade, but not in a monophyletic group. The varieties of *Cercocarpus betuloides* were also not supported. Only *Cercocarpus minutiflorus* was supported as a monophyletic group with weak bootstrap support. The *Cercocarpus betuloides* grade also included one individual of *C. fothergilloides* var. *mojadensis* (individual 18), one individual of *C. mexicanus* (individual 134A), and the two

individuals of the *C. montanus* complex, all highlighted by arrows in Figure 3.3. Clade 2 was well supported, containing only individuals representing the Mexican taxa *Cercocarpus rzedowskii*, *C. fothergilloides*, and *C. macrophyllus*, but not *C. traskiae*. The species *Cercocarpus rzedowskii* was supported as a monophyletic group. *Cercocarpus macrophyllus* and *C. fothergilloides* were found to be paraphyletic. Clade 2 was the only major clade that had a bootstrap score above 50. Clade 3 contained individuals representing the *Cercocarpus ledifolius* complex. There was no support for the monophyly of the species *Cercocarpus intricatus* or the varieties of *C. ledifolius*.

In the MP analysis, the *Cercocarpus montanus* clade and *C. betuloides* grade form a monophyletic clade, but very weakly supported (bootstrap score less than 50). The *Cercocarpus fothergilloides* clade 2 was sister to the *C. ledifolius* clade 3, which together was sister to the *C. montanus* clade 1 and *C. betuloides* grade.

The three analyses (NJ, ME and MP) of the full AFLP dataset that included all 153 individuals support the four species complexes as described by Lis (1992) (Figure 3.4). All three analyses also supported the *C. fothergilloides* complex clustering at the base of the tree. The three analyses differed in the overall relationships between clusters. The NJ cluster analysis and the MP analysis supported a close relationship between the *Cercocarpus montanus* complex and the *C. betuloides* complex in the most derived position. The ME

analysis, in contrast, grouped the *Cercocarpus montanus* and *C. ledifolius* clusters in the most derived position.

In all three analyses, the *Cercocarpus betuloides* cluster consistently contained three members of the *C. fothergilloides* complex as identified by Lis (1992) (*C. fothergilloides* var. *mojadensis* (18), *C. traskiae* 126A and 126E), *C. mexicanus* (134A), and two individuals identified as *C. montanus* (variety *glaber* 17 and variety *paucidentatus* 75A).

Relationships in *Cercocarpus* based on ETS characters

MP analysis

The aligned dataset of 977 characters contained 147 parsimony informative characters for the 53 sequences included (Appendix D). The heuristic search resulted in 15,000 equally most parsimonious trees of 456 steps (Figure 3.5). Clade A contained a diverse set of taxa including representatives of *Cercocarpus montanus* and all the individuals of the Mexican taxa *C. rzedowskii*. Clade B contained multiple members of *Cercocarpus montanus*, *C. betuloides*, and *C. macrophyllus* and one individual of *C. fothergilloides* (67B) and one individual of *C. ledifolius* (individual 108C). Clade C was highly supported, consisting of two individuals of the *Cercocarpus ledifolius* complex (individual 25 and 41B). Clades D and E were also highly supported, containing individuals of the *Cercocarpus betuloides* complex. Clade F was weakly supported,

containing individuals of the *Cercocarpus ledifolius* species complex, one individual of *C. macrophyllus* (26) and one individual of *C. fothergilloides* (82A). The ETS sequence from one individual of *Cercocarpus ledifolius* var. *intermontanus* (111A clone 5) was found in an unresolved position with clades C, D, and E.

Clades A and B were sister to each other and are in the most derived position in the MP analysis. Clades C, D, and E were in unresolved positions, sister to clades A and B. Clade F was found in the most basal position. The only deep node that was supported was the branch leading to clades A and B (Figure 3.5).

ML analysis

The maximum likelihood analysis did not swap to completion, but was stopped with two most likely trees with likelihood scores of $-\ln 4223.56020$ in memory (Figure 3.5). The dashed line shows where the two most likely trees differed. Overall, the clades (clades A -F) recovered in the maximum parsimony analysis were supported by the ML analysis, however, the relationships between the clades were not consistent between the MP and ML reconstructions. In the ML analysis, clades A and B were again found in the most derived position. Clade F was recovered as sister to clades A and B, and included the ETS sequence from *Cercocarpus ledifolius* var. *intermontanus* (111A clone 5). Clade C was found more basal to clade F followed by clade E and D. Although the MP and

ML trees differed in topology, the branches that differ between reconstructions were either unresolved or have bootstrap support below 50 in the MP tree. This suggests there is little reason to prefer one topology over another.

Comparison of ETS and AFLP trees

The maximum parsimony analysis on the AFLP presence/absence data subset, which included just the 49 accessions matching the individuals sampled for the ETS dataset, resulted in 24 equally parsimonious trees of length 802. The strict consensus of these 24 trees is in Figure 3.6 along side the strict consensus tree of over 10,000 most parsimonious trees recovered from the ETS dataset. The AFLP MP strict consensus tree contained 4 major clades. Clade A contained almost all the individuals identified as part of *Cercocarpus montanus*, but did not support the varieties as described by Martin (1950). Clade A also contained two individuals of the *Cercocarpus betuloides* complex (*C. betuloides* var. *betuloides* 93C, and *C. minutiflorus* 57B). Clade B contained all of the members of the *Cercocarpus ledifolius* complex, but did not support the monophyly of the species *C. intricatus* or the varieties of *C. ledifolius* (Holmgren 1992). Clade B also contained two individuals of the *Cercocarpus betuloides* complex (*C. betuloides* var. *betuloides* 65D, and *C. betuloides* var. *blancheae* 123G). Clade C contained all of the strictly Mexican taxa of *Cercocarpus*. Clade D contained 2 individuals of *Cercocarpus montanus* (*C. montanus* var. *glaber* 17 and *C. montanus* var

paucidentatus 75A), 2 individuals of the *C. betuloides* complex (*C. betuloides* var. *betuloides* 128A and 120C), *C. traskiae* 126E, and *C. mexicanus* 134A.

The MP AFLP strict consensus was not congruent with the MP or ML ETS strict consensus phylogenies (Figure 3.6). Members of clade A from the MP AFLP strict consensus tree are found in 3 clades of the MP ETS strict consensus tree. Members of clade B are found in 5 separate ETS clades. Members of clade C were found 2 of the ETS clades and members of clade D were found in 4 clades from the ETS strict consensus tree.

Incongruence results

The results of the compare-2, the Templeton, and the Kishino-Hasegawa tests can be seen in Table 3.6. For the compare-2 test, 100 permutations of each dataset were tested on each constraint tree. We found significant results using both datasets, where the difference in fit for the dataset on its own tree versus the constraint tree was significantly better than the difference with permuted data. The Templeton and Kishino-Hasegawa tests yielded highly significant results for each dataset, signaling significant incongruence between both datasets tested.

Table 3.6: Results of the incongruence tests conducted on the AFLP and ETS datasets. ETS 1, ETS 2, ETS 3, and ETS 4 datasets refer to Table 3.5. The constraint trees used for these tests were bootstrap consensus trees containing only branches that had support over 50%.

COMPARE-2		KISHINO-HASEGAWA			TEMPLETON				
p-value	tree length difference	s.d.	t	p-value	N	+ sign ranks	- sign ranks	p-value	
AFLP vs ETS 1									
AFLP dataset	p=0.01	133	24.73017	5.3780	p=0.0001	88	3188.5	727.5	p=0.0001
ETS 1 dataset	p=0.01	290	37.78070	7.6759	p=0.0001	98	4833.5	17.5	p=0.0001
AFLP vs ETS 2									
AFLP dataset	p=0.01	126	23.81775	5.2902	p=0.0001	88	3176.5	739.5	p=0.0001
ETS 2 dataset	p=0.01	287	37.60504	7.6320	p=0.0001	96	4639.5	16.5	p=0.0001
AFLP vs ETS 3									
AFLP dataset	p=0.01	117	24.04435	4.8660	p=0.0001	88	3069	847	p=0.0001
ETS 3 dataset	p=0.01	283	37.36917	7.5731	p=0.0001	95	4544	16	p=0.0001
AFLP vs ETS 4									
AFLP dataset	p=0.01	119	24.87939	4.7831	p=0.0001	87	2993.5	834.5	p=0.0001
ETS 4 dataset	p=0.01	283	37.87471	7.4720	p=0.0001	94	4448.5	16.5	p=0.0001

The effect of taxon sampling on tree topology

The three AFLP reduced datasets produced three different tree topologies (Figure 3.7 A, B, C), which differed with respect to the relationships between *Cercocarpus montanus*, *C. betuloides*, and *C. ledifolius*, and for which all possible topological relationships were recovered. The MP tree produced from AFLP reduced dataset A was the only one that did not have bootstrap support above 50

for all branches. The two ETS reduced datasets produced different tree topologies as well. We found no common clades between the trees produced from ETS datasets D and E.

DISCUSSION

The goals of this project were to generate molecular characters to help define species and varietal boundaries in *Cercocarpus* and to test some hypotheses generated by other authors about the evolution and taxonomic affinities of certain taxa. The three analyses of the full AFLP dataset generated support for four species complexes in *Cercocarpus* (Lis 1992). The ETS analyses did not support these complexes. Phylogenetic trees produced from the AFLP and ETS datasets were significantly incongruent. We also found that taxon sampling could have had a large effect on tree topology.

Taxonomic implications

The *betuloides* complex

Martin (1950) considered the taxa *Cercocarpus betuloides*, *C. minutiflorus* to be varieties of *C. montanus*. Lis (1992) reorganized these taxa after a detailed study of the leaf architecture. Lis recognized *Cercocarpus betuloides* as having three varieties, two of which are included in this study (*C. betuloides* var. *betuloides* and *C. betuloides* var. *blancheae*). He also recognized *Cercocarpus minutiflorus* at the specific level. We found that these taxa were distinct from the

C. montanus complex in the three AFLP analyses, however, they were found at the base of a clade including the *C. montanus* complex in the AFLP MP analysis and sister to the *C. montanus* cluster in the NJ and suggesting a close relationship. We only had two samples of *Cercocarpus minutiflorus* in our AFLP data, yet in all three AFLP analyses we found those two samples united with moderate bootstrap support. Lis (1992) found that *Cercocarpus minutiflorus* shared leaf architecture character states with both *C. traskiae* and *C. betuloides*, but was reluctant to assign it as a variety of either species. *Cercocarpus minutiflorus* consistently grouped with *C. betuloides* in our analyses. *Cercocarpus minutiflorus* differs from *C. betuloides* in its distribution (restricted range from San Diego to northern Baja California, Mexico) and glabrous leaf characteristics. The data presented here would suggest *Cercocarpus minutiflorus* may be a distinct group, but appears to be closely related to *C. betuloides*.

We found no support for the varieties of *Cercocarpus betuloides* defined by Lis (1993). This is a surprising result because Searcy (1967) found that these taxa retained their leaf traits in a common garden experiment, and hypothesized distinct genotypes for the two groups.

The *ledifolius* complex

Holmgren (1987) recognized two varieties of *Cercocarpus ledifolius*, *C. ledifolius* var. *intermontanus* and *C. ledifolius* var. *ledifolius*. The characters used to distinguish them consisted of leaf size, geographic location, and leaf vestiture

(Table 3.2). *Cercocarpus intricatus* has been raised to specific status in many of the western floras, a decision Holmgren (1987) agreed with, based on the divergent leaf characteristics and ecological requirements. Based on the AFLP and ETS data presented above, we have no support for recognizing the two varieties of *Cercocarpus ledifolius* or *C. intricatus*.

The continuous variation seen between *Cercocarpus ledifolius* and *C. intricatus* has been postulated as evidence of hybridization and hybrid swarms between the two taxa (Schneider 1905b, Brayton and Mooney 1966, Davis and Brotherson 1991). In his revision of *Cercocarpus*, Schneider (1905b) believed the intermediate forms may have been of hybrid origin, and he designated them as *Cercocarpus ledifolius* var. *intercedens*. This taxon was later synonymized with *Cercocarpus ledifolius* var. *ledifolius*. If we accept this hypothesis, we would expect each parental type (*Cercocarpus ledifolius* var. *intermontanus* and *C. intricatus*) to cluster independently in the NJ and ME phenograms. The intermediate leaf shape (*Cercocarpus ledifolius* var. *ledifolius*) would be expected to cluster in between or at the base of the parental taxa due to the presence of both sets of AFLP markers in the hybrid individual. Further, instead of clustering in between, the individual with intermediate morphology may group with one of the parents if considerable backcrossing had occurred. We do not see this distribution of individuals in any of our AFLP or ETS reconstructions. Instead we see a single cluster (with some exceptions) in the NJ and ME AFLP trees, and a single clade

in the MP AFLP tree, which would suggest leaf diversity seen in *Cercocarpus* might be the result of ecotype variation, where microhabitat factors have selected certain leaf shapes.

The *montanus* complex

Three of the four individuals of *Cercocarpus montanus* var. *montanus* (5, 8, and 10) form a moderately supported cluster (bootstrap value 65) in the AFLP NJ and ME trees and a moderately supported clade (bootstrap value 66) in the AFLP MP tree (Figure 3.1, 3.2, and 3.3). The fourth individual of *Cercocarpus montanus* var. *montanus*, individual 28A, was found in the larger *C. montanus* cluster in the NJ and ME trees and the larger clade (*C. montanus* clade 1) in the AFLP MP tree, but is not directly a part of a monophyletic group. In the phylogenies based on the ETS dataset, all members of *Cercocarpus montanus* var. *montanus* were found in an unresolved clade (part of clade B) in both the MP and ML analyses. These results suggest that *Cercocarpus montanus* var. *montanus* may indeed be a real taxonomic entity, although the support for this claim is weak. *Cercocarpus montanus* var. *montanus* is winter deciduous, a character not listed in Table 3.2, which separates it from the other varieties of *C. montanus*.

Cercocarpus montanus var. *paucidentatus* and *C. montanus* var. *glaber* do not unite in monophyletic groups in any analysis. As illustrated in Table 3.2, these taxa overlap in many of the characters used to differentiate them. Although

not studied in *Cercocarpus montanus*, leaf characters have been shown to vary based on environment in *C. betuloides* (Searcy 1977) and *C. ledifolius* (Davis and Brotherson 1991). We argue, based on the data presented here, that the varieties currently defined by Martin (1950) are not real taxonomic entities.

The *fothergilloides* complex

Richard Lis (1992) described a relationship between the taxa *Cercocarpus fothergilloides*, *C. mojadensis*, *C. pringlei*, *C. macrophyllus* and *C. traskiae* based on leaf characters. He proposed that these taxa were all part of one species complex he called the *C. fothergilloides* complex. James Henrickson is currently revising this group and has recognized the same overall taxa, making a few taxonomic changes (Madroño ined.), including the recognition of *Cercocarpus mojadensis* as a variety of *C. fothergilloides*. In our study, based on the AFLP data, *Cercocarpus fothergilloides* var. *mojadensis* (18) and *C. traskiae* (126A and E) consistently grouped with the *betuloides* complex. The other members of this complex, *Cercocarpus fothergilloides* var. *medranoanus*, variety *fothergilloides*, and *C. macrophyllus* consistently grouped together with high bootstrap support. This result gives strength to Lis' claim that the *Cercocarpus fothergilloides* complex exists, if one removes *C. traskiae*, and possibly, *C. fothergilloides* var. *mojadensis*, but more expanded sampling is needed before any taxonomic decisions are warranted.

The ETS dataset did not support this complex as currently defined. *Cercocarpus fothergilloides* var. *fothergilloides* (67B) and *C. macrophyllus* (69A and 69B) grouped with members of *C. montanus*. *Cercocarpus fothergilloides* var. *medranoanus* (82A) and *C. macrophyllus* (26) grouped with *C. ledifolius* and *C. intricatus*.

Cercocarpus rzedowskii* and *C. mexicanus

In 1987, Henrickson described two new species of *Cercocarpus*, *C. rzedowskii* from southern Tamaulipas, and *C. mexicanus* from southern Coahuila (see Figure 1.12). He suggested a close relationship between *Cercocarpus rzedowskii* and *C. ledifolius*, based on overall leaf morphology. He also postulated a possible relationship between *Cercocarpus mexicanus* and *C. montanus*. We did not find any support for these relationships. In the ETS phylogenies, *Cercocarpus rzedowskii* is nested deeply in a clade with members of the *C. montanus* complex, while in the AFLP trees, we find *C. rzedowskii* united with members of the *C. fothergilloides* complex. The one individual of *Cercocarpus mexicanus* (134A) included in this study is found deeply nested within the *C. betuloides* complex in all three analyses using the AFLP data.

ETS considerations

Within the ETS data set, we had six individuals that were represented by two cloned ETS sequences (Figure 3.4). In four of the six, the two clones united

in a monophyletic group in both the MP and ML analyses. The other two, *Cercocarpus betuloides* var. *betuloides* (62D clone 2 and 3) and *C. ledifolius* var. *intermontanus* (111A clone 4 and 5) did not unite, although in the ML analysis, the two clones from 111A did fall out in the same clade. Explaining these results depends on how concerted evolution and gene flow (hybridization) are operating in the ETS region of the nrDNA in *Cercocarpus*. If concerted evolution is operating quickly (Wendel 1989), the divergent clones are most likely evidence of recent gene flow, especially if the two clones reside in different, highly supported clades. If concerted evolution is proceeding more slowly, the divergent clones could be evidence of multiple ETS types (orthologous) due to mutation. In this case, we would expect the different clones to inhabit the same clade, but not necessarily as sister to each other.

The two clones of *Cercocarpus betuloides* var. *betuloides* (62D) resided in clades (clades D and E) that were almost entirely composed of members of the *C. betuloides* complex. The relationship between the two clades is not resolved using our ETS dataset, so it is possible that the two clades (clades D and E) are sister clades. However, each clade is highly supported as independent and, therefore, it could be that the position of the two clones is a result of gene flow between different geographic populations of *Cercocarpus betuloides* present in clades D and E. Members of *Cercocarpus betuloides* var. *betuloides* present in

clade D are located in the foothills of the Sierra, while members found in clade E are located in Arizona (Table 3.1) (please see Figure 1.10).

The two clones of *Cercocarpus ledifolius* var. *intermontanus* (111A clone 4 and 5) do not unite in the ETS MP tree, yet in the ETS ML analysis, the two clones reside in the same clade. Although unexpected, this result does not suggest hybridization as strongly as the results above. Multiple intra-specific ETS types have been reported in other studies (Linder et al. 2000) from *Argyranthemum* and *Helianthus*. The ETS types coalesced within a species, leading to the conclusion that concerted evolution was working, but not as rapid as in the ITS or coding regions of the rDNA repeat (Linder et al. 2000). The two clones of *Cercocarpus betuloides* var. *betuloides* (62D) appear to coalesce with *C. betuloides* species complex, but the clones from *C. ledifolius* var. *intermontanus* do not. We might explain the two clones recovered from *Cercocarpus ledifolius* var. *intermontanus* as evidence of multiple ETS types (orthologous) due to mutation, and the two clones were not homogenized by concerted evolution.

ETS versus AFLP

Direct comparison of the AFLP and ETS phylogenetic reconstructions for the same individuals showed a high degree of incongruence (Figure 3.5) (Table 3.6). Although the characters used to reconstruct the phylogenies came from different sources, such a high degree of incongruence was unexpected. The AFLP reconstruction recovered, for the most part, each of the species complexes that are

currently defined by morphology and geography (Lis 1992), although it did not recover the varieties. The ETS phylogenies did not recover monophyletic species or varieties. We have identified three possible sources for this incongruence between the ETS and AFLP analyses. First, it may just be an artifact of the lack of real support for most of the branches in either analysis. Few branches in either the NJ, ME, or MP reconstructions of the AFLP dataset and some branches in the ETS MP reconstruction have bootstrap support over 50.

Second, the incongruence may be due to an excess of homoplasy in the AFLP dataset. Since homoplasy can be generated in an AFLP analysis if non-identical markers of equal length are assumed to be homologous - this homoplasy should be magnified the greater the evolutionary distance is between samples - incongruence between the AFLP and ETS trees could have been caused by these factors. We found that when we compared the NJ, ME, and MP reconstructions of the AFLP dataset in this study, the topologies were incongruent. Koopman et al. (2001) suggests that in these cases, it is important to examine the strength of support for branches in the MP tree by bootstrap support, assuming the incongruence is due to homoplasy and will be evident by having low bootstrap scores. Branches that are incongruent between trees that have low support should be discarded and not used for conclusions about species relationships. Based on this approach, only the *Cercocarpus fothersgilloides* clade could be supported (Figure 3.4). It is clear that the AFLP dataset contains homoplasy and also

contains only a few characters that are supporting the deeper branches in the MP reconstruction. The combination of these two traits leads to low bootstrap scores, and may have led to the incongruence we have seen.

Third, the incongruence between the ETS and AFLP trees could be a result of hybridization and lack of concerted evolution, in which ETS sequences we assume to be orthologous are not. Hybridization has been documented in *Cercocarpus* (Rieseberg 1989, 1994). Unlike the AFLP markers, the ETS dataset is a representation of a small non-coding area in the nrDNA. The phylogenetic history of this small non-coding region may be quite different than the species in which it is found. We have already cited examples above where ETS clones from a single individual did not unite in a monophyletic group, suggesting that hybridization and the maintenance of orthologous ETS sequences does occur in *Cercocarpus*, which may make the ETS an unsuitable nuclear marker for phylogenetic reconstruction of species relationships in *Cercocarpus*.

Taxon and population sampling in recently diverged groups

A major issue in phylogenetic inference is taxon sampling, which is usually defined as the selection of a small number of representative specimens for higher-level phylogenetic analysis used to infer evolutionary history. Insufficient taxon sampling is often cited as a major source of error in phylogenetic analysis (Murphy et al. 2001, Saunders and Edwards 2000, Soltis et al. 1999), and can occur because of constraints on time, money, or rarity (Rosenberg and Kumar

2001). The major argument for increased taxon sampling is that inclusion of more taxa counteracts long-branch attraction, where long, unrelated branches associate erroneously in the phylogenetic analysis. We argue here that in recently evolved groups, or groups with high rates of gene flow between taxa, increased population sampling within taxa is important for understanding the total variation present and properly delimiting species. If variation exists in the marker chosen for phylogenetic reconstruction that is shared between species, relationships between taxa can be reconstructed erroneously, because the individuals chosen to represent them (Wendel and Doyle 1998). These errors can be lessened if multiple populations are sampled.

In analyses based on both the ETS and AFLP datasets, we found that relationships between taxa and within a defined taxon depended on the individuals used to represent the taxon (Figure 3.6). MP trees created from different subsets of the taxa from the AFLP dataset (Figure 3.6) showed that we could reconstruct all three trees that represent the possible relationships of *C. betuloides*, *C. ledifolius*, and *C. montanus*. Even more striking, we produced two highly incongruent, but well supported MP trees based on different subsets of ETS sequences. These incongruencies can only be discovered because we sampled more than one individual per taxon, and highlight how increased population sampling within taxa is very important for correct inferences of both evolutionary history and taxon delimitations.

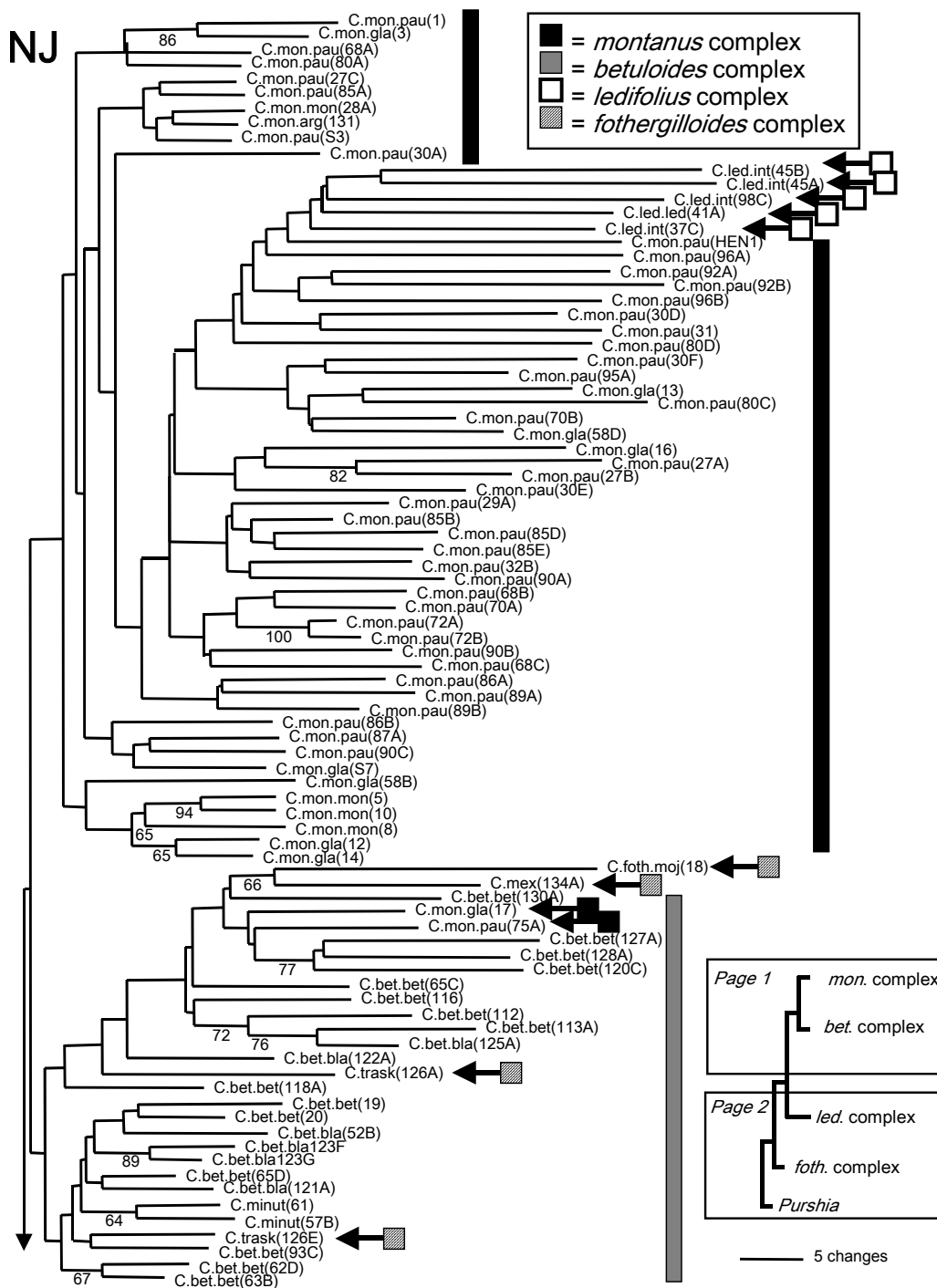


Figure 3.1. A neighbor joining (NJ) phenogram based on 376 presence/absence AFLP characters from three selective primer combinations for 153 individuals of *Cercocarpus*. The species names of the individuals have been abbreviated as in Figure 2.4. Numbers in parentheses refer to the collection number (Table 3.1). Numbers below the branches are a measure of bootstrap support. Only bootstrap values above 50% are reported.

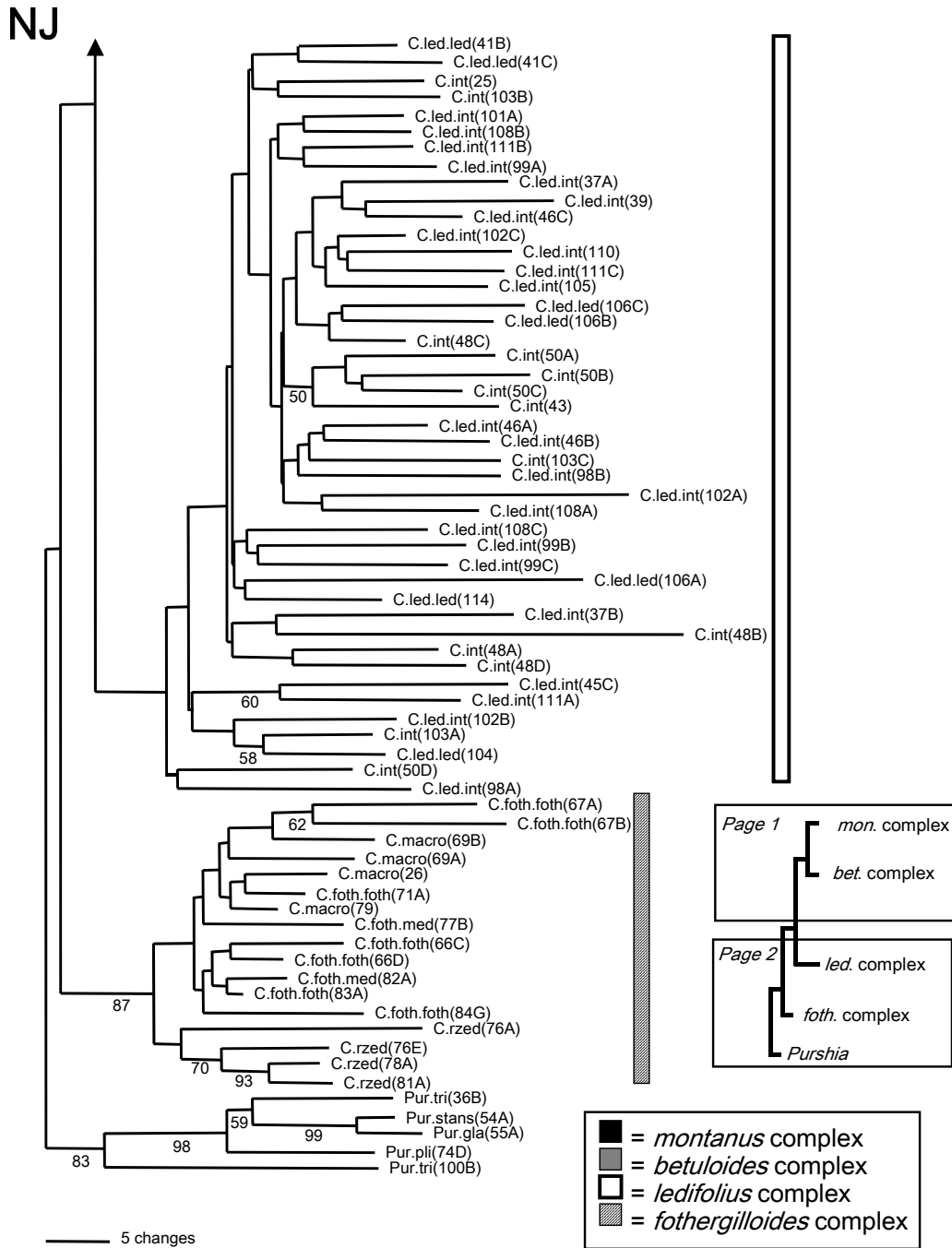


Figure 3.1: (continued)

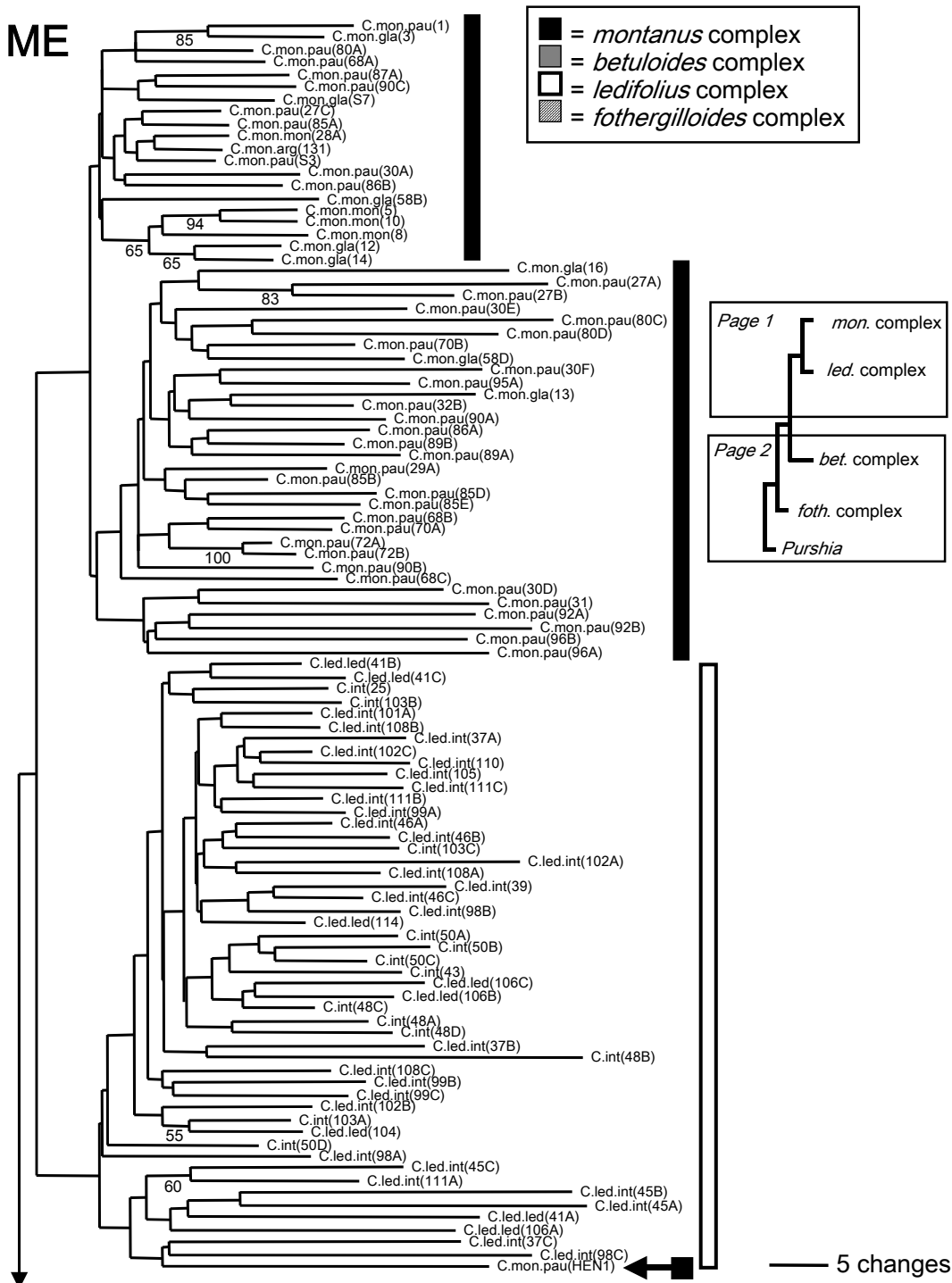


Figure 3.2. A minimum evolution (ME) phenogram based on 376 presence/absence AFLP characters from three selective primer combinations for 153 individuals of *Cercocarpus*. The species names of the individuals have been abbreviated as in Figure 2.4. Numbers in parentheses refer to the collection number (Table 3.1). Numbers below the branches are a measure of bootstrap support. Only bootstrap values above 50% are reported.

ME

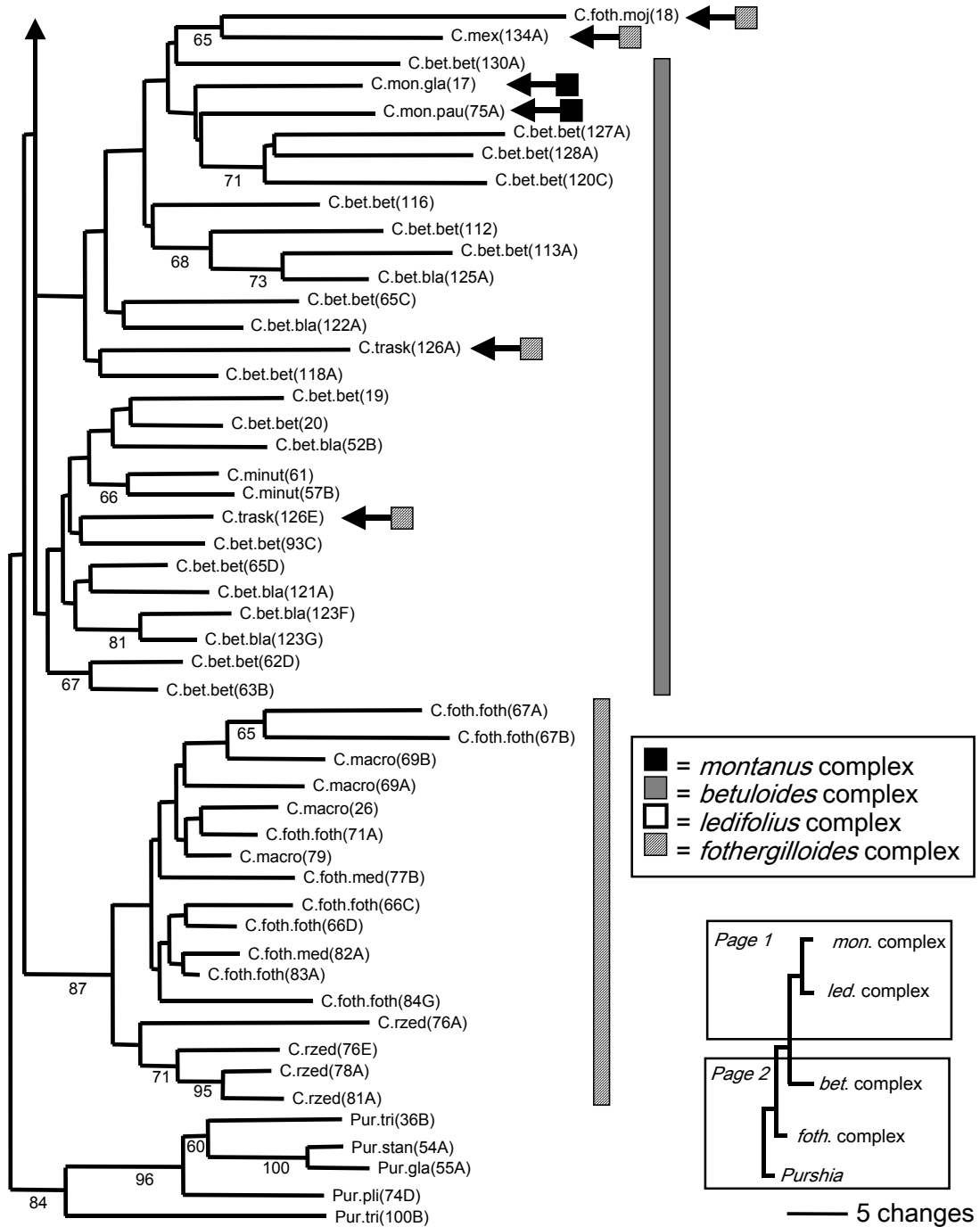


Figure 3.2: (continued)

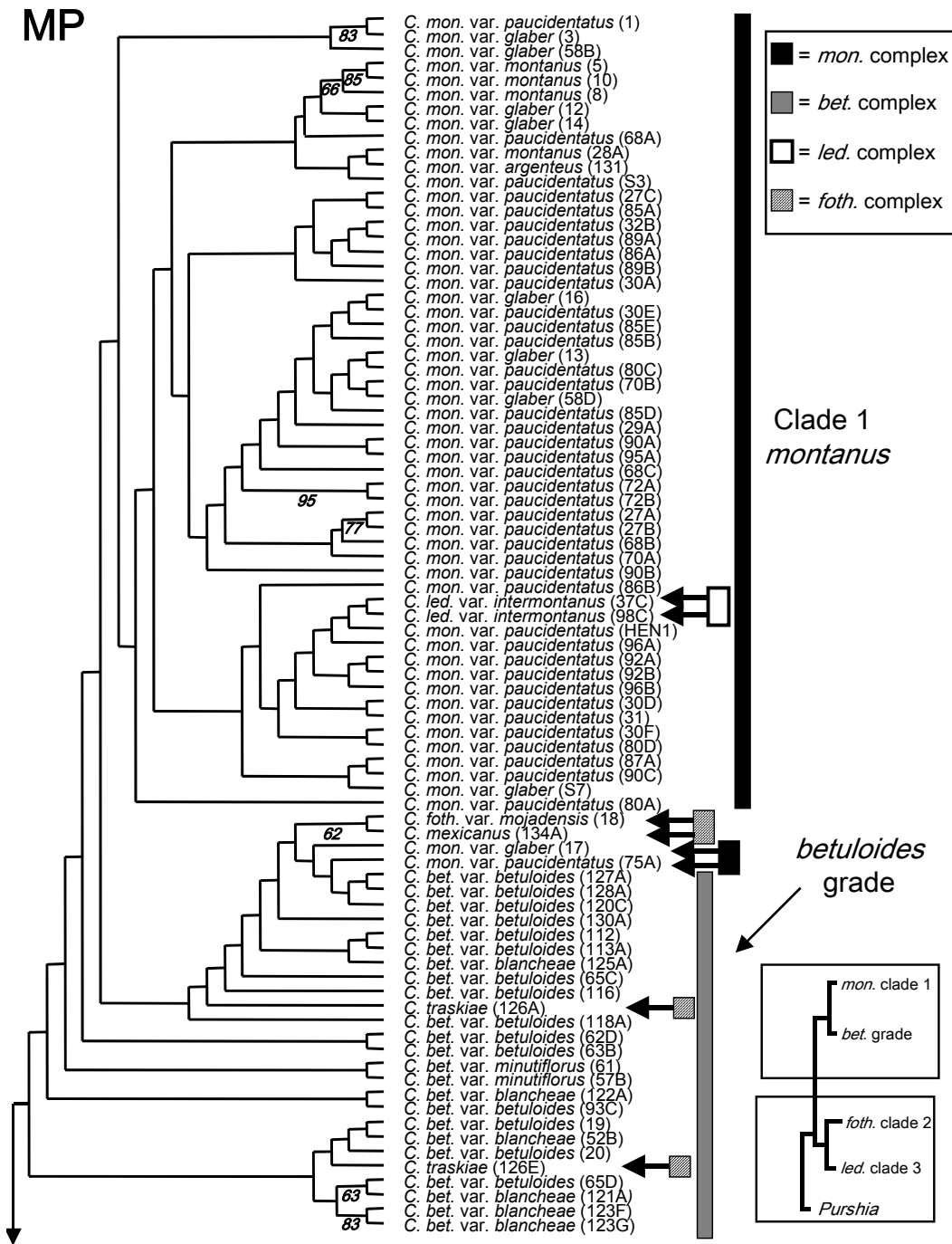


Figure 3.3. The single most parsimonious tree of length 2643 resulting from analysis of 376 presence/absence AFLP characters. Taxonomic identities have been abbreviated where C. = *Cercocarpus* and P. = *Purshia*. The three letters following are the first three letters of the species name. Numbers in parentheses refer to collection number (Table 3.1). The number of steps supporting the branch is shown above, the bootstrap support over 50 is shown below each branch.

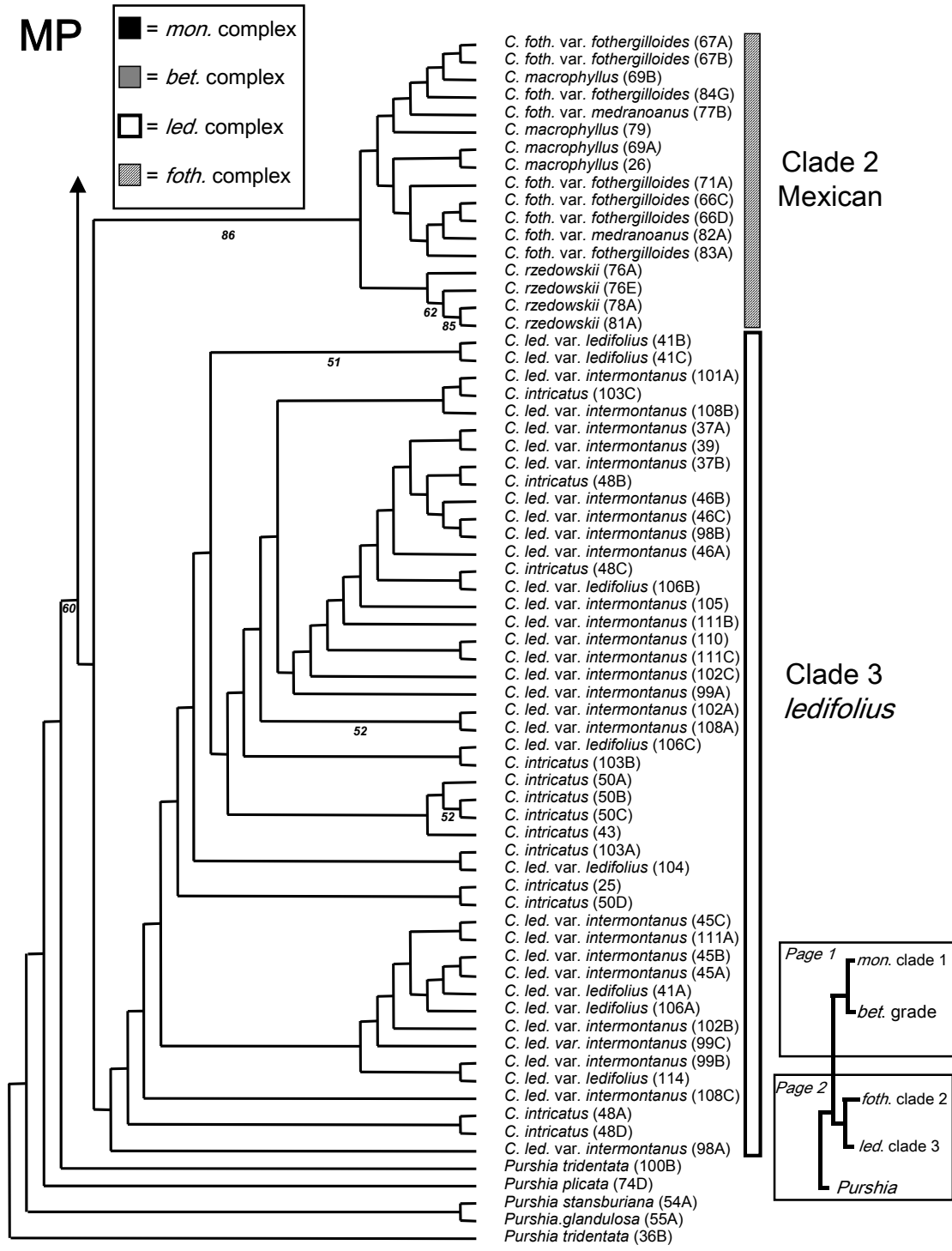


Figure 3.3: (continued)

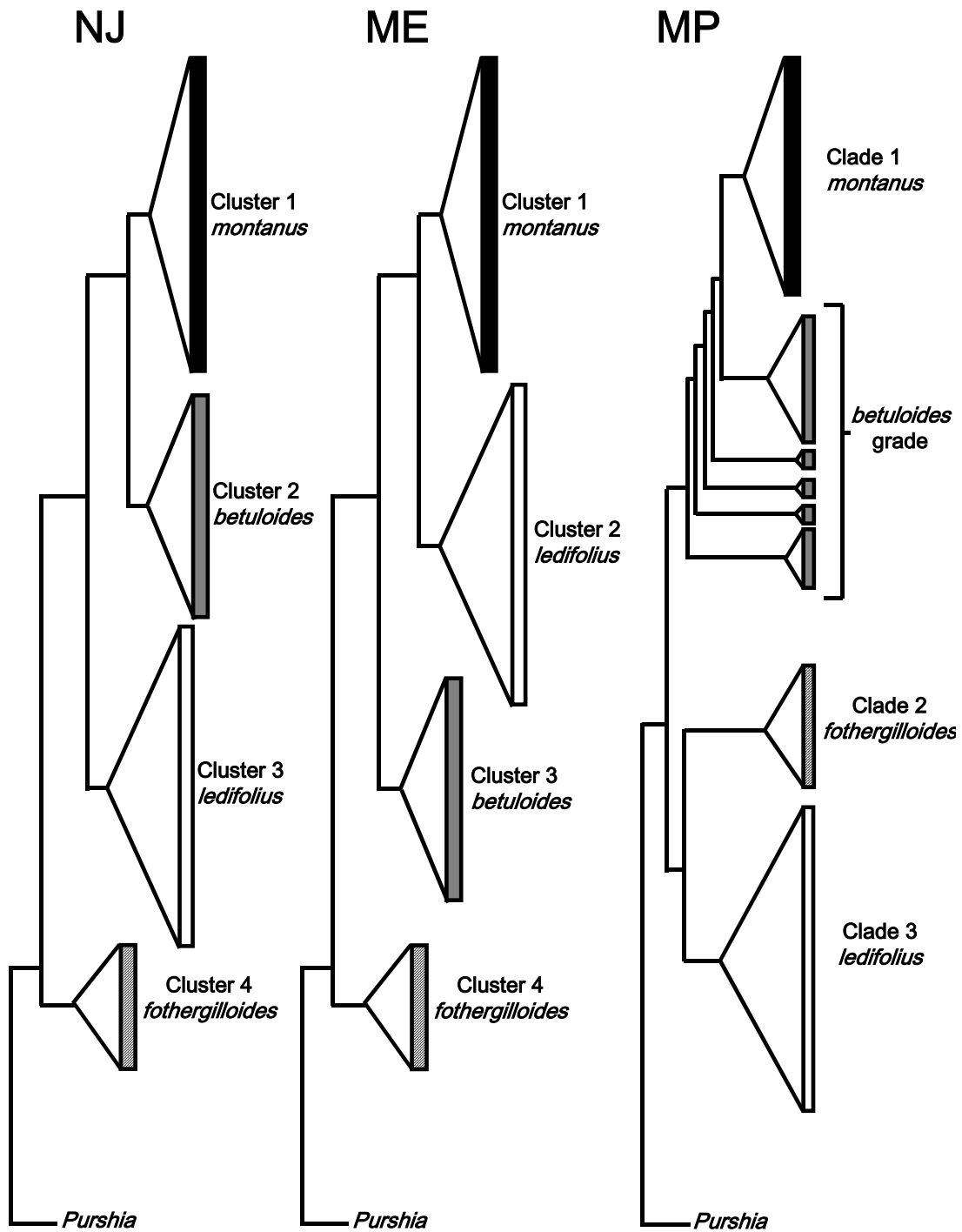


Figure 3.4. A comparison of the NJ phenogram, ME phenogram, and MP cladogram based on analyses of the AFLP characters. The three trees are stylized representations of Figures 3.1, 3.2, and 3.3. Colored bars are used to represent the 4 species complexes.

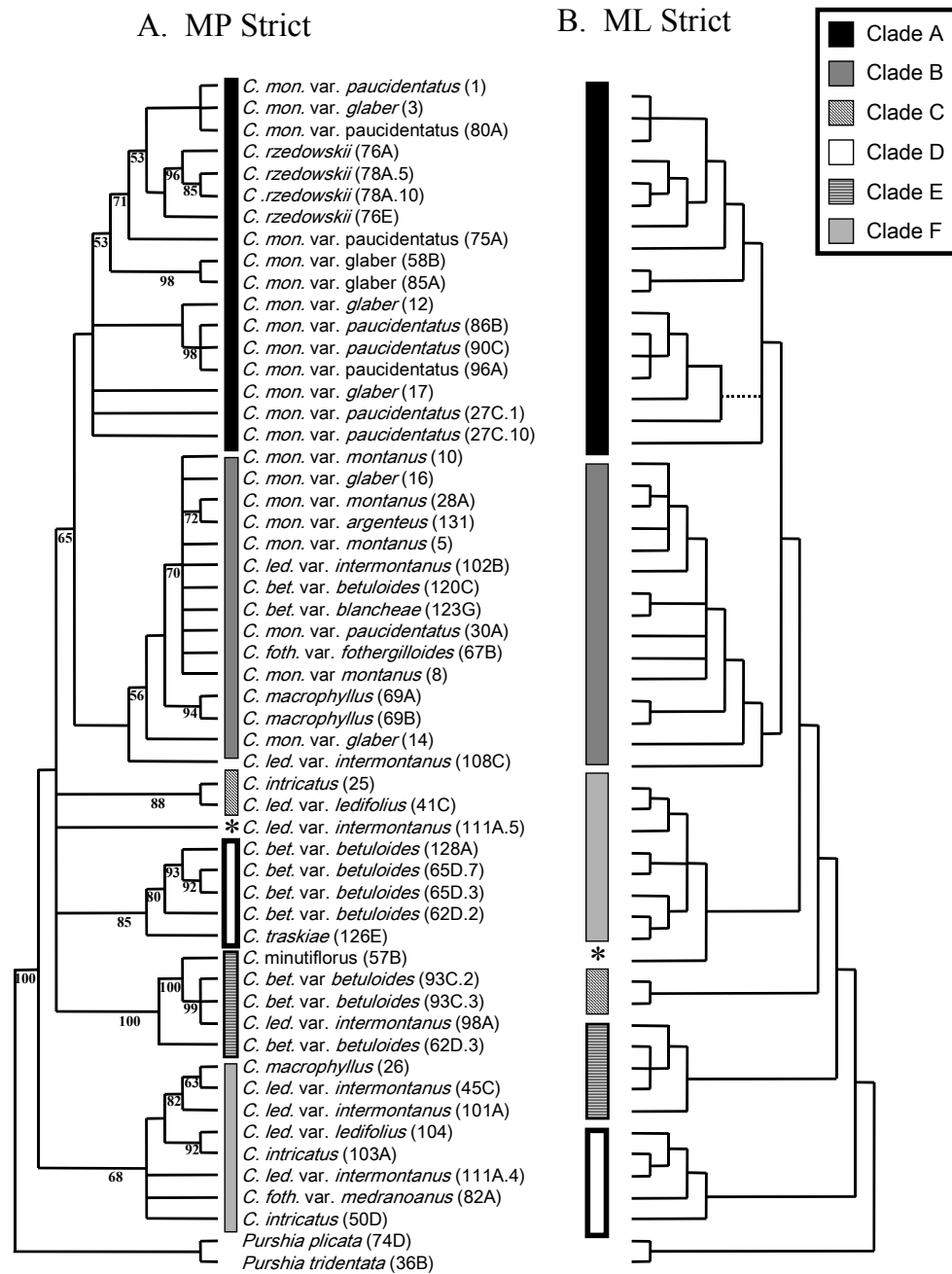


Figure 3.5. Strict consensus trees recovered from the MP and ML analyses conducted on the ETS dataset. A. The maximum parsimony (MP) strict consensus tree of over 15,000 most parsimonious trees of length 456. The numbers below the branches show bootstrap support above 50. The MP strict consensus tree has six major clades, labeled A through F, identified by the colored bars. The large asterisks denotes clone number 5 from *C. ledifolius* var. *intermontanus* 111A, which changes position between analyses. B. One of two most likely trees with a likelihood score of $-\ln 4223.56020$ recovered in the maximum likelihood analysis. The branch that collapses in the consensus tree is shown as a dotted line. Bootstrap support for the branches in the ML analysis were not estimated due to the computational time required.

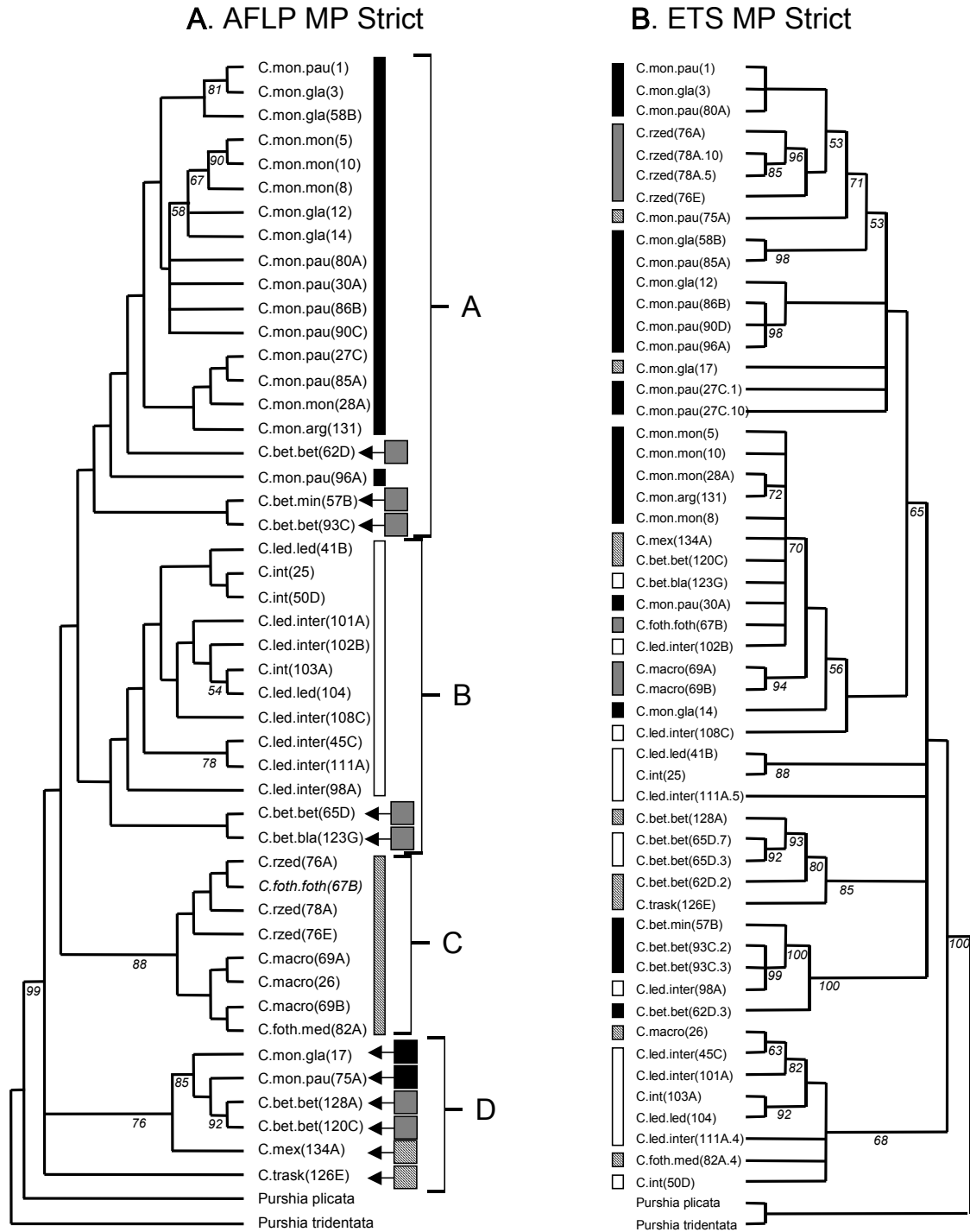


Figure 3.6. Comparison of the AFLP MP strict consensus tree and the ETS MP strict consensus tree. The names of the taxa have been abbreviated as in Figure 2.4. The number in parentheses at the end of the name refers to the collection number (Table 3.1). The colored bars indicate the species complex designations of Lis (1992).

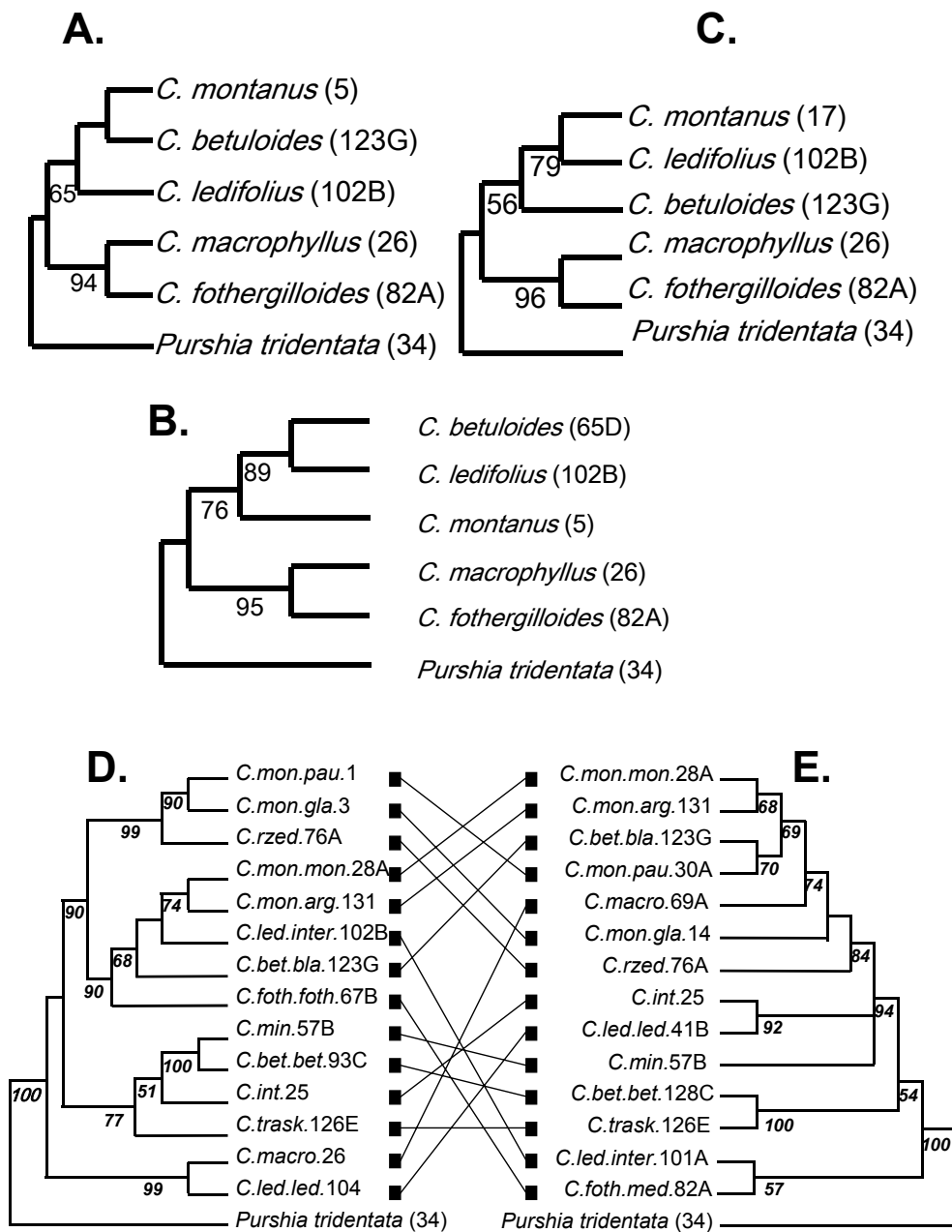


Figure 3.7. MP trees reconstructed using both AFLP and ETS characters showing the importance of taxon sampling in *Cercocarpus*. Figures A, B, and C are single most parsimonious trees based on exhaustive searches in PAUP, based on 376 presence/absence AFLP characters and individuals sampled in Table 3.3. Figure A, B, and C correspond to AFLP datasets A, B, and C in Table 3.3. Figure A had a length of 122. Figure B had a length of 114. Figure C had a length of 128. Numbers below the branches are a measure of bootstrap support based on 1000 replicates above 50. Figures D and E are trees that resulted from a heuristic MP search based on the aligned 977 ETS data set. Figures D and E correspond to ETS reduced datasets D and E in Table 3.4. Figure D is a strict consensus of 3 most parsimonious trees of length 201. Figure E is the most parsimonious tree of length 185. Numbers below the branches are measures of bootstrap support based on 100 replicates above 50.

Chapter 4: A phylogeny for *Cercocarpus* H.B.K (Rosaceae) based on cpDNA sequence data and the implications of congruence and incongruence with AFLP and ETS datasets

INTRODUCTION

The benefits of creating and comparing independent datasets of molecular characters for plant phylogenetic reconstruction are well discussed and follow two major lines of reasoning (Hillis 1987, Wendel and Doyle 1998). First, when phylogenetic topologies from independent molecular datasets are congruent, it is usually considered a reliable topology representing the evolutionary history of the taxa being studied (Hillis 1987). Under this condition, many researchers would support combining the datasets in order to improve support for particular branches or reconstruct groupings that were not present in the original, independent topologies (Bull et al. 1993, Kellogg et al. 1996). Second, multiple, independent datasets may show well-supported incongruence, suggesting different evolutionary histories for the molecular markers involved (Wendel and Doyle 1998). It is important to consider that most of the molecular characters used to reconstruct phylogenetic hypotheses come from DNA sequence data, representing small stretches of DNA. These small pieces of DNA may have different evolutionary histories from the species/genome within which they reside, as a result of past hybridization events or lineage sorting. Therefore, in both cases,

multiple independent molecular datasets are desirable. This study presents the results of a chloroplast-based phylogeny and compares it with two independent nuclear phylogenies for the genus *Cercocarpus*. Congruencies and incongruencies between the three phylogenies are examined.

Recently, sequence data from the external transcribed spacer (ETS) of the nuclear ribosomal DNA (nrDNA) and presence/absence data generated from AFLP markers were used to reconstruct a nuclear phylogenetic hypothesis for *Cercocarpus* (Chapter 3). Unexpectedly, these phylogenies, based on two different nuclear datasets, were not congruent (Chapter 3). Three proposed reasons for the incongruence between the ETS and AFLP trees were presented, including; 1) an artifact of the data due to lack of support for branches, 2) excess homoplasy in the AFLP dataset, and 3) hybridization and a lack of concerted evolution, whereby ETS sequences assumed to be orthologous were not. Due to the incongruence between nuclear markers, a comparison of both the ETS and AFLP phylogenies to a third, truly independent molecular dataset is required. For the first case, after comparison to an independent phylogeny, we may identify a few individuals that were incorrectly reconstructed in both the ETS and AFLP trees due to a lack of phylogenetically informative variation, but were fully resolved in the independent tree. We might find the significant incongruence between the ETS and AFLP trees disappears after the individuals causing the incongruence are excluded.

For the second case, where the incongruence was produced from excess homoplasy in the AFLP dataset, we might find that ETS tree topology to be congruent to an independent phylogeny for *Cercocarpus*, but the AFLP tree will be incongruent due to its inherent homoplasy. For the third and last case, significant incongruencies between AFLP, ETS and an independent phylogeny may suggest different evolutionary histories for the molecular markers involved and yield insight into past evolutionary processes, including hybridization and introgression (Rieseberg et al. 1996).

Plastid sequence data is the choice most studies of plants make for a molecular marker that is independent of nuclear datasets. Recently, many non-coding chloroplast markers have been developed for widespread use in plant systematics (Soltis and Soltis 1998) and have been shown to be informative at inter and intra-specific levels (Hamilton 1999). Given three independent phylogenies, areas of topological congruence and incongruence may be identified (considering only well supported branches) that may allow the identification of the source of significant incongruence between the AFLP and ETS, and also past evolutionary processes.

A chloroplast phylogeny will also help to test hypotheses generated by previous authors about taxon relationships and evolutionary trends in *Cercocarpus* discussed in Chapter 3. Briefly, we would like to address whether or not *Cercocarpus betuloides* is indeed a separate taxon from *C. montanus* as the

AFLP and ETS datasets suggested. These results would indicate that the taxonomic changes made by Lis (1993) were warranted. A second question asks if *Cercocarpus traskiae* actually has closer affinities to *C. betuloides* than to *C. fothergilloides* as the AFLP and ETS datasets implied. Third, *Cercocarpus rzedowskii*, a newly described species from Tamaulipas, Mexico (Henrickson 1987) was found to cluster with *C. fothergilloides* in the AFLP trees, yet with *C. montanus* in the ETS dataset. *Cercocarpus mexicanus*, the other newly described species was found to cluster with *C. betuloides*. A cpDNA tree may help us determine the true relationships for these two taxa. Lastly, Stutz (1974) hypothesized that both *Cercocarpus montanus* and *C. intricatus* were formed by peripheral isolation from *C. ledifolius*. I did not find any support for this claim in either AFLP or ETS phylogenies, yet I will address this claim again with an independent cpDNA phylogeny.

MATERIALS AND METHODS

Taxon Sampling and DNA extraction

The same DNA accessions were used as in Chapter 3 with a few additions. Sampling for this cpDNA study attempted to overlap with both the ETS and AFLP datasets previously sampled (Table 4.1). One sample from each of two species of *Purshia*, *P. tridentata* and *P. plicata*, and one sample of *Chamaebatia foliolosa* were used as outgroups in this study. These outgroups were chosen

based on the documentation of a close relationship between *Cercocarpus*, *Purshia*, and *Chamaebatia* in molecular phylogenies of the Rosaceae (Morgan et al. 1994, Evans et al. 2000). Fifty-four individuals, including the three outgroup individuals, are included in the cpDNA dataset. The 51 *Cercocarpus* individuals represent 46 populations of 17 taxa (Table 4.1).

Table 4.1: A list of the taxonomic identity, code, location, and collector and voucher number of the individual leaf samples used in this study. BVH is the author and JH is James Henrickson. All vouchers are housed at the plant resources center (UTEX)* denotes those individuals that are in common with the AFLP dataset and were used in the cpDNA comparison. † denotes those individuals that are in common with the ETS type 1 dataset and were used in the cpDNA comparison.

taxon	code	location	collector and voucher
<u>montanus complex</u>			
<i>C. montanus</i> var. <i>montanus</i>	†*5	Castle Rock, Douglas Co., CO USA	BVH 5
	†*8	Hartville, Platte Co., WY USA	BVH 8
	†*10	Dewey, Custer Co., SD USA	BVH 10
	†*28A	Bailey Canyon, Otero Co., NM USA	BVH 28
<i>C. montanus</i> var. <i>glaber</i>	†*3	Jame, Coahuila, MX	JH 20439b
	†*12	Chisos Mts., Brewster Co., TX USA	BVH and JH 20662
	†*14	Chisos Mts., Brewster Co., TX USA	BVH and JH 20663
	†*15	Chisos Mts., Brewster Co., TX USA	BVH and JH 20665
	†*58B	Leaky, Real Co., TX USA	JH 22275
	*S.7	Guadalupe Mts., Culberson Co., TX USA	CJ 18482
<i>C. montanus</i> var. <i>paucidentatus</i>	†*1	La Peridida, Tamaulipas, MX	JH 19218
	†*27C	Guadalupe Mts., Eddy Co., NM USA	BVH 27
	†*30A	Pinos Altos, Grant Co., NM USA	BVH 30A
	†*75A	Inturbide, Coahuila, MX	BVH and JH 22429
	†*80A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22457
	†*86B	Tunnel Springs, Bernalillo Co., NM USA	BVH 86
	*S.3	Guadalupe Mts., Culberson Co., TX USA	CJ 24804

<i>C. montanus</i> var. <i>argenteus</i>	†*131	Guadalupe Mts., Culberson Co., TX USA	McVaugh 7424
<i>betuloides</i> complex			
<i>C. betuloides</i> var. <i>betuloides</i>	†*62D	Porterville, Tulare Co., CA USA	BVH 127a
	†*65D	Camp Nelson, Tulare Co., CA USA	BVH 127b
	†*93C	Miami, Gila Co., AZ USA	BVH 93
	†*120C	American Canyon, San Luis Obispo Co., CA USA	BVH 120
	†*128A	Oakhurst, Madera Co., CA USA	BVH 128
<i>C. betuloides</i> var. <i>blancheae</i>	*121A	Parisima Hills, Santa Barbara Co., CA USA	BVH 121
	*123F	Santa Monica Mts., Los Angeles Co., USA	BVH 123
	†*123G	Santa Monica Mts., Los Angeles Co., USA	BVH 123
<i>C. minutiflorus</i>	†*57B	Del Mar, San Diego Co., CA USA	JH 22311
<i>fothergilloides</i> complex			
<i>C. traskiae</i>	†*126E	Santa Catalina Island, Los Angeles Co., CA USA	Thorne 36737
<i>C. fothergilloides</i> var. <i>fothergilloides</i>	†*67B	Los Lirios, Coahuila, MX	BVH and JH 22413
	*84G	San Roberto Junction, Nuevo Leon, MX	BVH and JH 22482
<i>C. fothergilloides</i> var. <i>medranoanus</i>	*77B	Valle Hermosa, Tamaulipas, MX	BVH and JH 22444
	†*82A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22456b
<i>C. pringlei</i>	C. pri	Matatlan, Oaxaca, MX	Stevens 1318
<i>C. macrophyllus</i>	†*26	Durango, MX	Crozier n/a
	†*69A	Sierra de la Viga, Coahuila, MX	BVH and JH 22415
	†*69B	Sierra de la Viga, Coahuila, MX	BVH and JH 22415
New taxa			
<i>C. mexicanus</i>	*134A	San Antonio Pena Nevada, Nuevo Leon, MX	JH 19295
<i>C. rzedowskii</i>	†*76A	Miquihuana, Tamaulipas, MX	BVH and JH 22434
	†*76E	Miquihuana, Tamaulipas, MX	BVH and JH 22434
	*81A	Valle Hermosa, Tamaulipas, MX	BVH and JH 22460
<i>ledifolius</i> complex			
<i>C. ledifolius</i> var. <i>intermontanus</i>	†*45C	Mantua, Box Elder Co., UT USA	BVH 45

	†*98A	Verni, Sierra Co., CA USA	BVH 98
	†*101A	Toiyabe Range, Lander Co., NV USA	BVH 101
	†*102B	Ruby Mts., Elko Co., NV USA	BVH 102
	†*108C	Spring Mts., Clark Co., NV USA	BVH 108
	†*111A	West Walker River, Mono Co., CA USA	BVH 111
<i>C. ledifolius</i> var. <i>ledifolius</i>	†*41B	Emery, Emery Co., UT USA	BVH 41
	†*104	Ely, White Pine Co., NV USA	BVH 104
<i>C. intricatus</i>	†*25	Escalante, Garfield Co., UT USA	BVH 25
	†*50D	Vernal, Uintah Co., UT USA	BVH 50
	†*103A	Ely, White Pine Co., NV USA	BVH 103
<i>Purshia tridentata</i>	†*36B	Casto Canyon, Garfield Co., UT USA	BVH 36
<i>Purshia plicata</i>	†*74D	Interbide, Coahuila, MX	Nesom 6198
<i>Chamebatia foliolosa</i>	129	Wakalu Hep Yo Campground, Calaveras Co., CA USA	BVH 129

CpDNA Amplification and sequencing

I amplified and sequenced two non-coding cpDNA markers for the DNA accessions in Table 4.1. The first was the entire inter-genic spacer between the *trnS* and *trnG* coding regions using the forward primer (5'-GCCGCTTTAGTCCACTCAGC-3') and the reverse primer (5'-GAACGAATCACACTTTTACCAC-3') (Hamilton 1999). The other non-coding region was the entire inter-genic spacer between the *rpl20* and *rps12* coding regions using the forward primer (5'-TTTGTCTACGTCTCCGAGC-3') and the reverse primer (5'-GTCGAGGAACATGTACTAGG-3') (Hamilton 1999). A negative control, which lacked template DNA, was included in all PCR runs. Fifty microliter PCRs were performed using 30mM Tricine, pH 8.4, 2mM MgCl₂, 50 mM KCl, 5% acetamide, 100 uM of each dNTP, 1.0 unit of Taq polymerase, and 30 nM of each amplification primer. Twenty nanograms of template DNA was

added, and the reactions were overlaid with mineral oil and run on a MJ Research Thermal Cycler programmed for a hot start (95° C, 5 min; 74° C, 7 min) and 30 cycles of 94° C for 30 sec, 50° C for 45 sec, and 72° C for 1 min. A final extension of 72° C for 7 min was also included. Three microliters of the reaction products were run on a 1.5 % agarose gel in 0.5X TBE buffer. PCR products were visualized with ethidium bromide under UV light, and purified using Sephadex columns (Princeton Separations, Inc.). DNA sequencing used standard Big Dye reagents (Applied Biosystems) and was performed on a MJ BaseStation (MJ Research). Each cpDNA region was proofread and then aligned using Clustal W in MegAlign (DNASTAR, 1998), followed by adjustment by eye, yielding two aligned datasets (Appendix E.).

Phylogenetic analysis

Phylogenetic analyses were performed with maximum parsimony (MP) optimality criterion. I treated the characters as unordered. The analyses were performed in PAUP version 4.0b10 (Swofford 1999). I used a heuristic search strategy using accelerated transformation (ACCTRAN), with 10,000-sequence addition replicates and TBR branch swapping. Support for particular branches was evaluated using 100 bootstrap replicates (Felsenstein 1985), using the simple addition of sequences.

I performed this search strategy on each of the aligned cpDNA regions and a combined dataset (see below). Indel events of 2 bp or more were coded as

presence/absence and added to the aligned sequence dataset. I also performed this search strategy on the 52 taxa common between the AFLP and cpDNA datasets, and on the 44 taxa common between the cpDNA and ETS datasets (Table 4.1). A description of the AFLP and ETS datasets can be found in Chapter 3. To compare the cpDNA dataset to the ETS dataset, I had to take into account the two cases of paraphyletic ETS clones from the individuals *Cercocarpus ledifolius* var. *intermontanus* 111A (clone 4 and 5) and *C. betuloides* var. *betuloides* 62DA (clones 2 and 3). Therefore, I constructed 4 ETS datasets which differed in the clones which represented the individuals 111A and 62D in all possible combinations (Table 4.2).

Table 4.2: A description of the clones included in each of the ETS datasets tested against the cpDNA dataset. 111A refers to the individual *Cercocarpus ledifolius* var. *intermontanus* collection 111A (table 1). 62D refers to *C. betuloides* var. *betuloides* collection 62D (table 1).

	111A clone 4	111A clone 5	62D clone 2	62D clone 3
ETS dataset 1	included	not included	included	not included
ETS dataset 2	included	not included	not included	included
ETS dataset 3	not included	included	included	not included
ETS dataset 4	not included	included	not included	included

Tests for incongruence

I tested for incongruence between the *trnS-trnG* inter-genic spacer region and the *rpl20* and *rps12* inter-genic spacer region using the partition homogeneity test (Farris et al. 1994, 1995) as implemented in PAUP (Swofford 1999). This

test first repartitions the characters found in the two original datasets into two temporary data subsets, finding the shortest tree length for each and adds them together. This was done for 100 replicates. Then, the sum of the shortest tree lengths for the original datasets is compared to the distribution of the repartitioned tree length sums. If there is a low probability of finding shorter tree length sums from the repartitioned data than from the original data, the original datasets are interpreted as incongruent. The decision to combine the cpDNA non-coding regions into one dataset is based on the results of this test.

To assess the incongruence between the cpDNA dataset and the ETS datasets, and the incongruence between the cpDNA dataset and the AFLP dataset, I conducted the "compare-2" test (Faith 1991), the Templeton test (Templeton 1983), and the Kishino-Hasegawa test as implemented in PAUP (Swofford 1999). The methods for these tests are the same as in Chapter 3. The constraint trees for all three tests were the MP bootstrap consensus trees, which only included branches with bootstrap support over 50%.

RESULTS

Full cpDNA phylogenetic analysis

The length of the aligned sequence dataset for the *trnS-trnG* spacer was 580 bp, and for the *rps12-rpl20* spacer was 757 bp. I recorded 6 indel events in the *trnS-trnG* spacer and recorded 8 indel events in the *rps12-rpl20* spacer that

were greater than 2 bp. The results of the partition homogeneity test for the congruence of the *trnS-trnG* spacer and the *rps12-rpl20* spacer yielded a non-significant result ($p=0.980$) (Table 4.3). Therefore, I combined the two datasets and continued with a MP heuristic search. For the full cpDNA dataset of 54 individuals, our analysis recovered 6369 equally parsimonious trees of length 251 (Figure 4.1). The tree statistics for this search can be seen in Table 4.4.

Table 4.3: Results of the partition homogeneity test for the *trnS-trnG* inter-genic spacer versus the *rpl20-rps12* inter-genic spacer. The asterisk denotes the sum of the tree lengths for the original, unpermuted partition.

sum of tree lengths	number of replicates
223	1
224	2
225	6
226	17
227	29
228	15
*229	18
230	2

Table 4.4: A comparison of the tree statistics for the multiple MP heuristic searches conducted on all three datasets. CpDNA refers to the complete cpDNA dataset, including all 54 taxa. CpDNA vs. AFLP includes the 52 taxa in common between datasets. CpDNA vs. ETS includes the 44 taxa in common between datasets. ETS 1 – 4 identities refer to Table 4.2.

	CpDNA	CPDNA vs AFLP	AFLP	CPDNA vs ETS	ETS 1	ETS 2	ETS 3	ETS 4
# of ind.	54	52	52	44	44	44	44	44
# of equally parsimonious trees	6369	84	126	9342	1598	1520	2561	2584
tree length	251	210	827	168	395	400	404	409
# of characters	1352	1352	376	1352	977	977	977	977
# of parsimony informative characters	72	69	76	61	133	133	134	134
CI	0.847	0.838	0.314	0.851	0.767	0.769	0.757	0.768
RI	0.824	0.827	0.503	0.836	0.808	0.799	0.794	0.797

The MP phylogeny produced from the full cpDNA dataset lacks overall resolution, and supports four main clades (Figure 4.1). *Cercocarpus* is found to be monophyletic with good bootstrap support. Within *Cercocarpus*, clade A is weakly supported (bootstrap score below 50) and contained members of all four species complexes defined by Lis (1992). Clade B was weakly supported (bootstrap score of 62), in an unresolved position with clade A. Clade B contained all members of the *Cercocarpus betuloides* complex, except *C. betuloides* var. *betuloides* (93C) and *C. minutiflorus* (57B), which were present in clade A. Two individuals of *Cercocarpus ledifolius* var. *intermontanus*

(collections 101A and 111A) were found nested within clade B. One individual of *Cercocarpus ledifolius* var. *intermontanus* (98A) was found in an unresolved position with clade B. Clades C and D were found in an unresolved position sister to clades A and B with moderate bootstrap support (score of 82). Clades C and D are shown as unresolved with respect to each other, yet 98% of the trees recovered in the MP search united them as sister to one another. Clades C and D are made up almost exclusively of members of the *Cercocarpus fothersgilloides* complex and *C. rzedowskii*. The one exception, *C. montanus* var. *paucidentatus* (75A), was positioned in an unresolved position within clade C.

CpDNA vs. AFLP phylogenetic analysis

We pruned two individuals from the full cpDNA dataset for a total of 52 individuals to compare to the AFLP dataset (Table 4.1). Our MP heuristic search using the reduced cpDNA dataset resulted in 84 most parsimonious trees of length 210 with similar topology to the full cpDNA dataset (Figure 4.2). The MP heuristic search using the same 52 accessions from the AFLP dataset resulted in 126 most parsimonious trees of length 827 (Figure 4.2) (Table 4.4).

Although MP trees produced from the AFLP and cpDNA datasets do not support the same relationships for the 52 individuals in common between them, I found some well supported congruent topologies. First, both the cpDNA and AFLP trees support a *Cercocarpus fothersgilloides* clade with high bootstrap support. Also, both the cpDNA and AFLP datasets support a close association

between the *Cercocarpus fothergilloides* complex and *C. rzedowskii*. Finally, both trees support *Cercocarpus traskiae* as being more closely related to the *C. betuloides* complex than to the *C. fothergilloides* complex.

Due to the lack of resolution in the cpDNA tree, we are unable to make many claims about relationships between species. The tree based on the AFLP dataset supports the separation of the *Cercocarpus montanus* complex, the *C. ledifolius* complex, although these groupings have bootstrap support values lower than 50. These groups were not supported in the cpDNA tree, where the *Cercocarpus montanus* and *C. ledifolius* complexes were grouped together in clade A and B. The individual of *Cercocarpus montanus* var. *paucidentatus* (75A) is the only exception for the monophyly of the *C. montanus* complex in the AFLP tree. This is the same individual that was divergent in the cpDNA analysis as discussed above. The *Cercocarpus fothergilloides* complex is the only clade that has high bootstrap support in the AFLP tree (Figure 4.2). The *Cercocarpus fothergilloides* complex did group together in cpDNA tree in clades C and D. Members of the *Cercocarpus betuloides* complex are found in clades A and B in the cpDNA tree, yet are not grouped together in the AFLP tree, where they are found scattered and unresolved.

CpDNA vs ETS phylogenetic analysis

I pruned 10 individuals from the full cpDNA dataset for a total of 44 individuals to compare to the ETS datasets (Table 4.1). The tree statistics for the

reduced cpDNA dataset and the four ETS datasets are illustrated in Table 4.4. The cpDNA phylogeny has the same structure as the previous two cpDNA phylogenies (Figure 4.3). The four ETS datasets differed in their topology due to the inclusion and exclusion of the different clones for individual 111A and 62D (Table 4.2). The MP strict consensus of the four ETS datasets is in Figure 4.3.

The ETS and cpDNA trees are similar in the placement of the *Cercocarpus montanus* complex as derived, although clade A is unresolved. Both trees also support the unity of the *Cercocarpus betuloides* complex (clades D and E in the ETS tree, clade B in the cpDNA tree). Overall, the cpDNA and ETS datasets do not support the same relationships for the 44 individuals. The ETS strict consensus tree contains 6 recognizable clades, A through F (Chapter 3). The most notable difference between these two trees is the placement of individuals of the *Cercocarpus fothersgilloides* complex. The cpDNA tree supports a single clade with one exception (*Cercocarpus fothersgilloides* var. *fothersgilloides* 84G), yet members of the *C. fothersgilloides* complex are present in three clades in the ETS tree (Clades A, B, and F)

Incongruence tests

The results of the compare-2, the Templeton, and the Kishino-Hasegawa tests can be seen in Table 4.5. For the compare-2 test, 100 permutations of each dataset were tested on each constraint tree. I found significant results with every dataset, where the difference in fit for the dataset on its own tree versus the

constraint tree was significantly better than the difference with permuted data. The Templeton and Kishino-Hasegawa tests yielded highly significant results for each dataset, signaling significant incongruence between all datasets tested.

Table 4.5: Results for the tests of congruence between the cpDNA trees and the trees produced from ETS and AFLP datasets.

	COMPARE-2	KISHINO-HASEGAWA			TEMPLETON		
	p-value	diff. in tree length	s.d.	t	p-value	N	p-value
cpDNA vs AFLP							
cpDNA dataset	p=0.01	66	14.31537	4.6104	p=0.0001	27	p=0.0001
AFLP dataset	p=0.01	319	38.5267	8.28	p=0.0001	109	p=0.0001
cpDNA vs ETS 1							
cpDNA dataset	p=0.01	65	18.71187	3.4737	p=0.0005	18	p=0.0002
ETS 1 dataset	p=0.01	296	40.12442	7.3771	p=0.0001	85	p=0.0001
cpDNA vs ETS 2							
cpDNA dataset	p=0.01	65	18.71187	3.4737	p=0.0005	18	p=0.0002
ETS 2 dataset	p=0.01	348	43.41185	8.0162	p=0.0001	103	p=0.0001
cpDNA vs ETS 3							
cpDNA dataset	p=0.01	61	18.01354	3.3863	p=0.0007	19	p=0.0002
ETS 3 dataset	p=0.01	340	42.47163	8.0053	p=0.0001	86	p=0.0001
cpDNA vs ETS 4							
cpDNA dataset	p=0.01	66	19.31461	3.4171	p=0.0007	18	p=0.0002
ETS 4 dataset	p=0.01	301	40.86533	7.3657	p=0.0001	86	p=0.0001

DISCUSSION

My goal was to develop an independent molecular phylogeny for *Cercocarpus* to compare with the two incongruent nuclear phylogenies presented in Chapter 3. The *trnS-trnG* spacer and the *rps12-rpl20* had congruent phylogenetic signal. The maximum parsimony phylogeny produced from the combined dataset yielded trees with 4 major clades, A through D. These trees strongly supported the monophyly of *Cercocarpus* using both *Purshia* and

Chamaebatia as outgroups. There was little resolution within the clades due to a lack of phylogenetically informative variation, and I was unable to resolve the relationships between the majority of *Cercocarpus montanus* and *C. ledifolius* samples in the dataset present in clade A. All but two members of the *Cercocarpus betuloides* complex were found in clade B, including *C. traskiae*. Clades C and D included mostly members of the *Cercocarpus fothergilloides* complex and the two newly described taxa, *C. rzedowskii* and *C. mexicana*. The chloroplast phylogeny was significantly incongruent to both the ETS and AFLP phylogenies for comparable individuals. Both the chloroplast and AFLP phylogenies suggest a well supported *Cercocarpus fothergilloides* complex and the close association between the *C. fothergilloides* complex and the two newly described taxa of *C. rzedowskii* and *C. mexicanus*. The cpDNA and ETS tree both support the *Cercocarpus betuloides* complex as unique from the *C. montanus* complex, with a few exceptions.

Taxonomic implications and testing previous hypotheses

I found all but two members of the *Cercocarpus betuloides* complex present in clade B, separate from the majority of *C. montanus* samples in clade A. This result was also found in the ETS and AFLP trees, and lends further support to the decision by Lis (1993) to recognize *C. betuloides* as a separate entity from *C. montanus*.

I also found *Cercocarpus traskiae* grouping with members of the *C. betuloides* complex. This result is consistent with both the AFLP and the ETS datasets. *Cercocarpus traskiae* had been hypothesized to be a part of the *C. fothergilloides* complex based on leaf characteristics (Lis 1991, Henrickson ined.). It appears based on all three datasets that *C. traskiae* belongs allied to the *C. betuloides* complex and not the *C. fothergilloides* complex.

The three individuals of *Cercocarpus rzedowskii* were found in the weakly supported clade C, along with one individual of *C. montanus* var. *paucidentatus* (75A) and *C. mexicanus* (134A). The relationship of clade C to clade D in the full cpDNA dataset, the *Cercocarpus fothergilloides* clade, is unresolved in the MP strict consensus tree, yet most (98%) of the equally parsimonious reconstructions supported a sister relationship. This implies that based on the cpDNA dataset, *Cercocarpus rzedowskii* and *C. mexicanus* both have affinities to the *C. fothergilloides* complex. This was an interesting result. Henrickson (1987) suggested in his description of these taxa that *C. rzedowskii* had similarities to *C. ledifolius* based on the revolute, entire margin and overall leaf shape. The ETS dataset strongly supported a close association between *Cercocarpus rzedowskii* and *C. montanus*, yet the AFLP suggested it was sister to the *C. fothergilloides* complex. Here I have more evidence for the inclusion of *Cercocarpus rzedowskii* into the *C. fothergilloides* complex. The similarities of leaf characters appear to be examples of convergent evolution.

In the description of *Cercocarpus mexicanus*, Henrickson (1987) hypothesized it was similar to the *C. montanus* complex, due to the serrate leaf margin and basic leaf shape displayed by this taxon. The AFLP dataset grouped the one individual of *Cercocarpus mexicanus* in this study deeply nested in the *C. betuloides* complex cluster. In the cpDNA tree, we see *Cercocarpus mexicanus* with *C. rzedowskii* in the *C. fothergilloides* cluster. More sampling of this taxon is required for a firm hypothesis on its affinities in *Cercocarpus*. Unfortunately, the one sample of *C. mexicanus* used in this study did not have an ETS type 1 to be included in the ETS analysis.

Stutz (1974) hypothesized that the origin of *Cercocarpus montanus*, with large, broad deciduous leaves arose as a peripheral isolate of *C. ledifolius* that became adapted to colder, higher altitudes in the Rocky Mountains. Stutz (1974) also hypothesized that *Cercocarpus intricatus*, with small, revolute leaves arose from adaptations by an ancestral *C. ledifolius* lineage that became adapted to drier, hotter conditions. I found limited support for this theory (Figure 4.1) with both *Cercocarpus montanus* and *C. intricatus* arising from a paraphyletic *C. ledifolius* clade. Although this is an interesting result, the lack of support for the cpDNA tree topology requires caution. Also, we have no support for these hypotheses with both the ETS and AFLP tree topologies.

Implications of incongruence and evidence for hybridization

I found significant incongruences for all comparisons between the cpDNA tree and the ETS and AFLP trees. This result might be explained as merely an artifact, i.e., a product of a lack of support for branches. As can be seen in Figure 4.1, the cpDNA MP phylogeny is unresolved, and contains few branches with bootstrap support greater than 50%. I attempted to reduce the possibility of this explanation by using the bootstrap consensus trees that included only branches with at least 50% bootstrap support as the constraint trees in the incongruence tests. The significant result using these bootstrap constraint trees implies that the incongruence goes beyond simple lack of support for branches, and suggests that the markers used in creating the phylogenetic trees had different evolutionary histories in *Cercocarpus* and may have been subject to horizontal transmission between distinct lineages. Hybridization has long been discussed as a probable occurrence in the genus *Cercocarpus* (Blauer et al. 1975). Schneider (1905b) invoked hybridization to explain the intergradations in leaf morphology he saw in some populations of *Cercocarpus ledifolius*, although the individuals with intermediate morphology might also be explained by either plastic responses to water availability or selection for ecotypes on a rainfall gradient. The only clear genetic evidence for hybridization in *Cercocarpus* comes from Santa Catalina Island, where Rieseberg (1989, 1994) documented introgression between *C. betuloides* and *C. traskiae* using both allozymes and RAPD markers.

The use of hybridization as the cause for the incongruence between the chloroplast and AFLP or ETS trees can only be used if geographic proximity of the taxa involved would allow it, although we must also consider possible changes in range limits for hybridizing taxa. I identified four individuals that are located in clades that I would not expect based on their morphology. These four taxa are *Cercocarpus montanus* var. *paucidentatus* (75A) placed in clade C sister to *C. rzedowskii* and *C. macrophyllus*, two individuals of *C. ledifolius* var. *intermontanus* (101A and 111A) placed in clade B with individuals of the *betuloides* complex, and *C. betuloides* var. *betuloides* (93C) in clade A, with individuals identified as *C. montanus* and *C. ledifolius*. Arrows in Figure 4.2 and 4.3 identify these taxa. As might be expected, we find the individuals which are “out of place” in the chloroplast tree are found in geographical areas of overlap between ranges of species complexes. *Cercocarpus montanus* var. *paucidentatus* (75A) was collected in southwestern Coahuila, near the species range of *C. rzedowskii* and *C. mexicanus*, whose members it was grouped with in the chloroplast tree. The two individuals of *Cercocarpus ledifolius* var. *intermontanus* (101A and 111A) that were placed in the same clade as members of the *C. betuloides* complex, were collected in the western end of the *C. ledifolius* distribution near the eastern edge of the range of *C. betuloides*. Lastly, the individual of the *Cercocarpus betuloides* complex, *C. betuloides* var. *betuloides* (93C), that was placed in clade A was collected in central Arizona,

well within the species range of the *C. montanus* complex. The anomalous placement of these individuals in the chloroplast tree based on their position in the AFLP tree, coupled with the geographic position of the collection sites supports hybridization and chloroplast capture as the explanation for the incongruences.

Molecular systematics of *Cercocarpus* illuminates the problems that may arise from attempting phylogenetic reconstruction in very closely related groups of plants. First, it can be difficult to obtain phylogenetically informative variation in DNA sequence regions, i.e., the chloroplast sequences presented here and the ITS region attempted in Chapter 3, due to a lack of sufficient time for substitutions to accumulate between taxa. Second, if you do indeed find DNA sequence variation, (for example the ETS region) issues of coalescent times can become paramount. If the coalescent time for the sequence variation obtained is older than the taxa trying to be reconstructed, many spurious relationships can be reconstructed (Chapter 3).

Solving the phylogenetic problems that accompany closely related groups of plants means finding a marker that is evolving at an appropriate rate, where homology is unambiguous, and is not subject to deep coalescence (Soltis and Soltis 1998). It may be that such a marker can be identified in *Cercocarpus*. Another possibility is that a robust phylogeny for *Cercocarpus* will never be developed because the history of the group contains an excess of reticulation, where identified taxa do not have bifurcating evolutionary histories. Plant groups

with such a history might be better understood if relationships between species were viewed as networks, rather than phylogenies. With the inclusion of more data, I hope to uncover further relationships in *Cercocarpus* with the full understanding that the evolutionary history of taxa in this group may not conform to traditional, bifurcating tree topologies.

This work has laid a foundation for my work present and future work on the coevolutionary relationships between *Cercocarpus* and *Frankia*. Only by understanding the evolutionary history of both members in an ecological association can we study coevolution in a meaningful way. I hope to use what I have learned with *Cercocarpus* and apply it to other genera in the Rosaceae that associate with *Frankia*, expanding the study of coevolution to include all members of nitrogen-fixing members of the Rosaceae.

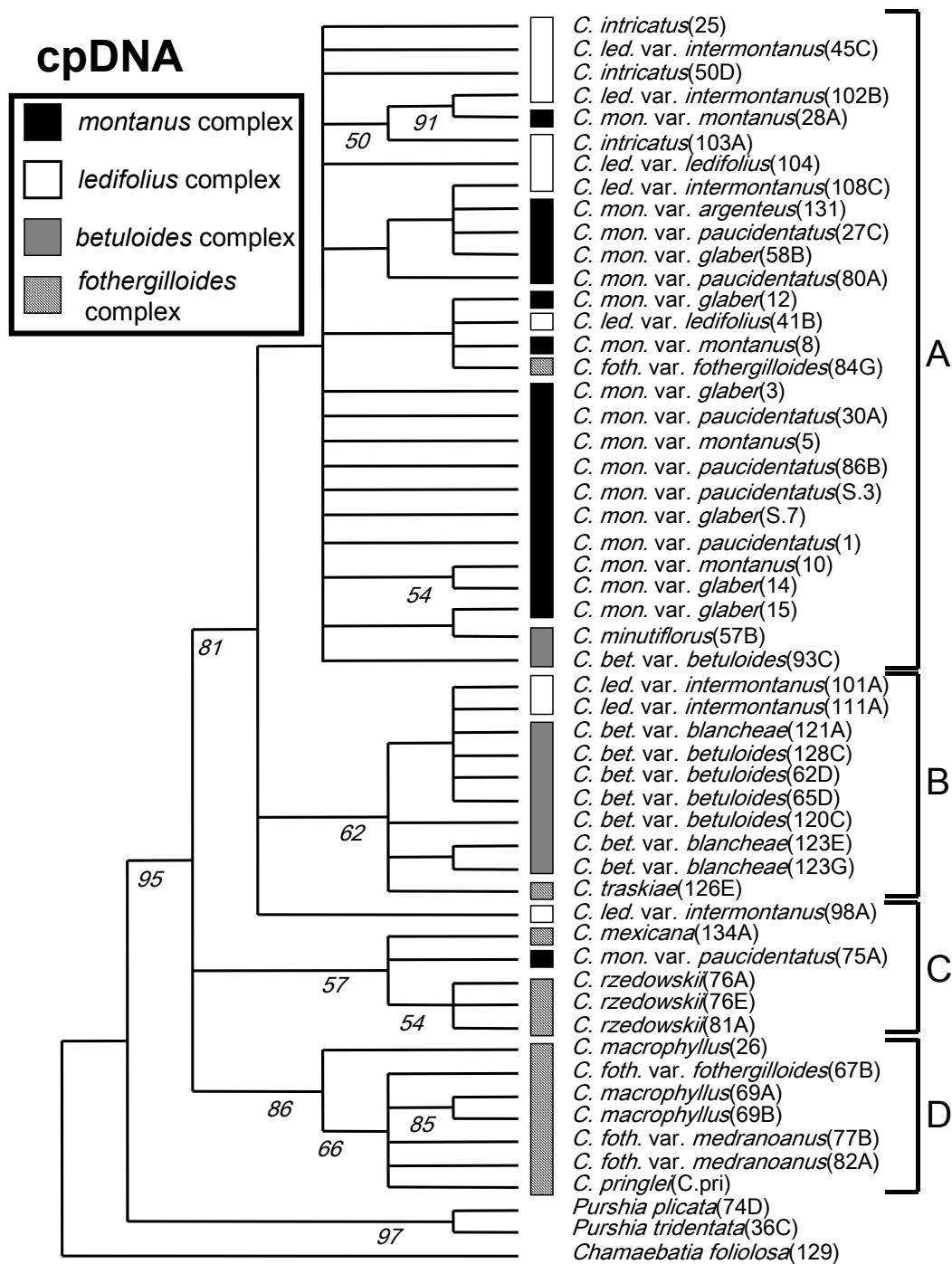


Figure 4.1: A strict consensus cladogram based on the combined cpDNA regions. It is composed of 6369 most parsimonious trees of length 251 steps. See table 4.4 for further tree statistics. Bars next to the taxon name designate species complex membership. Clades A-D are shown by the brackets.

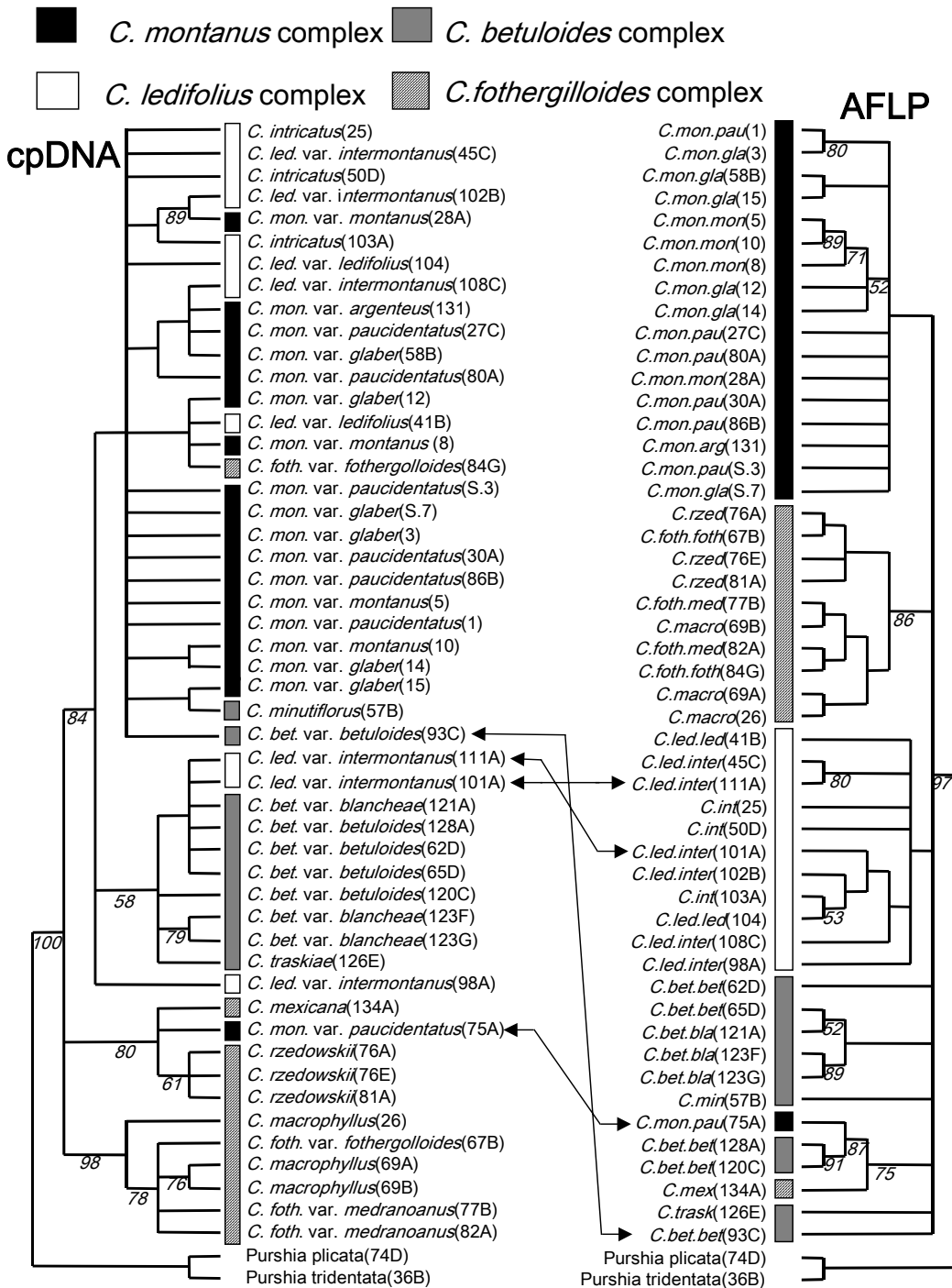


Figure 4.2: Comparison of the maximum parsimony strict consensus trees based on cpDNA sequences and AFLP data for 52 taxa. See Table 4.4 for further tree statistics. The cpDNA tree is the strict consensus of 84 most parsimonious trees of length 210 steps. The AFLP tree shown is the strict consensus of 126 most parsimonious trees of length 827 steps. Bars next to the taxon name designate species complex membership. Arrows refer to taxa in discussion.

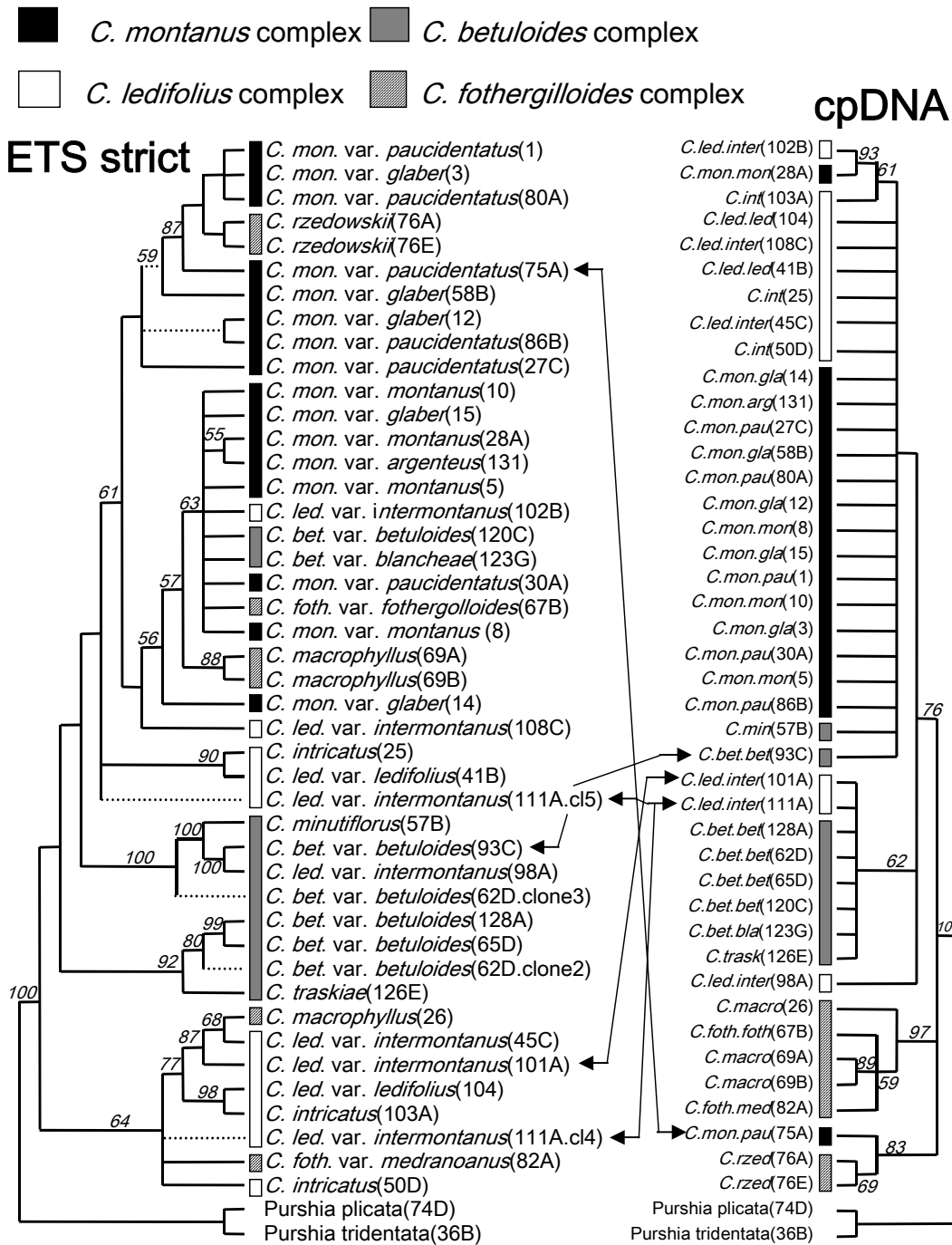


Figure 4.3: A comparison of maximum parsimony strict consensus trees based on cpDNA sequences and ETS sequence data for 44 taxa. See table 4.4 for tree statistics. The cpDNA tree shown is a strict consensus of 9342 most parsimonious trees of 168 steps. The ETS tree is the strict consensus of 2367 most parsimonious trees of 419 steps. Bars next to the taxon name designate species complex membership. Arrows refer to taxa in discussion.

Appendix A. ETS alignable DNA sequence data

This dataset contains the aligned sequences for the three alignable regions (5' alignable, 3' alignable, and the α and β regions) from the ETS types used in Chapter 2. The sequences for the three regions have been combined. The 5' alignable region is represented from positions 1-174. The 3' alignable region is represented from positions 175-382. The α and β regions are represented from positions 383-512. ID codes for the sequences correspond to those found in Table 1.1. ETS type 1, type 2, type 3, and type 4 have been separated and identified. Gaps in the sequence are indicated by a dash (-). Uncertainties are denoted by standard IUPAC/IUB codes as follows: N=unknown; R=A/G; Y=C/T; M=A/C; K=G/T; S=C/G; W=A/T.

Bases 1-60

5' Common region →

ETS type 1

1	TCT-C-AATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
10	TST-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
14	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
15	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
12	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
17	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
25	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
26	TCT-CCAATA	TT---CTTAC	GTGTTCTTGA	GTGTTTCATCT	ACAGTCCTTG	CA-TATTT-G
27C.1	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
27C.10	-CT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
28A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
3	TCT-C-AATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
41B	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
45	TCT-CCAATA	TT---CTTAC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
5	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
101A	TCT-CCAATA	TT---CTTAC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
102B	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
104	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
108C	-----	-----AC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCA	CACTACTTTG
111A.4	-----	-----C	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
111A.5	-----	-----C	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
120C	-----	-----TG	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
123G	-----	-----	-----	-----	-----	-----
131	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
30A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
57B	-----	-----GC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
58B	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
62D.2	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
62D.3	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
65D.7	TCT-GCA-TA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAATTCTTG	CA-TATTT-G
65D.3	-CT-GCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAATTCTTG	CA-TATTT-G
67B	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
69A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
69B	-----	-----C	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
76A	TC---CAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
76E	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
78A.10	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
78A.5	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
8	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
80A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
82A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
85A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
86B	-----	-----C	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
90D	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
93C.2	TCC----ATA	TT---CTTGC	GCGTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
93C.3	TCT-CCAACA	TT---CTTGC	GCGTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
98A	-----	-----GC	GCATTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
126E	-----	-----	-----	-TGTTTCATCT	ACAATTCTTG	CA-TATTT-G
103A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	GCAGTTCTTG	CA-TATTT-G
50D	-----	-----	-----TTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTTAG
68	----CC-ATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
75A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G

Bases 1-60

ETS type 2

121C	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
18	TCT-CTAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
19	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
20	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
21	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
122A	TCT-CCAACA	TTG--CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
126.29	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
26	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
52B	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
57B	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
65C.3	TCT-CCCACT	TT---CTTAC	GTGTTCTTGA	GTGTTTGTCT	ACAGTTCTCA	CA-TATTT-G
65C.4	TC---CCACT	TT---CTTAC	GTGTTCTTGA	GTGTTTGTCT	ACAGTTCTCA	CA-TATTT-G
93C	TCT-CCAACA	TCT--CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
118A	TCTCCA-CA	TCTCTCTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTAA	CA-TATTT-G
126.BC1	TCTACCAACA	TCTCTCTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
126E.1	TCT-CCAAGA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
126E.3	TCT-CCAAAA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
126E.4	TCT-CCAAAA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
126A	TCT-CCAAAA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
61	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G

ETS type 3

121B	TCT-CCAAAA	TT---CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
66C	TCT-CCAACA	TT---ATTTT	ATGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
66D	TCT-CCAACA	TT---CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
67A	-----	-----C	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
69B	-----	-----C	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
71A	TCT-CCAACA	TT---CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
77B	---CCAACA	TT---CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
82A	TCT-CCAACA	TT---CTTTC	ATGTTCTTAA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
83B	-----	-----CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
86B.24	TCT-CCAACA	TT---CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
86B.29	TCT-CCAACA	TT---CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
86B.5	-----	-----CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
87B	-CT-CCAACA	T--CTCTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G
90D	TCT-CCAACA	T----CTTTC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTCG	CA-TATTT-G

ETS type 4

118A	TCT-CCCACA	TT---CTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTAA	CA-TATTT-G
121B	TCT-CCCACA	TT---CTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTAA	CA-TATTT-G
123F	-----	TT---CTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTCA	CA-TATTT-G
126.29.1	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
126.29.3	TCT-CCAACA	TT---CTTAC	GTGATCTTGA	GTGTTTATCT	GCAGTTCTCA	CA-TATTT-G
63B	TCT-CCCACA	TT---CTTAC	GTGTTCTTGA	GTGTTTGTCT	ACAGTTCTCA	CA-TATTT-G
81A	TNT-CCAGCA	TT---CGTGN	GTGNTCNTGA	GTGTTTCATCA	ACAGTTCTGT	CA-TATTT-A
76A	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
76E	TCT-CCAATA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
78B	TCT-CCAGCA	TT---CTTGC	GTGTTCTTGA	GTGTTTCATCA	ACAGTTCTTG	CA-TATTT-A
84A.1	TCA-CCCACA	TT---CTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTAA	CA-TATTT-G
84A.2	TCT-CCCACA	TT---CTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTAA	CA-TATTT-G
83B	TCT-CTAATA	TT--TCTTG	GTGTTCTTGA	GTGTTTCATCT	ACAGTTCTTG	CA-TATTT-G
87A	TCT-CCAACA	TT---CTTAC	GTGTTCTTGA	GTGTTTCATCT	GCAGTTCTCG	CA-TATTT-G
126.WB3	TCT-CCCACA	TT---CTTAC	GTGTTCTTGA	GTGTTTATCT	ACAGTTCTCA	CA-TATTT-G
Pur-34	-----	-----	-----	-----	-----	-----
Pur-132	-----	-----	-----	-----	-----	-----

Bases 61-120

ETS type 1

1	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
10	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
14	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
15	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
12	AGCTG-TCTA	ATCTACTT-T	GGCAGGCR-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
17	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
25	AGTTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTAACT	TAGCTTGTTT	TTGGAGGTGA
26	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
27C.1	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
27C.10	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
28A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
3	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
41B	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTAACT	TAGCTTGTTT	TTGGAGGTGA
45C	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
50	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
101A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
102B	AGCTG-GCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
104	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
108C	AGCTACTCTG	ATCTA-TT-T	GGCAGGCACT	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
111A.4	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGATT	TAGCTTGTTT	TTGGAGGTGA
111A.5	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGATT	TAGCTTGTTT	TTGGAGGTGA
120C	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
123G	-----	-----	-----	-----	-----	-----
131	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
30A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
57B	AGTTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTAGCTTACT	TAGCTTGTTT	TTGGAGGTGA
58B	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
62D.2	AGCTG-TCTA	ATCTACTT-T	GGCAAGCG-T	TTAGCTAACT	TAGCTTGTTT	TTGGAGGTGA
62D.3	AGCTG-TCTA	ATCTACTT-T	GGCAAGCG-T	TTAGCTAACT	TAGCTTGTTT	TTGGAGGTGA
65D.7	AGCTG-TCTA	ATTTACTT-T	GGCAGGCG-T	TTGGATGACT	TAGCTTGTTT	TTGGAGGTGA
65D.3	AGCTG-TCTA	ATTTACTT-T	GGCAGGCG-T	TTGGATGACT	TAGCTTGTTT	TTGGAGGTGA
67B	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
69A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
69B	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
76A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCATGTTT	TTGGAGGTGA
76E	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
78A.10	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCATGTTT	TTGGAGGTGA
78A.5	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCATGTTT	TTGGAGGTGA
8	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
80A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TARCTTGTTT	TTGGAGGTGA
82A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCA-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
85A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
86B	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
90D	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
93C.2	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTAGCTAACT	TAGCTTGTTT	TTGGAGGTGA
93C.3	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTAGCTAACT	TAGCTTGTTT	TTGGAGGTGA
98A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTAGCTAACT	TAGCTTGTTT	TTGGAGGTGA
126E	AGCTG-TCTA	ATTTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
103A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
50D	AGCTG-GCTG	ATCTACYTCT	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
68A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGCTTGTTT	TTGGAGGTGA
75A	AGCTG-TCTA	ATCTACTT-T	GGCAGGCG-T	TTGGCTGACT	TAGSTTGTTT	TTGGAGGTGA

Bases 61-120

ETS type 2

121C AGATA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TGGCTTGTTT TTGGAGGTGA
18 GGCTG-TCTG ATCTACTT-T GGCAGGCG-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
19 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
20 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
21 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
122A AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
126.29 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
26 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
52B AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
57B AGATA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
65C.3 AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
65C.4 AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
93C AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
118A AGATA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TGGCTTGTTT TTGGAGGTGA
126.BC1 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
126E.1 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
126E.3 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
126E.4 AGCTA-TCTG ATCTATT--- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
126A AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA
61 AGCTA-TCTG ATCTATTT-- AGCAGGCA-T TTGGCAGACT TAGCTTGTTT TTGGAGGTGA

ETS type 3

121B AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
66C AGCTG-CCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
66D AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
67A AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
69B AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
71A AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
77B AGCTG-CCTG ATCTACTT-T GGCTGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
82A AGCTG-TCTG ATCTCTTT-T GGCAGACA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
83B AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
86B.24 AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
86B.29 AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
86B.5 AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
87B AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
90D AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA

ETS type 4

118A AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TCGGCTGACT TAGCTTGTTT TTGGAGGTGA
121B AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TCGGCTGACT TAGCTTGTTT TTGGAGGTGT
123F AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
126.29.1AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
126.29.3----- GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
63B AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
81A AGCTG-TCTG ATATACTT-T GGCAGGCA-T TTGGCTGACT TAGCTAGTTT TTGGAGGTGA
76A AGCTG-TCTA ATCTACTT-T GGCAGGCG-T TTGGCTGACT TAGCTAGTTT TTGGAGGTGA
76E AGCTG-TCTA ATCTACTT-T GGCAGGCG-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
78B AGCTG-TCTG ATCTACTT-T GGCAGGCA-T TTGGCTGACT TAGCTAGTTT TTGGAGGTGA
84A.1 AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TCGGCTGACT TAGCTTGTTT TTGGAGGTGA
84A.2 AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TCGGCTGACT TAGCTTGTTT TTGGAGGTGA
83B GGCTG-TCTG ATCTACTT-T GGCAGGCG-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
87A AGCTG-TCTG ATCT-TCTTT GGCAGGTG-T TTGGCTGACT TAGCTTGTTT TTGGAGGTGA
126.WB3 AGCTA-TCTG ATCTATTT-- GGCAGGCA-T TCGGCTGACT TAGCTTGTTT TTGGAGGTGA
Pur-34 -----
Pur-132 -----

Bases 121-174**ETS type 1**

1	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
10	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAAT	CATCCCATGA	CAGT
14	TCAAGTCTAA	GGTCACACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
15	TCAAGTCTAA	GGTCAYACAG	GTAGAGGACT	TCAAAACAAT	CATCCCATGA	CAGT
12	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
17	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
25	TTAAGTCTAA	GGTCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
26	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
27C.1	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
27C.10	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCATAACAGT	CATCCCATGA	CAGT
28A	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
3	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
41B	TTAAGTCTAA	GGTCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
45C	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
5	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
101A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
102B	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
104	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
108C	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
111A.4	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
111A.5	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
120C	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAAT	CATCCCATGA	CAGT
123G	-----TCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
131	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
30A	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAAT	CATCCCATGA	CAGT
57B	TCAAATCTAA	GGTTGCACGG	GTAGAGGACT	TCAAAATAGT	TATCCCATGA	CAGT
58B	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
62D.2	TCAAGTCTAA	GATCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCATGA	CAGT
62D.3	TCAAGTCTAA	GGTCGCACGG	GTAGAGGACT	TCAAAATAGT	TATCCCATGA	CAGT
65D.7	TCAAGTCTAA	GATCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCATGA	CAGT
65D.3	TCAAGTCTAA	GATCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCATGA	CAGT
67B	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAATAAT	CATCCCATGA	CAGT
69A	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
69B	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
76A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
76E	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
78A.10	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	TATCCCATGA	CAGT
78A.5	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	TATCCCATGA	CAGT
8	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
80A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
82A	TCAAGTCTAA	GGTCATACAG	GTAGAGGACT	TCAAAACAAT	CATCCCATGA	CAGT
85A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
86B	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAATAGT	CATCCCATGA	CAGT
90D	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAATAGT	CATCCCATGA	CAGT
93C.2	TCAAATCTAA	GGTCGCACGG	GTAGAGGACT	TCAAAATAGT	TATCCCATGA	CAGT
93C.3	TCAAATCTAA	GGTCGCACGG	GTAGAGGACT	TCAAAATAGT	TATCCCATGA	CAGT
98A	TCAAATCTAA	GGTCGCACGG	GTAGAGGACT	TCAAAATAGT	TATCCCATGA	CAGT
126E	TCAAGTCTAA	GATCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
103A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
50D	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
68A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT
75A	TCAAGTCTAA	GGTCGCACAG	GTAGAGGACT	TCAAAACAGT	CATCCCATGA	CAGT

Bases 121-174

ETS type 2

121C	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
18	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAGCAGT	CATCCCGTGA	CACT
19	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
20	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
21	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
122A	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
126.29	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
26	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
52B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
57B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
65C.3	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
65C.4	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
93C	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACGGT	CATCCCGTGA	CACT
118A	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126.BC1	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126E.1	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126E.3	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126E.4	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126A	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
61	TCAAGTTTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT

ETS type 3

121B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCGTGA	CACT
66C	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
66D	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
67A	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
69B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCGTGA	CACT
71A	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCGTGA	CACT
77B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
82A	TCAAGTCTAA	GGCTGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGCGA	CACT
83B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TCAAAATAGT	CATCCCGTGA	CACT
86B.24	TCAAGTCTAA	GGCCACACGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
86B.29	TCAAGTCTAA	GGCCACACGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
86B.5	TCAAGTCTAA	GGCCACACGG	GTAGAGGACT	TCAAAACAGT	TATCCCGTGA	CACT
87B	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	TATCCCATGA	CACT
90D	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	TATCCCATGA	CACT

ETS type 4

118A	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
121B	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
123F	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126.29.1	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
126.29.3	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
63B	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
81A	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	--AAAACAGT	CATCCCTTGA	CACT
76A	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TGAAAACAGT	CATCCCTTGA	CACT
76E	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	TGAAAACAGT	CATCCCTTGA	CACT
78B	TCAAGTCTAA	GGCCGCATGG	GTAGAGGACT	T--AAACAGT	CATCCCTTGA	CACT
84A.1	TTAAGTCTGA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
84A.2	TCAAGTCTGA	GGCCGCACGG	GTAGAGGACT	TCAAAACAGT	CATCCCGTGA	CACT
83B	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAGCAGT	CATCCCGTGA	CAGT
87A	TCAAGTCTA-	GGCCGCACGA	GTAGAGGACT	TCAAAACAAT	TATCCTGTAA	CAGT
126.WB3	TCAAGTCTAA	GGCCGCACGG	GTAGAGGACT	TCAAAACAAT	CATCCCGTGA	CACT
Pur-34	-----	-----	-----	-----	-----	----
Pur-132	-----	-----	-----	-----	-----	----

Bases 175-234

3' common region→

ETS type 1

1	ATGACCCTCA	CTCTCRGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
10	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
14	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
15	ATGACCCTCA	CTCTCGGGTR	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
12	ATGACCCTCA	STCTCGGGTG	CTTGTGAAGG	TGCAATGGGT	GCTATTCGAT	GTCGTATATG
17	ATGACCCTCA	CTCTCGGGCA	MTTGTGAAGG	TGCATTGGKT	GCTACTCGAT	MTCTTCTATG
25	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
26	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
27C.1	ATGACCCTTA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTTGAT	CTCTTCTATG
27C.10	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
28A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
3	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GGTAATCGGT	CTCTTCTATG
41B	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTGTATG
45C	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
5	ATGACCCTCA	CTGTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
101A	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
102B	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
104	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
108C	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
111A.4	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATA
111A.5	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCATATG
120C	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
123G	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GGTACTCGAT	CTTTTTTATG
131	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
30A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
57B	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
58B	ATGACCCTTA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
62D.2	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	CCTACTCGAT	CTCTCGTATG
62D.3	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
65D.7	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	CCTACTCGAT	CTCTCGTATG
65D.3	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	CCTACTCGAT	CTCTCGTATG
67B	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
69A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
69B	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
76A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCCACTCGAT	CTCTTCTATG
76E	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTAGTT	GCTACTCGAT	CTCTTCTATG
78A.10	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
78A.5	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
8	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
80A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTTTTCTATG
82A	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
85A	ATGACCCTTA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
86B	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
90D	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTTCTATG
93C.2	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
93C.3	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
98A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGTATG
126E	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	CCTACTCGAT	CTCTCGTATG
103A	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCAAT	CTCTTCTATG
50D	ATGACCCTCA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTAYTGGAT	CTCTTATATG
68A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCAATGATT	GATATATGAT	CTATTCTAWK
75A	ATGACCCTCA	CTCTCGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGRT	CTCTTCTATG

Bases 175-234

ETS type 2

121C	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
18	ATGACCCGCA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	GCTACTCGAT	CTCTCGGATG
19	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATA
20	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTAGAT	CTCTTGGATG
21	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTAGAT	CTCTTGGATG
122A	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
126.29	ATGACCCGTA	CTCTCAGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
26	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
52B	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
57B	GTGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
65C.3	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
65C.4	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
93C	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
118A	ATGACCCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
126.BC1	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
126E.1	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCCTGGATG
126E.3	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
126E.4	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
126A	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
61	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG

ETS type 3

121B	-----	-----	-----	-----	-----	-----
66C	-----	-----	-----	-----	-----	-----
66D	-----	-----	-----	-----	-----	-----
67A	-----	-----	-----	-----	-----	-----
69B	-----	-----	-----	-----	-----	-----
71A	-----	-----	-----	-----	-----	-----
77B	-----	-----	-----	-----	-----	-----
82A	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
86B.24	-----	-----	-----	-----	-----	-----
86B.29	-----	-----	-----	-----	-----	-----
86B.5	-----	-----	-----	-----	-----	-----
87B	-----	-----	-----	-----	-----	-----
90D	-----	-----	-----	-----	-----	-----

ETS type 4

118A	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
121B	ATGACCCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
123F	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
126.29.1	ATGACCCGTA	CTCTCAGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
126.29.3	ATGACCCGTA	CTCTCAGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATG
63B	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
81A	ATGACTCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCACTGGTT	GCTACTCGAT	CTCTTGGATG
76A	ATGACCCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCACTGGTT	GCTACTCGAT	CTCTTGGATG
76E	ATGACCCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCATTGGTT	GCTACTCGAT	CTCTCGGATG
78B	ATGACTCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCACTGGTT	GCTACTCGAT	CTCTTGGATG
84A.1	ATGACCCGTA	CTCTTGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
84A.2	ATGACCCGTA	CTCTTGGGTG	CTTGTGTGGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
83B	-----A	TTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	GCTACTCGAT	CTCTCGGATG
87A	-----	-----	-----	-----	-----	-----
126.WB3	ATGACCCGTA	CTCTCGGGTG	CTTGTGAAGG	TGCACTGGTT	TCTACTCGAT	CTCTTGGATT
Pur-34	AAGACCTGCA	TTGTTGGGTT	TTTGTGTCCG	TCCACGTGTT	GCTACTCGAT	CTCTCGGATG
Pur-132	AAGACTAGCA	TTGTTAGGTT	TTTGTGTCCG	TCCATGTGTT	GCTACTCGAT	CTCTCGGATG

Bases 235-294

ETS type 1

1	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
10	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCCY-GAM	GTC-AGATGC	AAAA-GCATT
14	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
15	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
12	CRGAACATAC	AAAGGGCGCG	GGGGTTTTCA	GCCCC-GAC	GTC-AGATGC	AAAW-GCATT
17	CGGAACATAC	AAAGGGCGCG	GGGGTCTTTA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
25	CAGAACATAC	AAAGGGTGCG	GGGGTCTTCA	GCCCTC-GAC	GTC-CGATGC	AAAA-GCATT
26	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
27C.1	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
27C.10	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
28A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCCT-GAM	GTC-AGATGC	AAAA-GCATT
3	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
41B	TAGAACATAC	AAAGGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTC-CAATGC	AAAA-GCATT
45C	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
5	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
101A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
102B	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
104	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCCT-GAC	GTC-AGATGC	AAAA-GCATT
108C	CGGAACATAC	AAAGGGCGCG	GTGGTCTTCA	GCCCC-GAC	GTC-AGAAGC	AAAA-GCATT
111A.4	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCC	GCCCC-GAC	GTC-AGACGC	AAAA-GCATT
111A.5	CAGAACATAC	AAAGGGTACG	GGGGTCTTCA	GCCCC-GAC	GTC-CGATGC	AAAA-GCATT
120C	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
123G	CGGAACATAC	AAAGGSGGG	GGGGT-----	-----	-----	-----
131	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCCT-GAA	GTC-AGATGC	AAAA-GCATT
30A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
57B	CGGAACATAC	AAAGGGCGAG	GGGGTCTTCA	GCCCTC-GAT	GTCC-GATGC	AAAA-CCATT
58B	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGT	AAAA-GCATT
62D.2	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-CGATGC	AAAA-CCATT
62D.3	CGGAACATAC	AAAGGGCGCG	GGGGTTTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-CCATT
65D.7	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-CGATGC	AAAA-CCATT
65D.3	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-CGATGC	GAAA-CCATT
67B	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCCA-GAC	GTC-AGATGC	AAAA-GCATT
69A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
69B	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
76A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCCN-TAC	GTC-AGATGC	AAAA-GCATT
76E	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
78A.10	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-TAC	GTC-AGATGC	AAAA-GCATT
78A.5	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-TAC	GTC-AGATGC	AAAA-GCATT
8	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGAT-C	AAAA-GCATT
80A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCTCC-GAC	GTC-AGATGC	AAAA-GCATT
82A	CAGAACATAC	AAAGGGTGCG	GGGGTCTTCA	GCCCTC-GAC	GTCC-GATGC	AAAA-GCATT
85A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGT	AAAA-GCATT
86B	CAGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT
90D	CAGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AAATGC	AAAA-GCATT
93C.2	CGGAACATAC	AAAGGGCGAG	GGGGTCTTCA	GCCCTC-GAT	GTC-CGATGA	GAA--CCATT
93C.3	CGGAACATAC	AAAGGGCGAG	GGGGTCTTCA	GCCCTC-GAT	GTCC-GATGC	AAAA-CCATT
98A	CGGAACATAC	AAAGGGCGAG	GGGGTCTTCA	GCCCTC-GAT	GTCC-GATGC	AAAA-CCATT
126E	CGGAACATAC	AAAGGGCGCG	AGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
103A	CGGAACATAC	AAAGGGCGCG	GGAGTCTTCA	GCCCCT-GAC	GTCCAGATGC	AAAA-GCATT
50D	CGGAACATAC	AAAGGGTGCG	GGGGTCTTCA	GCCCCM-GAC	GTC-AGAWGC	AAGA-GCATT
68A	CGGAACAATC	AAAGAGCKCG	GGGGTCT---	-----	-----	-----
75A	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTC-AGATGC	AAAA-GCATT

Bases 235-294

ETS type 2

121C	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAACCATT
18	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-GCATT
19	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
20	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
21	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
122A	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
126.29	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
26	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
52B	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
57B	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAACCATT
65C.3	CGGAACATAC	AAAAGGTGCG	GGGGTTTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-GCATT
65C.4	CGGAACATAC	AAAAGGTGCG	GGGGTTTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-GCATT
93C	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-TAAAA	ACCATTTCATT
118A	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAA--GCATT
126.BC1	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAASCATT
126E.1	CGGAACATAC	AAAAGGCG--	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
126E.3	CGGAACATAC	AAAAGGCG--	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
126E.4	CGGAACATAC	AAAAGGCG--	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
126A	CGGAACATAC	AAAAGGCG--	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT
61	CAGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAAGCATT

ETS type 3

121B	-----	-----	-----	-----	-----	-----
66C	-----	-----	-----	-----	-----	-----
66D	-----	-----	-----	-----	-----	-----
67A	-----	-----	-----	-----	-----	-----
69B	-----	-----	-----	-----	-----	-----
71A	-----	-----	-----	-----	-----	-----
77B	-----	-----	-----	-----	-----	-----
82A	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
86B.24	-----	-----	-----	-----	-----	-----
86B.29	-----	-----	-----	-----	-----	-----
86B.5	-----	-----	-----	-----	-----	-----
87B	-----	-----	-----	-----	-----	-----
90D	-----	-----	-----	-----	-----	-----

ETS type 4

118A	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAACCATT
121B	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	ACCCCC-GAC	GTC-GATGC	AAA--GCATT
123F	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-GCATT
126.29.1	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAACCATT
126.29.3	CGGAACATAC	AAAAGGCGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAAACCATT
63B	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-GCATT
81A	CGGAACATAC	AAAGGGCGTG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAAA-GTATT
76A	CGGAACATAC	AAAGGGCGTG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAAA-GTATT
76E	CGGAACATAC	AAAGGGCGTG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAAA-GTATT
78B	CGGAACATAC	AAAGGGCGTG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAAA-GTATT
84A.1	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAA--GCATT
84A.2	CGGAACATAC	GAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTC-GATGC	AAA--GCATT
83B	CGGAACATAC	AAAGGGCGCG	GGGGTCTTCA	GCCCC--GAC	GTCC-GATGC	AAAA-GCATT
87A	-----	-----	-----	-----	-----	-----
126.WB3	CGGAACATAC	AAAAGGTGCG	GGGGTCTTCA	GCCCC-GAC	GTCC-GATGC	AAAA-GCATT
Pur-34	CAGAACATGT	TGCGGGCGC-	AGGTCTTTCG	GCCCTTGAC	GTCCAAA---	ACAAAGCGTT
Pur-132	CGGAACATGT	TGCGGGTGT-	AGGTCTTTCG	GCCCTTGAC	GTCCAAA---	ACAAAGCGTT

Bases 295-354

ETS type 1

1	T-GTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
10	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
14	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
15	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
12	TTGTCCCTAK	ACACGAACGA	CAGTTGCGCG	CTGTTGCACG	GTTCTCGACC	TT-CACGG-C
17	TTGTCCCTAT	ACACGAACGG	CAGTCGCGCG	C-GTTGCAGG	TT-CTKGACC	TT-CACGG-C
25	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTTGACC	TT-CACGG-C
26	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
27C.1	TTGTCCCTAT	AC-----	-----	-----	-----	-----
27C.10	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
28A	TTGTCCCTAT	ACACGAACGA	CAGTSGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
3	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CATGG-C
41B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
45C	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
5Q	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
101A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CTT-TYGACC	TT-CACGAGC
102B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
104	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	WT-CACGG-C
108C	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
111A.4	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
111A.5	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
120C	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
123G	-----	-----	-----	-----	-----	-----
131	TTGTCCCTAT	ACACGAACGA	CAGTGGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
30A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
57B	TTGTCCCTAT	ACACGAATGA	TAGTTGTGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
58B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
62D.2	TTGTCCCTAC	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGGG-C
62D.3	TTGTCCCTAT	ACACGAATGA	CAGTCGTGCG	C-GCTGTAGG	CT-CTCGACC	TT-TACGG-C
65D.7	TTGTCCCTAC	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGGG-C
65D.3	TTGTCCCTAC	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
67B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
69A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
69B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
76A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGGG-C
76E	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
78A.10	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
78A.5	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGGG-C
8	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
80A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CTTCTCGACC	TT-CACGG-C
82A	TTGCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGGG-C
85A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CTCMTTGACC	TTGCRCGG-C
86B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	TACTCTCGACC	TTGCACGG-C
90D	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	TT-CTCGACC	TT-CACGG-C
93C.2	TTGTCCCTAT	ACACGAATGA	CAGTTGTGCA	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
93C.3	TTGTCCCTAT	ACACGAATGA	CAGTTGTGCA	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C
98A	TTGTCCCTAT	ACACGAATGA	CAGTTGTGCA	C-GTTGCAGG	CT-CTCGACC	TT-CTCGG-C
126E	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
103A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-A-----	-----	-----
50D	TTGTCCCTAT	ACA--AAC-A	TATT-----	-----	-----	-----
68A	-----	-----	-----	-----	-----	-----
75A	TTGTCCCTAT	MCACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGACC	TT-CACGG-C

Bases 295-354

ETS type 2

121C	TTGTCCCTAT	ACACGAACAA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
18	TTGTCCCTAA	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGT	CT-ATCGACC	TT-CAAGG-C
19	TTGTCCCTAT	ACACGAATGA	CAGTCGCGTG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
20	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
21	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
122A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126.29	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
26	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGGG-C
52B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
57B	TTGTCCCTAT	ACACGAACGA	CAGTCACGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
65C.3	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
65C.4	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
93C	TTGTCCCTAT	ACACGGACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
118A	TTGTCCCTAT	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126.BC1	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126E.1	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126E.3	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126E.4	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126A	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
61	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CTTCTGGGCC	TT-CACGGAC

ETS type 3

121B	-----	-----	-----	-----	-----	-----
66C	-----	-----	-----	-----	-----	-----
66D	-----	-----	-----	-----	-----	-----
67A	-----	-----	-----	-----	-----	-----
69B	-----	-----	-----	-----	-----	-----
71A	-----	-----	-----	-----	-----	-----
77B	-----	-----	-----	-----	-----	-----
82A	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
86B.24	-----	-----	-----	-----	-----	-----
86B.29	-----	-----	-----	-----	-----	-----
86B.5	-----	-----	-----	-----	-----	-----
87B	-----	-----	-----	-----	-----	-----
90D	-----	-----	-----	-----	-----	-----

ETS type 4

118A	TTGTCCCTAT	ACACGAACAA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
121B	TTGTCCCTAT	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
123F	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126.29.1	TTGTCCCTAT	ACACGAACTA	CAGTCGCGCG	T-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
126.29.3	TTGTCCCTAT	ACACGAACTA	CAGTCGCGCG	T-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
63B	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
81A	TTGTCCCTAT	ACACGAACGA	CAATCACCCG	C-GTTGCAGG	CT-MTYGACC	TT-CACAG-C
76A	TTGTCCCTAT	ACACGAACGG	CAGTCACCCG	C-GTTGCAGG	CT-CTCGACC	TT-CACAG-C
76E	TTGTCCCTAT	ACACGAACGA	CAGTCACCCG	C-GTTGCAGG	CT-CTCGACC	TT-CACAG-C
78B	TTGTCCCTAT	ACACGAACGA	CAATCACCCG	C-GTTGCAGG	CT-CTCGACC	TT-CACAG-C
84A.1	TTGTCCCTAT	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
84A.2	TTGTCCCTAT	ACAAGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
83B	TTGTCCCTAA	ACAAGAACGA	CAGTCGCGCG	C-GTTGCTGG	CT-ATCGACC	TT-CAAGG-C
87A	-----	-----	-----	-----	-----	-----
126.WB3	TTGTCCCTAT	ACACGAACGA	CAGTCGCGCG	C-GTTGCAGG	CT-CTCGGCC	TT-CACGG-C
Pur-34	TT-YCGCTAG	ACACGAACGA	CAGTCGCGTA	C-GTTGCGGG	CTTTT-GACC	TT-CGCGG-C
Pur-132	TT-TCGCTAG	ACACGAATGA	CTGTCGCGTA	C-GTTGCGGG	CTTGT-GACC	TT-CGCGG-C

Bases 355-382

ETS type 1

1	CC-YTAGTC-	TTGCACCGCG	TCGA-GT-
10	CC-TTAGTC-	TTGCACCTCG	TCGA-GTC
14	CC-TTAGTC-	TTGCACCKCG	TCGACGTC
15	CC-TTAGTC-	TTGCACCTCG	TCGACGTC
12	CC-TTAGTC-	TTGCACCTCG	TAGA-GTC
17	CC-TTAGTC-	WTGCAC----	-----
25	CC-TTAGTC-	TTGCACCGCG	TTGACGTG
26	CC-TTAGTC-	TTGCACCTCG	TCGACGTC
27C.1	-----	-----	-----
27C.10	CC-TTAGTC-	TTGCA-----	-----
28A	CC-TTAGTC-	TTGCACCTCG	TCGACGTC
3	CC-TTAGTC-	TTGCACCGCG	TCGACGTC
41B	CC-TTAGTC-	TTGCACCGTG	TCGACYTC
45C	CC-TTAGTC-	TTGCACCGCG	TCGA-GTC
5	CC-TTAGTC-	TTGCACCTCG	TCGACGTC
101A	CC-TTAGTTC	TTG-----	-----
102B	CC-TTAGTCG	TTGCAC----	-----
104	CC-TTAGTTC	TT-----	-----
108C	CC-TTAGTC-	TTGCACC---	-----
111A.4	CC-TTAGTC-	TTGCA-----	-----
111A.5	CC-TTAGTC-	TTGCACCGCG	T-----
120C	CC-TTAGT--	-----	-----
123G	-----	-----	-----
131	CC-TTAGT--	-----	-----
30A	CC-TTAGTC-	TTGCACCTCG	TCGACGTC
57B	CCCTTAGTC-	TTGCACCGCG	TCAACGTC
58B	CC-TTAGT--	-----	-----
62D.2	CC-TTA-TTC	TTGCACCGCA	TCGATGTA
62D.3	CC-TTAGT-C	TTGCACCGCG	TCAACGT-
65D.7	CC-TTA-TTC	TTGCACCGCA	TCGATGTC
65D.3	CC-TTA-TTC	TTGCACCGCA	TCGATGTC
67B	CC-TTAGTC-	TTGCACCGCG	TCGACGTC
69A	C-----	-----	-----
69B	C-----	-----	-----
76A	CC-TTAGTC-	TT-----	-----
76E	CC-TTAGTC-	TTGCACCGCG	TCGACGTC
78A.10	CC-TTAGTC-	TTGCACCGCG	TCGACGTC
78A.5	CC-TTAGTC-	TTGCA-----	-----
8	CC-TTAGTC-	TTGCACC-TG	TCGACGTC
80A	CC-TTAGTTC	TTGCAAC---	-----
82A	CCCTTAGTC-	TTGCACCGCG	TT-----
85A	CC-TTAGTTC	TTGC-----	-----
86B	CC-TTAGTC-	TTGCA-C---	-----
90D	CC-TTAGTC-	TTGCACCG--	-----
93C.2	CC-TTAGTC-	TTACACCGCG	TCAACGTC
93C.3	CC-TTAGTCC	TTACACCGCG	TCAACGTC
98A	CC-TTAGTC-	TTACACCGCG	TCCA----
126E	CC-TTAGTCC	-----	-----
103A	-----	-----	-----
50D	-----	-----	-----
68A	-----	-----	-----
75A	CC-TTAGTCT	-TGCAAC---	-----

Bases 355-382

ETS type 2

121C	CC-TTAGTCC	-TGCACCACG	TCAACGTC
18	CC-TTAGCCC	-TGCACCGCG	TCGACGTC
19	CC-TTAGTCC	-TGCACCGCG	TCAACATC
20	CC-TTAGTCC	-AGCACCGCG	TCAACGTC
21	CC-TTAGTCC	-AGCACCGCG	TCAACGTC
122A	CC-TTAGTCC	-TGCACCGCG	T--ACAT-
126.29	CC-TTAGTCA	-TGCACCGCG	TCAACGTC
26	CCCTTAGTCC	-AGCACCGCG	TCAA----
52B	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
57B	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
65C.3	CCCTTAGTCC	CTGCACCGCG	TCGACGTC
65C.4	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
93C	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
118A	CC-TTAGTCC	-TGCACCGCG	T-----
126.BC1	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
126E.1	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
126E.3	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
126E.4	CC-TTAGTCC	-TGCACCGCG	TCAACGTC
126A	CC-TTAGTCC	-TGCACCGCG	TCA---TA
61	CC-TTAGTCC	CTGC-----	-----

ETS type 3

121B	-----	-----	-----
66C	-----	-----	-----
66D	-----	-----	-----
67A	-----	-----	-----
69B	-----	-----	-----
71A	-----	-----	-----
77B	-----	-----	-----
82A	-----	-----	-----
83B	-----	-----	-----
86B.24	-----	-----	-----
86B.29	-----	-----	-----
86B.5	-----	-----	-----
87B	-----	-----	-----
90D	-----	-----	-----

ETS type 4

118A	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
121B	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
123F	CC-TTAGTCC	-----	-----
126.29.1	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
126.29.3	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
63B	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
81A	CC-TTAGTCC	-TGCACCGCA	TCGACGTC
76A	CC-TTAGTCC	-TGCACCGCA	TCGACGTC
76E	CC-TTAGTCC	-TGCATTGCA	TCGACGTC
78B	CC-TTAGTCC	-TGCACCGCA	TCGACGTC
84A.1	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
84A.2	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
83B	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
87A	-----	-----	-----
126.WB3	CC-TTAGTCC	-TGCACCGCG	TCGACGTC
Pur-34	CA-MTAGTCT	-CGC-----	-----
Pur-132	CA-TTAGTCC	-CGT-----	-----

Bases 383-442

α and β region→

ETS type 1

1	TT-TGTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
10	TT--GTTGCC	TTT-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
14	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
15	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
12	TT--GAWGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CCTGMTTCTG
17	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
25	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
26	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
27C.1	-----	-----ATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
27C.10	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
28A	TT--GTTGCC	TTT-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
3	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
41B	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
45C	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
5	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
101A	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
102B	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
104	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
108C	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
111A.4	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
111A.5	TT--GTTGCC	TTC-GAAATG	GTGGATTAGA	GACGAATTGG	ATTGCCTATG	GTTGGTTGTG
120C	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
123G	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
131	TT--GTTGCC	TTT-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
30A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
57B	TT-TGTTGCC	TTC-GAAATG	GTGGATGAGA	GGCAAATTGG	ATTGCCTATG	CTTGCTTGTG
58B	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GATGAATTGG	ATTGCCTATG	CTTGCTTGTG
62D.2	TT--CTTGCC	TTC-AAAATG	GTGGATGAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
62D.3	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GGCAAATTGG	ATTGCCTATG	CTTGCTTGTG
65D.7	TT--CTTGCC	TTC-AAAATG	GTGGATGAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
65D.3	TT--CTTGCC	TTC-AAAATG	GTGGATGAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
67B	TTGTCTCGCC	TTCCGAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
69A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
69B	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
76A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
76E	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
78A.10	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGT	ATTGCCTATG	CTTGCTTGTG
78A.5	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGT	ATTGCCTATG	CTTGCTTGTG
8	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
80A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
82A	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
85A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GATGAATTGG	ATTGCCTATG	CTTGCTTGTG
86B	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACCAATTGG	ATTGCCTATG	CTTGCTTGTG
90D	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACCAATTGG	ATTGCCTATG	CTTGCTTGTG
93C.2	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GGCAAATTGG	ATTGCCTATG	CTTGCTTGTG
93C.3	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GGCAAATTGG	ATTGCCTATG	CTTGCTTGTG
98A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GGCAAATTGG	ATTGCCTATG	CTTGCTTGTG
126E	TT--CTTGCC	TTC-AAAATG	GTGGATGAGC	GGCAAATTGG	ATTGCCTATG	CTTGCTTGTG
103A	TT--GTTGCC	TTC-AAAATG	GTGGATAAGC	GGCGAATTGG	ATTGCCTATG	CTTGCTTGTG
50D	TT--GTTGCC	TTC-AAAATG	GTGGATAAGA	GGCGAATTGG	ATTGCCTATG	CATGCTTGTG
68A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG
75A	TT--GTTGCC	TTC-GAAATG	GTGGATGAGA	GACGAATTGG	ATTGCCTATG	CTTGCTTGTG

Bases 383-442

ETS type 2

121C	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
18	-----	-----	-----	-----	-----CTATG	CTTGCTTGTG
19	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
20	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
21	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
122A	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
126.29	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
26	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
52B	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
57B	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
65C.3	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
65C.4	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
93C	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
118A	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
126.BC1	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
126E.1	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
126E.3	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
126E.4	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
126A	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG
61	-----	-----	-----	-----	-----CTATG	CCTGCTTGTG

ETS type 3

121B	-----	-----	-----	-----	-----	-----
66C	-----	-----	-----	-----	-----	-----
66D	-----	-----	-----	-----	-----	-----
67A	-----	-----	-----	-----	-----	-----
69B	-----	-----	-----	-----	-----	-----
71A	-----	-----	-----	-----	-----	-----
77B	-----	-----	-----	-----	-----	-----
82A	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
86B.24	-----	-----	-----	-----	-----	-----
86B.29	-----	-----	-----	-----	-----	-----
86B.5	-----	-----	-----	-----	-----	-----
87B	-----	-----	-----	-----	-----	-----
90D	-----	-----	-----	-----	-----	-----

ETS type 4

118A	-----	-----	-----	-----	-----	-----
121B	-----	-----	-----	-----	-----	-----
123F	-----	-----	-----	-----	-----	-----
126.29.1	-----	-----	-----	-----	-----	-----
126.29.3	-----	-----	-----	-----	-----	-----
63B	-----	-----	-----	-----	-----	-----
81A	-----	-----	-----	-----	-----	-----
76A	-----	-----	-----	-----	-----	-----
76E	-----	-----	-----	-----	-----	-----
78B	-----	-----	-----	-----	-----	-----
84A.1	-----	-----	-----	-----	-----	-----
84A.2	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
87A	-----	-----	-----	-----	-----	-----
126.WB3	-----	-----	-----	-----	-----	-----
Pur-34	TTG-GTTGCC	TTC-AGAATG	GTGGATGTGC	GGCGAATCGG	ATTGCTTATG	CTTGCTTGGG
Pur-132	-----	-----	-----	-----	-----	-----

Bases 443-502**ETS type 1**

1	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
10	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
14	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
15	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
12	CGGAGCCCTT	ACGGGCAGTG	CACAA-GSTT	GTCGACGATG	GCACCCCGAG	GAGTTGCGAT
17	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
25	CGGAGCCCTT	ACGAGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
26	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCCGAG	GAGCTGCGAT
27C.1	CGAAGCCCTT	GCGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
27C.10	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
28A	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
3	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
41B	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
45C	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCCGAG	GAGCTGCGAT
5	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
101A	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCCGAG	GAGCTGCGAT
102B	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
104	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCTGAG	GAGCTGCGAT
108C	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
111A.4	CGAAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCCGAG	GAGCTGCGAT
111A.5	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAC
120C	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGCCCATG	GCACCCCGAG	GAGCTGCGAT
123G	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
131	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
30A	CGGAGCCCTT	ATGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
57B	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGTGAT
58B	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
62D.2	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
62D.3	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGATGATG	GCACCCCGAG	GAGCTACGAT
65D.7	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
65D.3	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
67B	CGGAGCCATT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
69A	CGGAGCCCTT	ACGGGCAGTG	TACAA-GCTT	GTTGACGATG	GCGCCCCGAG	GAGCTGCGAT
69B	CGGAGCCCTT	ACGGGCAGTG	TACAA-GCTT	GTTGACGATG	GCGCCCCGAG	GAGCTGCGAT
76A	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
76E	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
78A.10	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
78A.5	CGGAGCCCTT	ACTGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
8	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTTGACGATG	GCACCCCGAG	GAGCTGCGAT
80A	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
82A	CGGAGCCCTT	ACGGGCAACG	CGCAA-GCTT	GTCGACAATG	GCACCCCGAG	GAGCTGCGAT
85A	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT
86B	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAA	GAGCTGCGAT
90D	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAA	GAGCTGCGAT
93C.2	CAGAGCCCTT	ATGGGTAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTACGAT
93C.3	CAGAGCCCTT	ACGGGTAATG	CACAA-TGTC	GACGGTGGCA	CCCCGAGGAG	CTACGATGAT
98A	CAGAGCCCTT	ACGGGTAATG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTACGAT
126E	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACTATG	GCACCCCGAG	GAGCTGCGAT
103A	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCTGAG	GAGCTGCGAT
50D	CGGAGCCCTT	ACGGGCAATG	CACAA-GCTT	GTCGACAATG	GCACCCCAAG	GAGCTGCGAT
68A	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATC	GCACCCCGAG	GAGCTGCGAT
75A	CGGAGCCCTT	ACGGGCAGTG	CACAA-GCTT	GTCGACGATG	GCACCCCGAG	GAGCTGCGAT

Bases 443-502

ETS type 2

121C	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
18	CAAAGCCCTC	ACGGGCAGAG	CACAA-GCTT	GTCGAGGATG	GCACTCCGAG	GAGCTGCGAT
19	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
20	CAAAGCCCAC	CTATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
21	CAAAGCCCAC	CTATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
122A	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
126.29	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
26	CAAAGCCCAC	CTATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
52B	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
57B	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
65C.3	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	GTCGACGATG	GCACCCCGAG	GAGCTTCAGT
65C.4	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	GTCGACGATG	GCACCCCGAG	GAGCTTCAGT
93C	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
118A	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
126.BC1	CAAAGCCCAC	C-ATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT
126E.1	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTTGACGATG	GCACTCCGAG	GAGCTGCAGT
126E.3	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTTGACGATG	GCACTCCGAG	GAGCTGCAGT
126E.4	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTTGACGATG	GCACTCCGAG	GAGCTGCAGT
126A	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTTGACGATG	GCACTCCGAG	GAGCTGCAGT
61	CAAAGCCCAC	CCATGCAATG	CACAAAGCTT	TTCGACGATG	GCACTCCGAG	GAGCTGCAGT

ETS type 3

121B	-----	-----	-----	-----	-----	-----
66C	-----	-----	-----	-----	-----	-----
66D	-----	-----	-----	-----	-----	-----
67A	-----	-----	-----	-----	-----	-----
69B	-----	-----	-----	-----	-----	-----
71A	-----	-----	-----	-----	-----	-----
77B	-----	-----	-----	-----	-----	-----
82A	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
86B.24	-----	-----	-----	-----	-----	-----
86B.29	-----	-----	-----	-----	-----	-----
86B.5	-----	-----	-----	-----	-----	-----
87B	-----	-----	-----	-----	-----	-----
90D	-----	-----	-----	-----	-----	-----

ETS type 4

118A	-----	-----	-----	-----	-----	-----
121B	-----	-----	-----	-----	-----	-----
123F	-----	-----	-----	-----	-----	-----
126.29.1	-----	-----	-----	-----	-----	-----
126.29.3	-----	-----	-----	-----	-----	-----
63B	-----	-----	-----	-----	-----	-----
81A	-----	-----	-----	-----	-----	-----
76A	-----	-----	-----	-----	-----	-----
76E	-----	-----	-----	-----	-----	-----
78B	-----	-----	-----	-----	-----	-----
84A.1	-----	-----	-----	-----	-----	-----
84A.2	-----	-----	-----	-----	-----	-----
83B	-----	-----	-----	-----	-----	-----
87A	-----	-----	-----	-----	-----	-----
126.WB3	-----	-----	-----	-----	-----	-----
Pur-34	CGGAGCCTTT	GTGGGCAATG	CACAA-CATG	TCGACGATGG	CACCTCGAGC	AGCCGCGATT
Pur-132	-----	-----	-----	-----	-----	-----

Bases 503-511

ETS type 1

1 GATTCCCAA
10 GATTCCCAA
14 GATTCCCAA
15 GATTCCCAA
12 GATTCCCAA
17 GATTCCCAA
25 GATTCCCAA
26 GATTCCCTAA
27C.1 GATTCCCAA
27C.10 GATTCCCAA
28A GATTCCCAA
3 GATTCCCAA
41B GATTCCCAA
45C GATTCCCTAA
5 GATTCCCAA
101A GATTCCCTAA
102B GATTCCCAA
104 GATTCCCTAA
108C GATTCCCAA
111A.4 GATTCCCTAA
111A.5 GATTCCCAA
120C GATTCCCAA
123G GATTCCCAA
131 GATTCCCAA
30A GATTCCCAA
57B GATTCCCAA
58B GATTCCCAA
62D.2 GATTCCCAA
62D.3 GATTCTCAA
65D.7 GATTCCCAA
65D.3 GATTCCCAA
67B GATTCCCAA
69A GATTCCCAA
69B GATTCCCAA
76A GATTCCCAA
76E GATTCCCAA
78A.10 GATTCCCAA
78A.5 GATTCCCAA
8 GATTCCCAA
80A GATTCCCAA
82A GATTCCCTAA
85A GATTCCCAA
86B GATTCCCAA
90D GATTCCCAA
93C.2 GATTCCCAA
93C.3 GATTCCCAA
98A GATTCCCAA
126E GATTCCCAA
103A GATTCCCTAA
50D GATTCCCTAA
68A GATTCCCAA
75A GATTCCCAA

Bases 503-511
ETS type 2
 121C GATTCCCAA
 18 GATTCCCAA
 19 GATTCCCAA
 20 GATTCCCAA
 21 GATTCCCAA
 122A GATTCCCAA
 126.29 GATTCCCAA
 26 GATTCCCAA
 52B GATTCCCAA
 57B GATTCCCAA
 65C.3 GATTCCCAA
 65C.4 GATTCCCAA
 93C GATTCCCAA
 118A GATTCCCAA
 126.BC1 GATTCCCAA
 126E.1 GATTCCCAA
 126E.3 GATTCCCAA
 126E.4 GATTCCCAA
 126A GATTCCCAA
 61 GATTCCCAA
ETS type 3
 121B -----
 66C -----
 66D -----
 67A -----
 69B -----
 71A -----
 77B -----
 82A -----
 83B -----
 86B.24 -----
 86B.29 -----
 86B.5 -----
 87B -----
 90D -----
ETS type 4
 118A -----
 121B -----
 123F -----
 126.29.1-----
 126.29.3-----
 63B -----
 81A -----
 76A -----
 76E -----
 78B -----
 84A.1 -----
 84A.2 -----
 83B -----
 87A -----
 126.WB3 -----
 Pur-34 GACATCCCA
 Pur-132 -----

Appendix B. AFLP presence/absence data matrix

ID codes correspond to those found in Table 3.1. Presence of the fragment is coded as '1' and absence as a '0'. The AFLP selective primer pair that produced the fragment and the size of the individual fragments can be found in Appendix C.

Fragments 1-50

1	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
3	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
5	00000	00000	00010	00000	00001	10001	01000	00000	00000	00000
8	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
10	00000	00000	00010	00000	00001	10001	01000	00000	00000	00000
12	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
13	10000	00000	00010	00000	00001	10111	01000	10000	00000	00110
14	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
16	10000	00000	00000	00000	00001	10011	01011	00000	00000	00000
17	00000	00001	00000	01001	00001	10001	00000	00001	10000	00000
18	00000	00000	00000	00000	00001	10000	01011	00001	10001	00001
19	00000	00000	00000	00001	00001	10000	01000	00000	00000	00000
20	00000	00100	00000	00000	00001	10000	01000	00000	00000	00000
25	00000	00000	00000	00000	00001	00001	01000	00000	00000	00000
26	00010	00000	00000	00000	00001	10000	00000	00000	01000	00000
27A	10000	00000	00000	01001	00001	10011	01110	00000	00000	00000
27B	10000	00000	00000	01101	00001	10001	01110	00000	00000	00000
27C	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
28A	00000	00000	00000	00000	10001	10001	01000	00000	00000	00000
29A	00000	00000	00000	00000	00001	10001	01100	00000	00000	00000
30A	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
30D	10000	00000	00010	01001	00001	10001	01000	00000	00000	00100
30E	00000	00000	00000	00000	00001	10011	01011	00000	00000	00000
30F	10000	00100	00000	00001	00001	10001	01000	00000	00010	00000
31	10000	10101	00010	01001	00001	00101	01001	00000	00000	00100
32B	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
36B	00010	00000	00000	00000	00001	01001	00000	00000	00000	00001
37A	00000	00000	00000	00000	00001	10010	01011	00000	00000	00000
37B	10000	00000	00000	00000	00001	10011	01010	00000	00000	00000
37C	10001	01000	10100	01101	00001	00000	01000	00000	00000	00000
39	00000	00000	00000	00101	00001	00101	01011	00000	00000	00000
41A	10000	01000	00010	01001	00001	10000	01000	00000	00000	00000
41B	00000	00000	00000	00001	00001	10000	01000	00000	00000	00000
41C	00000	00000	00000	00001	00001	10000	01001	00000	00000	10000
43	00000	10000	00000	00000	00001	10000	01010	00000	00000	00000
45A	00000	10011	00100	01000	00011	00000	01000	00000	00000	10100
45B	10000	01001	01100	01101	00001	00010	01001	00000	00000	00000
45C	00000	00000	00000	00100	00011	10000	01000	00000	00000	00000
46A	00000	00000	00000	00010	00001	10101	01010	00000	00000	00000
46B	00000	00000	00000	00000	00001	10101	01000	00000	00000	00000
46C	00000	00000	00000	00000	00001	10101	01000	00000	00000	00000
48A	10000	00000	00000	00000	00001	10000	01011	00000	00000	00000
48B	10001	00001	00010	10010	00001	10011	01010	00000	00000	00000
48C	00000	00000	00000	00000	00001	10111	01010	00000	00000	00000
48D	00000	10000	00000	10000	00001	10000	01000	00000	00000	00000
50A	10000	00000	00000	00000	00001	10001	01010	11000	00000	00000
50B	00100	00000	00000	00000	00001	00000	00000	00000	00000	00000

Fragments 1-50

50C	00000	01000	00000	00000	00001	10001	01010	00000	10000	00000
50D	00000	00000	00000	00000	00001	00001	01000	00000	00000	00000
52B	00010	00000	00000	00000	00001	10000	01000	00000	00000	00000
54A	00010	00000	00000	00000	00001	01000	00000	00000	00000	00000
55A	00010	00000	00000	00000	00001	01000	00000	00000	00000	00000
57B	00010	00000	00000	10000	00001	10000	01000	00000	00000	00000
58B	00000	00000	00000	00000	00001	10101	01000	00000	00000	00000
58D	00010	00000	00000	00000	00001	10001	01010	00000	00000	00000
61	00010	00100	10000	00001	00001	10010	01000	00000	00000	00000
62D	00000	00000	00000	00010	00001	10000	01000	00000	00000	00000
63B	00000	00000	00000	00010	00001	10000	01000	00000	00000	00000
65C	00000	00000	01000	00000	00001	10000	01000	00001	00000	00000
65D	00000	00000	00000	00000	00001	10000	01000	00000	00000	00000
66C	00010	00000	00000	00000	00001	10000	01000	00000	01000	00000
66D	00010	00000	10000	00000	00001	10000	01000	00000	01000	00000
67A	00010	00000	00000	00000	00001	10000	00000	00000	01101	00000
67B	00000	00000	10000	00000	00001	10000	00000	00000	01001	01000
68A	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
68B	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
68C	00000	00000	00000	00000	00001	10011	01010	00000	00000	00000
69A	00010	00000	00000	00000	00001	10000	00000	00000	01000	00000
69B	00010	00000	00000	00000	00001	10000	00000	01000	01000	00000
70A	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
70B	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
71A	00010	00000	00000	00000	00001	10000	00000	00000	01000	00000
72A	10000	00000	00000	00000	00001	10001	01000	00000	00000	00000
72B	10000	00000	00000	00000	00001	10001	01000	00000	00000	00000
74D	00000	00000	00001	00000	01001	01000	01000	00000	00000	00000
75A	00000	00000	00010	00000	00001	10001	00000	00001	10000	00100
76A	00000	00000	00000	00000	00001	10101	01000	00000	01001	00000
76E	00000	00000	10000	00000	00001	10001	01000	00000	01000	00000
77B	00010	00000	00000	00000	00001	10001	00000	00000	01000	01000
78A	00010	00000	00100	00000	00001	10000	00000	00000	00000	01000
79	00010	00000	00000	00000	00001	10000	00000	00000	01000	00000
80A	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
80C	10000	00000	00001	00001	00001	10011	01000	10000	00000	00000
80D	10000	00100	00000	01001	00001	10001	01000	00000	00000	00000
81A	01010	00000	00000	00000	00001	10000	00000	00000	00000	01000
82A	00010	00000	00000	10000	00001	10000	00000	00000	01000	00000
83A	00010	00000	00000	00000	00001	10000	00000	00000	01000	00000
84G	00010	00000	00000	10000	00001	10000	00000	00000	01000	00000
85A	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
85B	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
85D	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
85E	00001	10000	00000	00000	00001	10011	01010	00000	00000	00000
86A	10000	00000	00000	00000	00001	10011	01010	00110	00000	00000
86B	00000	00000	00000	01000	00001	10001	01000	00000	00000	00000

Fragments 1-50

87A	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
89A	10000	00000	00000	00000	00001	10001	01000	00000	00000	00000
89B	10000	00000	00000	00000	00001	10011	01100	00000	00000	10000
90A	00000	00000	00000	00000	00001	10011	01000	00000	00000	00000
90B	10000	10000	00000	00000	00001	10001	01010	00000	00000	00010
90C	00000	00000	00000	01000	00001	10001	01000	00000	00000	00000
92A	10000	00101	00100	01000	00001	10001	01000	00011	00000	00000
92B	10000	00000	00100	01001	00001	10001	01000	00010	00000	00000
93C	00000	00000	00000	00010	00001	10000	01000	00000	00000	00000
95A	10000	00000	00000	00000	00001	10011	01000	00000	00000	00000
96A	10000	00001	00100	01001	00000	00010	01000	00100	00001	10000
96B	10001	00000	00100	01111	00001	10011	01000	00000	00000	00000
98A	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
98B	00000	00000	00000	00000	00001	10101	01000	00000	00000	00000
98C	10001	00011	11100	00001	00001	00000	01000	00001	10000	10100
99A	00000	00000	00000	00000	00001	10000	01011	00000	00000	00000
99B	10000	10000	00000	00000	00001	10010	01001	00000	00000	00000
99C	00000	10000	00000	00000	00001	00100	01000	00010	00000	00000
100B	10100	00000	00000	01001	00001	10000	01000	00000	00000	00000
101A	00000	01000	00000	00000	00001	10001	01010	00000	00000	00000
102A	00000	00000	00000	00000	00001	10100	01010	00001	00000	00000
102B	00100	01000	00000	00000	00001	00010	01000	00000	00000	00000
102C	00000	00000	00000	00000	00001	10000	01011	00000	00000	00000
103A	00000	01000	00000	00000	00001	00000	01000	00000	00000	00000
103B	00000	00000	00000	00000	00001	10101	01010	00000	00000	00000
103C	00000	00000	00000	00000	00001	10001	01000	01000	00000	00000
104	00000	00001	00000	00000	00001	00010	01000	00000	00000	00000
105	00000	10000	00000	00000	00001	10111	01011	00000	00000	00000
106A	10000	00100	00000	00001	00011	00100	01010	00001	00000	00100
106B	00000	00000	00000	00000	00001	00111	01010	00000	00000	00000
106C	00000	01000	00000	00000	00001	10000	01010	00001	00000	00000
108A	00000	00000	00000	00000	00001	10010	01000	00000	00000	00000
108B	00000	00000	00000	00000	00001	10000	01010	10000	00000	00000
108C	00000	00000	00000	00000	00001	00000	01000	00000	00000	00000
110	01000	10000	00000	00000	00001	10110	01011	00000	00000	00000
111A	00000	00000	00000	00000	00001	10100	01010	00000	00000	00000
111B	00000	10000	00000	00000	00001	10111	01011	00000	00000	00000
111C	00000	10000	00000	00000	00001	10110	01001	00000	00000	00000
112	00000	00000	00000	00010	00001	10000	01000	00000	00000	00100
113A	00000	00000	00000	01001	00001	10000	01000	00001	00000	00100
114	00000	00000	00000	00000	00001	10000	01010	00000	00000	00000
116	10000	00000	00010	00011	00001	10000	01000	00000	00000	00000
118A	00000	00000	00000	00000	00001	10000	01000	00000	00000	00000
120C	00000	00000	00000	00000	00111	10000	00000	00101	10000	00100
121A	00000	00000	00000	00010	00001	10000	01000	00000	00000	00000
122A	00000	00000	00000	00000	00001	10000	01000	00000	00000	00000
123F	00000	00000	00000	00000	00001	10000	01000	00000	00000	00000

Fragments 1-50

123G	00000	00000	00000	00000	00001	10000	01000	00000	00000	00000
125A	00000	00000	00010	01000	00001	00000	01000	00001	00000	00100
126A	00000	00000	00000	00000	00001	10010	01000	00000	00000	00000
126E	00000	00000	00000	00010	00001	10000	01000	00000	00000	00000
127A	00000	00000	00000	01000	01001	10000	00010	00001	10000	00001
128A	00000	00001	00010	01000	00001	10000	01000	00001	00000	00100
130A	00000	00000	00000	10001	00001	10100	01000	00001	00000	00000
131	00000	00000	00000	00000	00001	10001	01000	00000	00000	00000
134A	00000	00000	00000	10000	00001	10100	01010	00001	10001	00000
HEN1	10000	00101	11000	01101	00011	00100	00001	00000	00000	00000
S3	00000	00000	00100	00000	00001	10001	01000	00000	00000	00000
S7	00000	00100	00000	00000	00001	10001	01000	00000	00000	00000

Fragment 51-100

1	00000	00000	00000	00000	01000	00010	00000	00100	00000	00000
3	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
5	00010	00000	01000	00000	00000	00010	00000	00100	00000	00000
8	00000	00000	01000	00000	00000	00010	00000	00100	00000	00000
10	00000	00000	01000	00000	00000	00010	00000	00100	00000	00000
12	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000
13	00100	00000	00000	00000	00000	00010	00000	00100	00000	00000
14	00010	00000	00000	00000	00000	00010	00000	00110	00000	00000
16	00000	00000	10000	00000	00000	00110	00000	01110	00000	00100
17	00000	00001	00000	00000	00000	00000	00000	00000	00010	00000
18	00000	00001	00000	00000	00001	01100	00000	00000	01100	00010
19	00000	00000	10000	00000	00000	00000	00010	00000	00000	00100
20	00000	00000	01000	00000	00000	00000	00000	00000	00000	00000
25	00000	00000	01000	00000	00000	00100	00000	00000	00000	00010
26	00000	00000	01000	00000	00000	01000	00000	00000	00000	00010
27A	00010	00000	00000	00000	10000	00010	00000	01110	00000	00100
27B	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000
27C	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
28A	00000	00000	10000	00000	00000	00010	00000	00100	00000	00000
29A	00010	00000	00000	00000	00000	00010	00000	00100	00000	00100
30A	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000
30D	00000	00000	00000	00000	00000	00010	00100	00100	00000	00000
30E	00000	00000	00000	00000	00000	00010	00000	00100	00000	00100
30F	00000	00000	00000	00000	00000	00010	00000	00100	00000	00100
31	10000	00000	00000	00000	00000	00010	00000	00100	00000	00000
32B	00000	00000	00000	00000	00000	00010	00000	00000	00000	00000
36B	00000	00000	00000	00000	00000	00000	00001	00000	00000	00000
37A	00000	00000	10000	00000	00000	00100	00000	00000	00001	10000
37B	00000	00000	10000	00000	00000	00100	00000	00000	00000	00000
37C	00000	00100	00000	00000	00000	00010	00000	01000	00000	00000
39	00000	00100	10000	00000	00000	00100	00000	00000	00000	00000
41A	00000	10000	00000	10000	00000	00000	00000	00000	00000	00100
41B	00000	00000	01000	00000	00000	00100	00000	00000	00000	00000
41C	00000	00000	10000	00000	00000	00100	00000	00000	00000	00000
43	00000	00000	01000	00000	00000	00110	00000	00000	00000	00000
45A	00000	00000	01000	00100	00000	00100	00000	00000	00000	00000
45B	00000	00010	00100	00001	10000	00100	00000	01000	00000	00000
45C	00000	00000	01000	00000	00100	00100	00000	00000	10000	00010
46A	00000	01000	10000	00000	00000	00100	00000	00000	00000	00000
46B	00000	01000	10000	00000	00000	00100	00000	00000	00000	00000
46C	00000	00000	10000	00000	00000	00100	00000	00000	00000	00000
48A	00000	00000	10000	00000	00000	00110	00000	00000	00000	00000
48B	00000	00000	10000	00000	00000	00100	00000	00000	00000	00010
48C	00000	00000	01000	00000	00000	00100	00000	00000	00000	00000
48D	00000	00000	01000	00000	00000	00110	00000	00000	00000	00010
50A	00000	00000	10000	00010	00000	00100	00000	00000	00000	00000
50B	00000	00000	10011	10000	00000	00110	00000	00000	00000	00000

Fragments 51-100

50C	00000	00000	10000	00000	00000	00100	00000	00000	00000	00000
50D	00000	00000	01000	10000	00000	00100	00000	01000	00000	00000
52B	00000	00000	10000	00000	10001	00000	00000	00000	00000	00000
54A	00100	00000	00000	00010	00000	00000	00001	00000	00000	00000
55A	00100	00000	00000	00000	00000	00000	00001	00000	00000	00000
57B	00000	00000	10000	00000	00000	00000	00001	00000	00000	00100
58B	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
58D	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
61	00000	00000	10000	00000	00000	00010	00000	00000	00000	00100
62D	00000	00000	10000	00000	10000	00000	00000	00000	00000	00000
63B	00000	00000	10000	00001	00000	00000	00000	00000	00000	00000
65C	00000	00000	10000	00001	00000	00000	00000	00000	00000	00100
65D	00000	00000	10000	00001	00000	00000	00000	00000	00000	00000
66C	00000	00000	10000	00001	00000	01000	00000	00000	00000	00010
66D	00000	00000	10000	00001	00000	01000	00000	00000	00000	00010
67A	00000	00000	00000	00000	00000	01000	00000	10001	00000	00010
67B	00000	00000	01000	00000	10000	01000	00000	00000	00100	00010
68A	00000	00000	00000	00000	11000	10010	00000	00100	00000	00000
68B	00000	00000	00000	00000	01000	00010	00000	00100	00000	00000
68C	00000	00000	00000	00000	01000	00010	00000	00100	00000	00000
69A	00000	00000	01000	00000	00000	01000	00000	00000	00000	00010
69B	00000	00000	01000	00000	00000	01000	00000	00000	00000	00010
70A	00000	00000	00000	00000	01100	00010	00000	00100	00000	00000
70B	00000	00000	00000	00000	01000	00010	00000	00100	00000	00000
71A	00000	00100	01000	00000	00000	01000	00000	00000	00000	00010
72A	00000	00000	00000	00000	01000	00010	01000	00100	00000	00000
72B	00000	00000	00000	00000	01000	00010	01000	00100	00000	00000
74D	00100	00000	00000	00000	00000	00001	00000	00000	00000	00000
75A	00000	00000	00000	00000	00000	00000	01000	00000	00000	00000
76A	00000	00000	00000	00000	00000	10000	00000	01000	00000	00010
76E	00000	00000	01000	00001	00000	10000	00000	01000	00000	00000
77B	00000	00000	01000	00001	00000	01000	00000	00000	00000	00000
78A	00000	00000	01000	00000	00000	10000	00000	01000	00000	00010
79	00000	00000	01000	00000	00000	01000	00000	00000	00000	00010
80A	00000	00000	00000	00000	01000	00010	00000	00100	00000	00000
80C	00000	00000	00000	00000	01100	00010	00000	00100	00000	00000
80D	00000	00000	00000	00000	01000	00010	00000	00100	00000	00000
81A	00000	00000	00000	00000	00000	10000	00000	01000	00000	00010
82A	00000	00000	10000	00001	00000	01000	00000	00000	00000	00010
83A	00000	00000	10000	00000	00000	01000	00000	00000	00000	00010
84G	00000	00000	10000	00000	00100	01000	00000	00000	00000	00010
85A	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
85B	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
85D	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
85E	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
86A	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
86B	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000

Fragments 51-100

87A	00000	00000	01000	00000	00000	00010	00000	00100	00000	00100
89A	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000
89B	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000
90A	00000	00000	00000	00000	00000	00010	00000	00100	00000	00100
90B	00000	00000	00000	00000	00000	00000	00000	00100	00000	00000
90C	00010	00000	00000	00000	00000	00010	00000	00100	00000	00100
92A	00000	00000	00000	00000	00000	00011	00000	00000	00000	00010
92B	00000	00000	00000	00000	00001	00010	00000	00100	00000	00000
93C	00000	00000	01000	00000	10000	00010	00000	00000	00000	00000
95A	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000
96A	00000	00000	00000	01000	00000	00000	10000	00000	00000	00000
96B	00000	01001	00000	00000	00000	00110	00010	00000	00000	00010
98A	00000	00001	01000	00000	00000	00100	00000	00000	00000	00010
98B	00000	00001	10000	00000	00000	00100	00000	00000	00000	00010
98C	00000	00000	10000	00000	00000	00000	01000	00000	00000	00001
99A	00000	00000	10000	00000	00000	00100	00000	00000	00000	00010
99B	00000	00000	01000	00000	00000	00100	00001	00000	00000	00010
99C	00000	00000	01000	00000	00000	00100	00000	00000	00000	00010
100B	00000	00000	00000	00010	00000	00000	00001	00000	00000	00000
101A	00000	00010	10000	00000	00000	00110	00000	00000	00000	00010
102A	00000	00000	10000	00000	00001	00100	00000	01000	00000	00000
102B	00000	00000	10000	00000	00000	00100	00000	00000	00000	00010
102C	00000	00000	10000	00000	00000	00100	00000	00000	00000	00010
103A	00000	00000	10000	00000	00000	00110	00000	00000	00000	00010
103B	00000	00000	10001	00000	00000	00100	00000	00000	00000	00010
103C	00000	01000	10000	00000	00000	00110	00000	00000	00000	00010
104	00000	00000	10000	00000	00000	00100	00000	00000	00000	00010
105	00000	00000	10000	00000	00000	00100	00001	00000	00000	00010
106A	00000	00000	01000	00000	00000	00100	00000	00000	00000	00010
106B	00000	00001	01000	00000	00000	00100	00000	00000	00000	00000
106C	00000	00000	10000	00000	00000	00100	00001	00000	00000	00000
108A	00001	00000	10000	00001	00000	00100	00000	00000	00000	00000
108B	00000	00010	10000	00000	00000	00100	00001	00000	00000	00010
108C	00000	10010	01000	00000	00000	00100	00001	00000	00000	00010
110	00000	00000	10010	00000	00000	00100	00000	00000	00000	00010
111A	00000	00000	01000	00000	00000	00100	00000	00000	00000	00010
111B	00000	00000	10000	00000	00000	00100	00000	00000	00000	00010
111C	00000	00000	00000	00000	00000	00100	00000	00000	00000	00010
112	10000	00000	00000	00000	10000	00000	00000	00010	10000	00000
113A	00000	10001	00000	10000	00000	00000	00010	00000	00100	01000
114	00000	00000	01000	00000	00000	00100	00000	00000	00000	00000
116	00000	00000	00000	00001	00000	00000	00000	00000	00000	00100
118A	00000	00000	10000	00001	00000	00000	00000	00100	00000	00100
120C	01001	00001	00000	00001	00010	00010	00010	00100	00010	00000
121A	01000	00000	10000	00000	00010	00000	00000	00000	00000	00000
122A	00000	00000	01000	00000	10000	00000	00000	00000	00000	00100
123F	00000	00000	01000	00000	00010	00000	00000	00000	00000	00000

Fragments 51-100

123G	00000	00000	01000	00000	00000	00000	00000	00000	00000	00000	00000
125A	00000	01000	00000	10000	00000	00000	00010	00000	00100	00000	00000
126A	00000	00000	00000	10000	00000	00000	00000	00100	00000	00010	00000
126E	00000	00000	01000	00000	10000	00010	00000	00000	00000	00000	00000
127A	00001	00010	00000	00000	00000	00000	01000	00000	10010	00000	00000
128A	01000	00010	00000	00001	00000	00000	00100	00000	10001	00000	00000
130A	01000	00000	01000	00000	10000	00100	00000	00000	00000	00000	00000
131	00000	00000	01000	00000	00000	00010	00000	00100	00000	00000	00000
134A	10000	00001	01000	10000	10000	00010	00100	00000	00100	01000	01000
HEN1	10000	00000	00010	00000	00000	00000	00000	00000	00000	00000	00000
S3	00010	00000	00000	00000	00000	00010	00000	00100	00000	00000	00000
S7	00000	00000	00000	00000	00000	00010	00000	00100	00000	00000	00000

Fragments 101-150

1	00000	00000	00000	00001	00000	00000	00000	00000	01010	00010
3	00000	00000	00000	00000	00000	00000	00000	00001	01000	10010
5	00000	10000	00000	00010	00000	00000	00000	00000	00100	10010
8	00000	10000	00000	00000	00000	00000	00000	00000	00100	10010
10	00000	10000	00000	00000	00000	00000	00000	00000	00100	10010
12	00000	11000	00000	00000	00000	00000	00000	00000	00100	10010
13	00000	00000	00000	00001	00000	00000	00010	00010	00010	01011
14	00000	10000	00000	00000	00000	00000	00000	00000	00100	10010
16	00000	01000	00000	00001	00000	00000	00000	00000	00010	01010
17	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
18	00000	00010	00000	00000	00000	00010	00000	00010	10110	00000
19	00001	00001	00000	00001	00100	00000	00000	00001	00110	00100
20	00001	00000	00000	00000	00100	00000	00000	00001	00010	10100
25	00010	00001	00000	00000	00001	01000	00000	00100	00011	11011
26	00000	00000	00000	00000	00000	00000	00000	00001	00010	10010
27A	00000	10000	00000	00001	00000	00000	00001	00000	01010	01010
27B	00000	00000	00000	00001	00000	00000	00001	00000	00010	01011
27C	00000	10000	00000	00000	00000	00000	00000	00000	00010	01011
28A	00000	10000	00000	00000	00000	00000	00000	00000	00010	11011
29A	00000	00000	00000	00000	00000	00100	00010	00000	00011	00010
30A	00000	00000	00000	00000	00000	00000	00000	00000	00010	01010
30D	00000	00000	00000	00000	00000	00000	01000	00000	00011	10011
30E	00000	00000	00000	00001	00000	00000	00000	01000	00010	01010
30F	00000	00000	00000	00000	00000	00000	01001	00000	00010	00011
31	00000	00000	00000	00001	00000	00000	01001	00000	00010	11011
32B	00000	01000	00000	00001	00000	00000	01010	00010	00010	01010
36B	00000	00000	00010	00000	00000	00000	00000	00000	00100	00100
37A	00000	00001	10000	10000	00000	01000	00000	00000	00011	00010
37B	00000	00001	00000	10100	01000	01000	00010	00000	00011	00010
37C	00000	00001	00000	00000	00000	00000	01000	00000	00011	00000
39	00000	00001	00000	10000	00001	10000	00001	00000	00011	01010
41A	11100	00000	00000	00000	00000	01000	01000	00000	00110	11010
41B	00000	00000	00000	00000	00000	01000	00001	00000	00111	10010
41C	00000	00001	00000	00000	00000	01000	00000	00100	00001	11010
43	00000	00001	00000	00100	00000	01001	00000	00100	00010	00010
45A	00000	00000	00000	00100	00000	00000	01000	10000	00011	01010
45B	10000	00001	00000	10100	01000	01000	01000	00000	00001	01010
45C	10000	00001	00000	11000	00001	01000	00000	00000	00100	10010
46A	00000	00001	10000	00100	00000	01000	00000	00000	00010	01010
46B	00000	00001	00000	10100	00001	01000	01000	00000	00011	00010
46C	00000	00001	00000	10100	00000	01000	01000	00000	00011	01010
48A	00000	00001	00000	10100	01001	10000	00000	00000	00011	11010
48B	10000	00001	00000	10100	01001	00000	00000	00000	00001	01011
48C	00000	00001	00000	00001	00001	00000	00000	00000	00001	01010
48D	10000	00001	00000	10100	00001	00000	00000	00000	00010	01010
50A	00000	00001	00000	10000	00000	01000	00000	00000	00011	01010
50B	00000	00001	00000	00000	00001	01000	00000	00100	00011	01010

Fragments 101-150

50C	00000	00000	00000	00000	00001	01000	00000	00100	00011	00010
50D	00000	00001	00000	00000	00001	01000	00000	00000	00010	00000
52B	00001	00000	00000	00010	00100	00000	00000	10001	00011	00010
54A	00000	00000	00000	00000	00000	00000	00000	00000	00010	00100
55A	00000	00000	00000	00000	00000	00000	00000	00000	00000	00111
57B	00001	00000	00000	00001	00100	00000	00000	00000	00010	10010
58B	00000	01000	00000	00000	00000	00000	00000	00000	00100	00100
58D	00000	01000	00000	00000	00000	00000	00000	00000	00010	10011
61	00001	00000	00000	00000	00000	00100	00000	00000	00000	00011
62D	00000	00000	00000	00000	00000	00000	00000	00000	00010	01010
63B	00000	00000	00000	00000	00010	00000	00000	00000	00010	00010
65C	00000	00000	00000	00000	00010	00000	01000	00000	00110	10010
65D	00000	00000	00000	00000	00100	00000	00000	00000	00110	10010
66C	00000	00000	00000	00000	00100	00000	00000	10000	10010	10010
66D	00000	00000	00000	00000	00000	00000	00000	00000	00010	10010
67A	00000	00000	00000	00000	00000	00000	00000	00000	00000	11010
67B	00000	00000	00000	00001	00000	00000	00000	00000	10110	11010
68A	00000	00000	00000	00001	00000	00000	00000	00000	00011	11010
68B	00000	00000	00000	00001	00000	00100	00011	00000	00010	01011
68C	00000	00000	00000	00000	00000	00000	00000	00000	00010	00010
69A	00000	00000	00000	00000	00000	00000	00000	00001	10000	10010
69B	00000	00000	00000	00001	00000	00000	00000	00000	10010	10010
70A	00000	00000	00000	00000	00000	00000	00000	00000	00010	01011
70B	00000	00000	00000	00000	00000	00000	00000	00000	00010	00011
71A	00000	00000	00000	00000	00000	00000	00000	00000	00010	10010
72A	00000	00000	00000	00000	00000	00000	00000	00000	00011	01010
72B	00000	00000	00000	00001	00000	00000	00000	00000	00011	01010
74D	00000	00000	00100	00000	00000	00100	00000	00000	00000	00110
75A	00000	00000	00000	00000	00000	00000	01000	00000	00000	00011
76A	00000	00000	00000	00001	00000	00000	00000	00000	00000	00000
76E	00000	00000	00000	00000	00000	00000	00000	00000	00111	11011
77B	00000	00000	00000	00001	00000	00000	00000	00001	00011	01011
78A	00000	00000	00000	00000	00000	00000	00000	00000	00100	11010
79	00000	00000	00000	00000	00000	00000	00000	00000	00010	10010
80A	00000	00000	00000	00001	00000	00000	00000	00000	00010	01010
80C	00000	00000	00000	00001	00000	00000	00010	00000	00010	10011
80D	00000	00000	00000	00001	00000	00000	01000	00000	00010	01011
81A	00000	00000	00000	00000	00000	00000	00000	00000	00100	11010
82A	00000	00000	00000	00001	00000	00000	00000	00000	00010	10010
83A	00000	00000	00000	00000	00000	00000	00000	00000	00010	10010
84G	00000	00000	00000	00000	00000	00000	00000	10000	10010	10010
85A	00000	10000	00000	00000	00000	00000	00000	00000	00010	01001
85B	00000	01000	00000	00000	00000	00000	00000	00000	00010	00010
85D	00000	00100	00000	00000	00000	00000	00010	00000	00010	00010
85E	00000	10000	00000	00000	00000	00000	00010	00000	00010	00010
86A	00000	10000	00000	00000	00000	00000	01010	00000	00010	01010
86B	00000	00000	00000	00000	00000	00000	00000	00000	00010	01011

Fragments 101-150

87A	00000	10000	00000	00000	00000	00000	00000	00000	00000	00000	10011
89A	00000	10000	00000	00000	00000	00000	00000	01010	00000	00010	01010
89B	00000	10000	00000	00000	00000	00000	00000	01000	00000	00010	01011
90A	00000	10000	00000	00000	00000	00000	00000	01010	00000	00010	00010
90B	00000	00000	00000	00000	00000	00000	00000	00000	00000	00010	01010
90C	00000	00000	00000	00000	00000	00000	00000	00000	00000	00010	01010
92A	00000	00000	00000	00000	00000	00000	00000	01010	00000	00010	00011
92B	00000	10000	00000	00000	00000	00100	00000	01010	00000	00010	10011
93C	00001	00000	00000	00001	00010	00000	00000	10001	00010	11010	
95A	00000	00000	00000	00000	00000	00000	00000	01011	10000	00010	00011
96A	00001	00000	00000	00000	00000	00000	00000	01010	00000	00010	01110
96B	00000	00001	00000	00001	00000	01000	01010	00000	10010	00011	
98A	00000	00001	00000	10000	00000	00000	01010	00000	00010	10000	
98B	00000	00001	00000	10000	00000	01000	01000	00000	00010	01010	
98C	00000	00001	00001	00000	00000	01000	01010	00000	00010	11000	
99A	00000	00001	00000	00000	00000	01000	00000	00000	00010	00010	
99B	00000	00001	00000	00000	00000	01000	00001	00000	00001	00110	
99C	00000	00001	00000	00000	00000	01000	00000	00000	01010	01010	
100B	00000	00000	00010	00000	00000	00000	00000	00001	00000	10010	
101A	00000	00001	00000	00000	00000	11000	00000	00000	00010	01010	
102A	00000	00001	00010	00000	00000	01000	00000	00000	00010	01010	
102B	00000	00001	00000	00000	00000	10000	00000	00000	00010	01010	
102C	00000	00001	00000	00000	00000	01000	00010	00000	00011	01010	
103A	00000	00001	00000	00000	00001	00000	00000	00000	00010	01010	
103B	00000	00001	00000	00000	00000	11000	00000	00000	00010	11011	
103C	00000	00001	00000	00000	00000	01000	01000	00000	00010	01010	
104	00000	00001	00000	10001	00000	10000	00000	00000	00010	01010	
105	00000	00001	10000	10000	00000	01000	00000	00000	00011	01010	
106A	00010	00001	10000	00000	00000	01000	01001	00000	00000	01010	
106B	00000	00001	00000	00100	00001	01000	00000	00000	00011	01010	
106C	00000	00001	00000	00000	00001	01000	00000	00000	00001	01010	
108A	00000	00001	00000	00000	00000	01000	00000	00000	00010	01010	
108B	00000	00001	00000	00000	00000	01000	00000	00100	00010	01010	
108C	00000	00101	00000	10000	00000	01000	00000	00000	00000	01010	
110	00000	00001	00000	00000	00000	01000	00010	00100	00011	01010	
111A	00000	00001	00000	00000	00000	01000	00100	00000	00010	00010	
111B	00000	00001	00000	00000	00000	01000	01000	00000	00010	11010	
111C	00000	00001	00000	00000	00000	01000	00000	00000	00011	01010	
112	00000	00000	00000	00001	00100	00000	00000	00001	00110	00010	
113A	00000	00000	00000	00000	00000	00000	10000	00001	00110	00010	
114	00000	00001	00000	10000	00000	01000	00000	00000	00010	00010	
116	00001	00000	00000	00001	00000	00000	00000	00000	00000	00010	
118A	00000	00000	00000	00001	00000	00000	00000	00000	00010	11010	
120C	00000	00000	00000	00000	00000	00000	00000	00000	00010	10010	
121A	00001	00000	00000	00000	00100	00000	01000	00000	00110	10010	
122A	00001	00000	00000	00000	00010	00000	00000	00001	00010	11010	
123F	00001	00000	00000	00001	00100	00000	00000	00001	00010	00010	

Fragments 101-150

123G	00001	00000	00000	00000	00100	00000	00000	00001	00011	10010
125A	00000	00000	00000	00000	00000	00000	10000	00001	00010	00010
126A	00001	00000	00000	00001	00100	00000	00000	00001	00110	00010
126E	00001	00000	00000	00000	00100	00000	00000	00000	00000	00000
127A	00000	00000	00000	00000	00000	00000	10000	00000	00000	00001
128A	00000	00000	00000	00100	00010	00000	00010	00000	00110	10000
130A	00000	00000	00000	00001	00010	00000	00000	00000	00000	00001
131	00000	10000	00000	00000	00000	00000	00000	00000	00011	01010
134A	00010	00000	00000	00000	00000	00000	10000	00000	00000	00000
HEN1	00000	00000	00000	00000	00000	00000	01010	00000	00111	01000
S3	00000	00000	00000	00000	00000	00000	00000	00000	00010	11010
S7	00000	10000	00000	00000	00000	00000	00000	00000	00010	01010

Fragments 151-200

1	00100	10000	00000	01010	00000	00011	00000	10100	00001	00100
3	00100	00000	00000	00010	00000	00011	00000	10100	00000	00000
5	00000	00000	00000	00000	10000	00011	00000	00000	00000	00100
8	00100	00000	00000	00000	10000	00001	00000	00101	00000	00000
10	00000	00000	00000	00000	10000	00001	00100	10000	00000	00000
12	00000	00000	00000	00000	10000	00001	00000	00000	00010	00100
13	00011	00000	10000	00010	10010	01001	11000	10100	00010	10000
14	00000	00000	00000	00000	10000	00001	00000	00000	00000	10000
16	11001	01000	00010	00110	10010	00011	01000	00101	00000	00010
17	00100	00000	00000	00000	00000	00000	00100	00000	00000	00000
18	01000	10000	00000	00000	00000	00011	10010	00010	00000	00000
19	00100	00010	00000	00000	10000	00110	01000	00100	00000	00000
20	00100	00000	00000	00010	10000	00011	01000	00000	00000	00000
25	00100	00000	00000	00100	10010	10011	01000	00000	00000	00110
26	00000	00000	00000	00010	10100	00011	00000	00100	00000	00010
27A	10111	00000	00000	00010	10010	00001	00000	10100	00000	00010
27B	00101	00000	00000	00010	10010	00001	00000	10100	00000	00010
27C	00101	00000	00000	00010	10000	00001	00000	10000	00000	00010
28A	00000	00000	00000	00000	10000	00001	00000	10000	00000	00000
29A	10001	00000	00000	00010	10010	10001	00100	00100	00010	00010
30A	00000	00110	00000	00010	10000	00001	00100	00001	00000	10010
30D	00100	01000	01000	00010	10010	00001	01000	01000	10010	10000
30E	00111	01010	00000	00010	10000	01001	00000	00101	00010	10000
30F	10101	00100	10000	00010	10000	00001	00110	00100	00010	10010
31	00000	01100	01000	01010	10000	01001	00100	00000	00000	00010
32B	00001	00000	00000	00010	10000	01001	01000	00100	00000	10000
36B	00000	11001	00000	00000	00000	00001	10010	00000	00000	00000
37A	00000	00000	00000	00100	10010	00000	00010	01000	00000	00110
37B	10000	00100	00000	00100	10010	00001	00100	01001	00000	00110
37C	00100	00100	00010	00100	11010	00011	00000	01001	00000	00010
39	00100	00000	00000	00100	10010	10001	00000	01001	00010	00110
41A	00000	01000	00100	00100	10010	00001	10000	00000	00000	00110
41B	00100	00000	00000	00100	10000	00011	00000	00000	00000	00100
41C	00100	00000	00000	00100	10010	00011	01000	00000	00000	00100
43	00100	00000	00000	00100	10010	00011	00000	01001	00000	00100
45A	00000	00000	00000	10100	11010	00101	10010	01001	00010	00100
45B	10000	00010	00000	00100	10010	00010	11010	01000	10000	00100
45C	00100	00000	00000	00000	10010	00011	00000	00001	00000	00100
46A	00100	00010	00000	00100	10010	00011	00000	00001	00000	00110
46B	00100	00000	00000	00100	10010	00011	00000	01001	00000	00110
46C	00000	00000	00000	00100	10010	10001	00000	01001	00000	00010
48A	00000	01000	00000	00100	10010	10011	00000	01000	00000	00110
48B	11101	01000	00000	00100	10010	10011	01010	00001	10100	10110
48C	00100	00000	00000	00100	10010	00011	00000	00000	00000	00110
48D	00000	00000	01000	00100	10010	01001	00010	00000	00010	00100
50A	00100	00010	00000	00100	10010	00011	00000	00000	00000	00100
50B	00100	00010	00000	00100	10010	00011	00000	00001	00000	00100

Fragments 151-200

50C	00100	00000	00000	00100	10010	00011	00000	00000	00000	00100
50D	00100	00000	00000	00000	10010	00011	00000	00000	00000	00000
52B	00100	00010	00000	00000	10000	00010	01000	00000	00000	00000
54A	00100	00001	00000	00000	10010	00011	10010	00000	00000	00000
55A	00100	10001	00001	00000	10010	00011	10010	00000	00000	00000
57B	00000	00010	00000	00010	10000	00010	11000	00000	00000	00000
58B	10100	00100	00000	00010	10000	00011	01000	00100	00010	00000
58D	00111	00110	00000	00010	10000	11001	11100	00101	00010	00000
61	00100	00000	00000	00000	10000	00011	01000	00000	00000	00000
62D	00000	00000	00000	00000	10000	00011	01000	00001	00000	00000
63B	00000	00000	00000	00000	10000	00011	01000	00000	00000	00000
65C	00100	00000	00000	00000	10000	00011	01000	00000	00000	00000
65D	00100	00000	00000	00000	10000	00011	01000	00000	00000	00000
66C	00000	10001	00000	00010	10100	00010	00010	00000	00000	00010
66D	00000	00000	00000	00010	10100	00011	00000	00000	00000	00010
67A	00000	00000	00000	00010	00100	00011	00001	00010	00000	01010
67B	10101	00000	00000	00011	00100	00011	00000	00010	00000	00010
68A	00100	00000	00000	00000	10000	00011	00000	00001	00001	00000
68B	00000	00100	00000	00010	10000	00001	00000	10000	00000	00010
68C	00111	00000	10000	00000	10000	01001	10000	10100	00000	10000
69A	00100	00000	00000	00010	10000	00011	00000	00000	00000	00000
69B	10001	00000	00000	00010	10100	00011	00000	00010	00000	01000
70A	00000	00000	10000	00010	10000	11001	00100	10000	00000	00010
70B	01111	01000	00000	00010	10000	11011	00000	00101	00010	00010
71A	00000	00000	00000	00010	10100	00011	00000	00000	00000	00000
72A	00001	00000	10000	00010	10000	00001	10000	10000	00000	00010
72B	00001	00000	10000	00010	10000	00001	10000	10000	00000	00010
74D	00000	00001	00000	00010	10010	00011	10010	00000	00000	00000
75A	00111	00000	10100	00010	10010	00011	10100	00000	00000	10000
76A	00100	00000	00000	00000	00000	00000	00000	00010	00000	00000
76E	00000	00000	00000	00001	00000	00011	00000	00010	00000	00000
77B	00000	00000	00000	00010	10100	00011	00000	00010	00000	00010
78A	00100	00000	00000	00001	00000	00011	00000	00010	00000	00000
79	00000	00000	00000	00010	10000	00011	00000	00000	00000	00000
80A	00000	00000	00000	00010	10000	00001	00000	00000	00000	00000
80C	00111	01100	00000	00010	10000	11001	11000	10100	00010	10110
80D	10111	01100	10000	00010	11001	11001	01000	10001	00000	00000
81A	00000	00000	00000	00001	00000	00001	00001	00010	00000	00000
82A	00000	00000	00000	00010	10000	00011	00000	00000	00000	00000
83A	00000	00000	00000	00010	10000	00011	00000	00000	00000	00000
84G	00000	10000	00000	00010	10100	00011	00000	00010	00000	00000
85A	00001	00000	00000	00000	10000	00001	00000	00001	00000	00010
85B	00001	00000	00000	00010	10010	01001	01000	00001	00000	00010
85D	10111	01000	00000	00010	10010	11001	01000	00101	00000	00010
85E	00001	01000	00000	00010	10010	01001	01000	00101	00000	00010
86A	00001	00100	00000	00010	10010	00001	00000	00100	00000	00010
86B	00000	00100	00000	00000	10000	00001	00000	10000	00000	00010

Fragments 151-200

87A	00100	00000	00000	00010	10000	00011	00000	00001	00000	00010
89A	00101	00000	00000	00010	10010	01001	00000	10100	00000	10000
89B	00000	00100	00000	00010	10000	00001	00000	10000	00000	00010
90A	00000	00000	00000	00010	10010	01001	11000	10100	00000	00010
90B	00000	00000	10000	00010	10000	00001	01000	00000	00000	00010
90C	00100	00000	00000	00000	10000	00001	00100	00001	00000	00000
92A	00000	00100	10000	00100	00001	00001	01010	10000	00000	10000
92B	11000	00100	00000	00010	10001	11011	10000	10000	00010	10000
93C	00100	10000	00000	00000	10000	00011	00000	00000	00000	00010
95A	00101	00100	00000	00010	10010	01001	10000	10000	00000	00010
96A	10100	00100	00000	00010	00000	00001	00000	00000	00000	00000
96B	01100	00000	00000	00010	10000	00001	01000	00001	00010	00010
98A	00000	00000	00000	00000	10100	00011	01000	00000	00100	00110
98B	00000	00000	00000	00100	10010	10001	00100	00001	00010	00010
98C	00100	01010	00000	00000	11010	00011	01000	00000	00100	00110
99A	01100	01000	00000	00100	10010	00011	01000	00001	00000	00111
99B	00100	00010	00000	00100	10010	01011	00100	00000	00000	00010
99C	00000	00000	01000	00100	10010	00001	00000	00000	00000	00010
100B	00100	01000	00000	00000	10010	10011	10001	00000	00000	00000
101A	00100	00000	00000	00100	10010	00011	01000	00000	00000	00110
102A	00100	00000	10000	00101	10010	00011	11010	00001	00000	00010
102B	00100	00000	00000	00100	10010	00001	00000	00000	00000	00010
102C	00100	00000	00000	00100	10010	00011	01000	01000	00000	00110
103A	00100	00000	00000	00100	10010	00011	00000	00000	00000	00100
103B	00000	00000	00000	00100	10010	00011	01000	00000	00000	00100
103C	00100	00000	00000	00100	10010	10011	01000	00001	00000	00110
104	00000	00000	00000	00010	10010	00011	00000	00000	00000	00100
105	00100	00000	00000	00000	10010	10010	00000	00000	00000	00111
106A	00101	00000	00000	00100	10010	01001	00110	00000	00000	00100
106B	00000	00000	01000	00100	10010	00011	00000	00000	00000	00110
106C	00000	01000	00000	00100	10010	00011	11001	00000	00000	00100
108A	00100	00000	00000	00001	10010	00001	01010	01001	00000	00110
108B	00100	00000	00000	00100	10010	00001	01000	00000	00000	00110
108C	00000	00000	00000	00100	10100	00011	00000	00001	00000	00110
110	00000	00000	00000	00100	10010	00001	01000	01001	00000	00110
111A	00100	00010	00000	00100	10100	00011	11000	00000	00010	00110
111B	00100	00000	00000	00100	10010	00001	01000	00000	00000	00010
111C	00100	00000	00000	00000	10010	00010	01000	01001	00000	00111
112	00100	00010	01010	00010	10000	00011	01000	00000	00000	00110
113A	00100	00010	01010	00010	10000	00011	01000	00000	00000	00010
114	00101	00000	01000	00100	10010	00001	00100	00001	00000	00010
116	00100	00010	00000	00000	00000	00011	01000	00000	00000	00000
118A	00000	00010	01000	00010	00000	00011	00000	00000	00000	00000
120C	00110	00010	01010	00010	10000	00111	01100	00000	00000	10000
121A	00100	00000	00000	10000	10000	00011	01000	00000	00000	00000
122A	00000	00000	00000	00000	10000	00011	01000	00001	00000	00000
123F	00000	00010	00000	00000	10000	00011	01000	00000	00000	00000

Fragments 151-200

123G	00000	00010	00000	00000	10000	00011	01000	00000	00000	00000
125A	00000	00010	00000	00000	10000	00011	01000	00000	00000	00000
126A	00000	00000	10000	00010	10000	00010	00000	00000	00000	00000
126E	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
127A	00110	00110	01010	10010	11000	00111	00100	00000	00000	00000
128A	00110	00000	01010	00010	11000	00011	00100	00000	00000	10000
130A	00100	00000	01010	00000	00000	00011	01100	00000	00000	10000
131	00000	00000	00000	00000	10000	00011	00000	10001	00000	00000
134A	10101	00000	00000	00000	10100	00011	00000	00000	00000	00000
HEN1	00100	00100	00100	00000	10000	00000	00000	00000	00000	00011
S3	00000	00000	00000	00000	10000	00001	00000	10000	00000	00000
S7	00100	00000	00000	00000	10000	00001	01000	00100	00000	00100

Fragments 201-250

1	00000	01010	00000	00100	00010	00010	00000	00001	00000	00001
3	00000	01010	00000	00000	00010	00010	00000	00001	00000	00001
5	00000	10010	01000	00000	00000	00010	00000	00101	00100	00001
8	00000	10010	01000	00000	00000	01010	00000	00011	00001	00000
10	00000	10010	01000	10000	00000	00010	00001	00001	00100	00001
12	00000	01010	00000	00000	00011	00010	00000	00001	00000	00001
13	00000	01010	00000	10100	00010	00010	00000	10001	00000	00000
14	00000	01010	00000	00000	00011	00010	00000	00001	00000	00001
16	00010	01010	00000	10100	00011	00010	00000	10001	00000	00011
17	00000	00010	00000	00000	00011	00000	00000	00000	00000	00000
18	00001	00100	00000	00000	00000	00000	00000	00000	00000	00000
19	00010	00100	00000	00000	00000	00010	00000	00000	00000	00000
20	00010	10100	00000	00000	00000	00010	00000	00000	00000	00000
25	00000	10010	00000	00000	00000	00010	00000	10000	00010	00000
26	00001	00100	00000	00000	00000	00100	00000	01001	00000	00000
27A	00000	01000	00000	10100	10010	00010	00000	10011	00000	00001
27B	00000	01010	00000	10100	10010	00010	00000	10001	00000	00100
27C	00000	01010	00000	00000	00010	00010	00000	00001	00100	00000
28A	00000	01000	01000	00000	00000	00010	00000	00001	00000	00001
29A	00000	01010	00100	00000	00010	00010	00000	10001	00000	00000
30A	00000	01010	00000	00000	00010	00010	00000	10001	00100	00001
30D	00000	01010	00000	00000	00010	00011	00000	10001	00000	00000
30E	00000	01000	00010	00000	00010	00010	00000	10011	00000	00011
30F	00000	01010	01000	00010	00010	00010	00001	10001	00100	00011
31	00000	01000	01000	00000	00010	00010	00000	10001	00100	00000
32B	00000	01010	00000	10000	00011	00010	00000	00001	00100	00000
36B	00000	00000	00000	00000	00000	00100	00100	00000	00000	01000
37A	00000	10110	00000	01000	00000	00010	00000	10000	00010	00000
37B	00000	10110	00000	01000	00000	00010	00000	10000	00010	00000
37C	00000	10000	00000	01000	00000	00010	00000	10000	00010	00000
39	00000	10110	00000	01001	00000	00010	00000	10000	00010	00000
41A	10000	10110	00000	01011	00000	00010	10000	10000	00010	00000
41B	00000	10100	00000	00000	00000	00010	00000	10000	00010	00001
41C	00000	10010	00000	00000	00000	00010	00000	10000	00010	00001
43	00000	10110	00000	01000	00000	00010	00000	10000	00010	00010
45A	00000	10100	10000	01000	01000	00010	10000	10000	01000	00011
45B	00000	10110	00000	01000	00000	00010	00000	10000	00010	00000
45C	00000	10110	00000	00000	00000	00010	00000	10000	00010	00001
46A	00001	10100	00000	00000	00000	00010	00000	10000	00010	00000
46B	00000	10100	00000	01000	00000	00010	00010	10000	00010	00001
46C	00000	10010	10000	01001	00000	00010	00000	10000	00010	00000
48A	00000	10010	00000	00000	00000	00010	00000	10000	00010	00011
48B	00000	10010	00000	01001	00000	00010	00000	10000	00010	00001
48C	00000	10110	00000	00000	00000	00010	00000	10000	00010	00000
48D	00000	10010	00000	01000	00000	00010	00000	10000	00010	00011
50A	00000	10110	00000	01000	00000	00010	00010	10000	00010	00010
50B	00000	10010	00000	01000	00000	00010	00010	10000	00010	00010

Fragments 201-250

50C	00000	10010	00000	00000	00000	00010	00010	10000	00010	00010
50D	00000	10010	00000	00000	00000	00000	00000	10000	00000	00000
52B	00000	10100	00001	00000	00000	00010	00000	00001	00000	10000
54A	00000	10100	00000	00000	00001	00100	00100	00000	00000	00000
55A	00000	10000	00000	00000	00001	00100	00100	00000	00000	00000
57B	00000	01010	00000	00000	00000	00010	00000	00001	00000	00000
58B	00000	10100	00000	00000	00011	00010	00000	00011	00000	00001
58D	00100	01010	00000	10000	00010	00010	00000	10001	00000	00001
61	00000	01010	00000	00000	00000	00010	00000	00001	00000	00000
62D	00000	01010	00000	00000	00000	00010	00000	00001	00000	00010
63B	00000	01010	00000	00000	00000	00010	00000	00001	00000	00010
65C	00000	10100	00000	00000	00000	00010	00000	00000	00000	00010
65D	00000	10100	00000	00000	00000	00010	00000	00000	00000	00000
66C	00001	00100	00000	00000	00000	00100	00000	00001	00000	00000
66D	00001	00100	00000	00000	00010	00100	00000	00001	00000	00000
67A	00011	00100	00000	00000	00000	00100	01000	01001	00000	00000
67B	00011	00100	00000	00000	00010	00100	01000	00001	00000	00000
68A	00000	01010	00000	00000	00010	00010	00000	00001	00000	00001
68B	00000	01000	00000	10100	00010	00010	00000	10001	00000	00001
68C	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
69A	00001	00100	00000	00000	00000	00000	00000	00001	00000	00000
69B	00001	00100	00000	00000	00000	00100	00000	00001	00000	00000
70A	00000	01000	00000	10000	00010	00010	00000	00001	00000	00001
70B	00000	01010	00000	10000	00010	00010	00000	10001	00100	00001
71A	00001	00100	00000	00000	00010	00100	00000	00001	00000	00000
72A	00000	01010	00000	00000	00010	00010	00000	10001	00000	00001
72B	00000	01010	00000	00000	00010	00010	00000	00000	00000	00001
74D	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
75A	00000	00000	00000	00000	00001	00000	00000	00000	00000	00000
76A	00011	00100	00000	00000	00010	00100	01000	01001	00000	00000
76E	00001	00100	00000	00000	00000	00100	00000	00001	00000	00000
77B	00001	00100	00000	00000	00000	00100	00000	00001	00000	00000
78A	00001	00100	00000	00000	00000	00100	00000	00001	00000	00000
79	00001	00100	00000	00000	00000	00100	00000	00001	00000	00000
80A	00000	01000	00000	00000	00011	00010	00000	00001	01000	00001
80C	00001	01000	00000	10000	01010	00010	00010	10001	00000	00001
80D	00001	01000	00000	00000	00010	00010	00000	00001	01000	00001
81A	00001	00000	00000	00000	00000	00100	00000	00001	00000	00000
82A	00001	00100	00000	00000	00010	00100	00000	00001	00000	00000
83A	00001	00100	00000	00000	00010	00100	00000	00001	00000	00000
84G	00001	00100	00000	00000	00010	00100	00000	00001	00000	00000
85A	00000	01010	00000	00000	00001	00000	00000	00001	00000	00000
85B	00000	01010	00000	00000	00001	00010	00001	10001	00000	00000
85D	00001	01010	01000	10000	00011	00010	00001	00001	10000	00000
85E	00000	01010	00000	00000	00011	00010	00000	00001	00000	00001
86A	00000	01010	00000	00000	00010	00010	00000	00001	00100	00001
86B	00000	01010	00000	00000	00001	00010	00000	00001	00000	00001

Fragments 201-250

87A	00000	01010	00000	00000	00010	00010	00000	10001	00000	00001
89A	00000	01010	00000	10000	00000	00010	00010	00000	00100	00001
89B	00000	01010	00000	00000	00010	00010	00000	00001	00100	00001
90A	00000	01000	00000	00000	00011	00010	00001	00001	00101	00000
90B	00000	01000	00000	00000	00011	00010	00001	00001	00000	00001
90C	00000	01000	00000	00000	00010	00010	00000	10001	00101	00001
92A	00100	01010	00000	00000	00000	00000	00000	10000	10101	00001
92B	00000	01000	00000	00010	00000	01010	00000	00001	00100	00000
93C	00000	01010	00000	00000	00000	00010	00000	00001	00000	00000
95A	00000	01000	10000	00000	00011	00010	00000	10001	00100	00000
96A	00000	01010	00010	00000	00010	00000	00000	00011	00000	00000
96B	00001	01011	00000	10000	00000	00011	00010	10010	00010	00001
98A	00000	10110	00010	00000	00000	00010	00010	10000	00000	00000
98B	00001	10111	10000	01000	00000	00010	00010	10000	00010	00000
98C	00000	10110	00010	00000	00000	00010	00010	10000	00000	00000
99A	11000	10010	10000	00000	00000	00010	10000	10000	00010	00000
99B	00000	10010	10000	00000	00000	00010	00000	10000	00110	00000
99C	00000	10110	00000	00000	00000	00010	00000	10000	00010	00000
100B	00000	01010	00000	00000	00001	00010	10100	10000	00000	01000
101A	00001	10110	00000	00000	00100	00010	00000	10000	00010	00001
102A	00000	10100	00000	00001	00101	00010	10011	10011	00010	00100
102B	00000	10100	00000	00000	00000	00010	00000	10000	00010	00000
102C	00000	10110	00000	01000	00000	00010	00000	10000	00010	00000
103A	00000	10110	00000	00000	00000	00010	00010	10001	00010	00000
103B	00000	10010	00000	00000	00000	00010	00000	10000	00010	00000
103C	10000	10110	00000	00000	00000	00010	00010	10000	00010	00011
104	00000	10110	00000	00000	00000	00010	00000	10000	00010	00000
105	00000	10110	00000	00000	00000	00010	10000	10000	00010	00011
106A	00000	10110	10000	01001	00000	00010	00000	10000	00010	00000
106B	00000	10110	00000	01000	00000	00010	00000	10000	00010	00000
106C	00000	10110	00000	00000	00000	00010	00000	10000	00010	00000
108A	00000	10110	00000	00000	00000	00010	00000	10000	00010	00010
108B	00000	10110	00000	00000	00000	00010	00000	10000	00010	00000
108C	00000	10100	00000	00000	00000	00010	00010	10000	00010	00001
110	00000	10110	00000	01000	00000	00010	00000	10000	00010	00000
111A	00000	10010	00000	00000	00000	00010	00000	10000	00010	00001
111B	00000	10010	10000	00000	00000	00010	00000	10000	00010	00000
111C	00011	10110	00000	01000	00000	00010	00000	10000	00010	00000
112	00000	10010	10100	00000	00000	10010	01000	00001	00000	00000
113A	00000	10010	10100	00000	00000	10010	01000	00001	00000	00000
114	00000	10010	10000	01000	00000	00010	00000	10000	00010	00000
116	00000	01010	00000	00000	00000	00010	00000	10000	00000	00000
118A	00000	01010	00000	00000	00000	00010	00000	00001	00000	00000
120C	00000	00000	00000	00000	00001	00000	00000	00000	00000	00000
121A	00000	10010	00000	00000	00000	00000	00000	00001	00000	00000
122A	00000	10010	00000	00000	00000	00010	00000	10001	00000	00000
123F	00000	10010	00001	00000	00000	00010	01000	10000	00000	00000

Fragments 201-250

123G	00010	10010	00000	00000	00000	00010	01000	00000	00000	00000
125A	00010	10010	00000	00000	00000	10010	00000	00001	00000	00000
126A	00000	01010	00000	00000	00000	00010	00000	00001	00000	00000
126E	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
127A	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
128A	00000	10010	00000	00000	00001	00000	00000	00000	00000	00000
130A	00000	10010	00000	00000	00000	00000	00000	00001	00000	00000
131	00000	01010	01000	00000	00000	00010	00000	00001	00100	00000
134A	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
HEN1	00000	00000	00000	00000	00000	00000	00000	00001	00000	00000
S3	00000	01000	00000	00000	00010	00000	00000	00001	00000	00000
S7	00000	01010	00000	00000	00011	00010	00000	00001	00000	00001

Fragments 251-300

1	00000	00000	00000	01010	00000	00000	00000	00100	00101	00000
3	00000	00000	00000	00010	00000	00010	10000	00010	00001	00010
5	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
8	01000	00000	00000	00000	00000	00000	00000	00010	00100	00000
10	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
12	01000	00000	00000	00010	00000	00000	00000	00010	00101	00000
13	01000	00000	00000	00010	00000	00000	00000	00010	00110	00000
14	00000	00000	00000	00010	00000	00000	10000	00010	00110	00000
16	01001	00000	00000	01010	00010	00000	00000	00100	00100	00000
17	00000	00000	00000	00010	00000	00000	10000	00010	00110	00001
18	00000	00000	00010	00010	00000	00010	10000	00011	00111	00001
19	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
20	00000	00000	00000	00010	00000	00000	00000	00010	00111	00000
25	01100	00000	00000	00000	00000	00000	00000	00010	00100	00000
26	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
27A	00000	00000	00000	00010	00000	00010	00000	00011	00101	00000
27B	01000	00000	00000	00010	00001	00001	00000	00011	10010	00010
27C	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
28A	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
29A	01000	00000	00000	00010	00000	00000	00000	00010	00110	00000
30A	00100	00000	00000	00010	00000	00000	00000	00010	00100	00000
30D	00001	00000	00000	01000	00000	00001	01000	00011	00110	00000
30E	01110	00100	00000	00010	00000	00000	10000	00010	00101	00001
30F	01000	00000	00000	00000	00000	00000	00000	00010	00100	00001
31	01000	01000	00000	01010	00000	00000	10000	00011	00110	01001
32B	01000	00000	00000	00010	00001	00000	00000	00011	00110	00000
36B	00000	00000	10000	00001	00000	00000	00000	00010	00101	00000
37A	01000	00000	00000	00000	00000	00000	00000	00010	00110	00000
37B	01000	00000	00000	00010	00100	00000	00000	00011	00110	00000
37C	00000	00000	00000	00010	10000	00000	01000	00011	10110	00001
39	01000	00000	00000	00000	00000	00000	00000	00100	00110	00000
41A	01000	00000	00000	00010	00000	00000	00000	00011	00110	00001
41B	01000	00000	00000	00000	00000	00000	00000	00010	00101	00000
41C	01000	00000	00000	00000	00000	00000	00000	00000	00001	00000
43	10000	00000	00000	00000	00010	00000	00000	00100	10100	00000
45A	01000	01000	00000	01010	00000	00000	10000	00011	01100	00000
45B	01000	00000	00000	00000	00000	00000	10000	00011	00110	00000
45C	01000	00000	00000	00010	00000	00010	10000	00011	00111	00001
46A	01000	00000	00000	00000	00000	00000	00000	00010	00110	00000
46B	01000	00000	00000	00000	00000	00000	00000	00010	00110	00000
46C	01000	00000	00000	00000	00000	00000	00000	00010	00100	00000
48A	00000	00000	00000	00000	00000	00000	00000	00010	00100	00000
48B	10000	00000	00000	01010	00000	00010	11000	00010	00111	00000
48C	01000	00000	00000	00000	00000	00000	00000	00100	00100	00000
48D	00000	00000	00000	00000	00000	00000	00000	00010	00100	00000
50A	10000	00000	00000	00000	00000	00000	00000	00110	00100	00000
50B	10000	00000	00000	00000	00000	00000	00000	00100	00101	00000

Fragments 251-300

50C	10000	00000	00000	00000	00000	00000	00000	00100	00101	00000
50D	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
52B	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
54A	00000	10000	10000	00001	00000	00001	00000	00100	00100	00000
55A	00000	10010	10000	00001	00000	00001	00000	00100	00100	00000
57B	00000	00000	00000	00010	00000	00000	00000	00010	00111	10000
58B	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
58D	00000	00000	00000	00010	00000	00000	00000	00010	00101	01000
61	00000	00000	00000	00010	00000	00000	00000	00010	00111	00100
62D	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
63B	00000	00000	00000	00000	00000	00000	00000	00010	00101	00000
65C	00000	00000	00000	00010	00000	00000	10000	00010	00111	00001
65D	00000	00000	00000	00010	00000	00000	00000	00010	00100	00001
66C	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
66D	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
67A	00000	00000	00000	00010	00000	00000	00000	00010	00110	00100
67B	00000	00000	00000	00010	00000	00100	00000	00010	00100	00100
68A	00000	00000	00000	00010	00000	00000	10000	00010	00100	00010
68B	00000	00000	00001	00010	00000	00000	00000	00010	00111	00000
68C	00000	00000	00001	00110	00000	00000	00000	00110	00110	00000
69A	00000	00000	00000	00010	00000	00000	00000	00010	00101	00100
69B	00000	00000	00000	00010	00000	00000	10000	00010	00110	00100
70A	01000	00000	00000	00000	00000	00000	00000	00110	00100	01000
70B	01000	00000	00000	00010	00000	00000	00000	00110	00111	00000
71A	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
72A	01000	00000	00000	00000	00000	00000	00000	00010	00110	00000
72B	01000	00000	00000	00000	00000	00000	00000	00010	00111	00000
74D	00000	10010	10000	00001	00000	00000	00000	00100	10100	00000
75A	00000	00000	00000	00010	00000	00000	00000	00011	10110	00001
76A	00000	00000	00000	00010	00000	00100	00000	00110	00111	00100
76E	00000	00000	00000	00010	00000	00100	00000	00010	00100	00000
77B	00000	00000	00000	00010	00000	00000	10000	00010	00110	00000
78A	00000	00000	00000	00000	00000	00100	00000	00010	00100	00000
79	00000	00000	00000	00010	00000	00000	10000	00010	00110	00000
80A	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
80C	01000	00000	00000	00010	00000	00000	10000	00111	00110	01000
80D	01000	00000	00000	00000	00000	00000	10000	00011	10110	01010
81A	00000	00000	00000	00010	00000	00100	00000	00010	00100	00000
82A	00000	00000	00000	00010	00000	00000	00000	00010	00110	00000
83A	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
84G	00000	00000	00000	00010	00000	00000	00000	00010	00110	00001
85A	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
85B	01000	00000	00000	00010	00000	00000	00000	00110	00110	00000
85D	01000	00000	00000	00000	00000	00000	00000	00010	00110	00000
85E	01000	00000	00000	00010	00000	00000	00000	00110	00110	00000
86A	01100	00000	00000	00010	00000	00000	10000	00111	00110	00000
86B	00000	00000	00000	00010	00000	00000	00000	00011	00110	00001

Fragments 251-300

87A	00000	00000	00000	00010	00000	00000	10000	00010	00110	00001
89A	01000	00000	00000	00100	00000	00010	00000	00010	00100	00001
89B	01000	00000	00000	00000	00000	00000	00010	00010	00100	00000
90A	00010	00000	00000	00000	00000	00000	00000	00010	00110	01000
90B	00010	00000	00000	00010	00000	10000	10000	00110	00110	00000
90C	00000	00000	00000	00010	00000	00000	10000	00010	00010	00001
92A	00000	00000	00000	01110	00000	00000	00000	00011	00100	00000
92B	01100	00000	00000	00000	00010	00010	00000	00010	01010	00011
93C	00000	00000	00000	00010	00000	00000	00000	00010	00101	00000
95A	01000	00000	00000	00010	00000	00000	00000	00110	00100	00000
96A	00000	00100	00000	00000	00010	00000	01000	00011	01100	00001
96B	01000	00000	00000	00000	00000	00010	00000	00011	00110	00001
98A	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
98B	01000	00000	00000	00000	00000	00000	00000	00111	00110	00000
98C	00000	00001	00000	00000	00000	00000	10000	00011	10110	00000
99A	01000	00000	00000	00000	00000	00000	00000	00110	00110	00000
99B	01000	00000	00100	00010	00000	00000	00000	00011	00110	00000
99C	01000	00000	00000	00010	00000	01000	00000	00111	01111	00000
100B	00000	10000	10000	00000	00000	00010	01000	00110	00110	00000
101A	00000	00000	00000	00010	00000	00000	00000	00010	00110	00000
102A	01000	00000	00000	00000	00000	00000	00000	00110	00110	00000
102B	00000	00000	00000	00010	00000	00000	10000	00010	00110	00001
102C	01000	00000	00000	00000	00000	00000	00000	00110	00110	00000
103A	01000	00000	00000	00010	00000	00000	10000	00010	00100	00000
103B	11000	00000	00000	00100	00000	00000	00000	00010	00110	00000
103C	00100	00001	00000	00000	00000	00000	00000	00000	00010	00000
104	01000	00000	00000	00010	00000	00000	10000	00010	00110	00000
105	00000	00000	00000	00000	00000	00000	00000	00110	00110	00000
106A	10000	00000	00000	00000	00000	00000	00000	00010	00100	00001
106B	10000	00001	00000	00000	00000	00000	00000	00100	00110	00000
106C	01000	00000	00000	00000	00000	00000	00000	00100	00110	00000
108A	01000	00000	00000	00000	00000	00000	00000	00100	00110	00000
108B	00000	00000	00000	00000	00000	00000	00000	00110	00110	00000
108C	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
110	10000	00000	00000	00100	00000	10000	00000	00110	00110	00000
111A	01000	00000	00000	00010	00000	00000	00000	00011	00110	00001
111B	01000	00000	00000	00000	00000	00000	00000	00110	00111	00000
111C	01000	00000	00000	00000	00000	00000	00100	10100	00100	00000
112	00000	00000	00000	00010	00000	00000	10000	00010	00101	00001
113A	00000	00000	00010	00010	00000	00000	00000	00011	00110	00001
114	01000	00000	00000	00000	00000	00000	10000	00010	00110	00000
116	00000	00000	00000	00010	00000	00000	00000	00011	00111	00001
118A	00000	00000	00000	00010	00000	00000	10000	00010	00101	00000
120C	00000	00000	00000	00110	00000	00000	00000	00011	00110	00001
121A	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
122A	00000	00000	00000	00010	00000	00010	10000	00010	00111	00001
123F	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000

Fragments 251-300

123G	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
125A	00000	00000	00000	00010	00000	00000	10000	00011	00100	00001
126A	00000	00000	00000	00010	00000	00010	10000	01011	10111	10001
126E	00000	00000	00000	00010	00000	00000	00000	00010	00101	10000
127A	00000	00000	00000	01010	00000	00000	10000	00011	00100	00001
128A	00000	00000	00000	00010	00000	00000	10000	00011	00110	00001
130A	00100	00000	00000	01010	00000	00000	10000	00011	00110	00001
131	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
134A	00000	00000	00000	00010	00000	00000	10000	00010	00111	00001
HEN1	00000	00000	00000	00010	00000	00010	10000	00011	00110	00001
S3	00000	00000	00000	00010	00000	00000	00000	00010	00100	00000
S7	00000	00000	00000	00010	00000	00000	00000	00011	00110	00001

Fragments 301-350

1	00100	00010	00000	00000	10000	00010	00001	00000	00000	00000
3	00100	00010	00000	00000	11000	00010	00000	00001	10000	00000
5	00000	00100	00000	00000	10000	00011	01000	00000	10000	00010
8	00100	00100	00100	00000	10000	00011	00000	00000	10010	00000
10	00000	00100	00000	00000	10000	00011	00000	00000	10000	00010
12	00000	00100	00000	00000	10000	00010	00000	00000	10010	00000
13	00000	00100	00000	01000	10000	01011	00001	11010	10000	00000
14	00100	00100	00000	00000	10000	00010	00001	00000	10010	00000
16	00001	00100	00000	00000	10000	00010	00001	10000	10000	00000
17	01100	00100	00001	00100	00000	01101	00001	00001	00000	00000
18	01100	00010	00001	00100	00000	01100	00001	10001	00000	00001
19	00000	00110	00000	00000	10000	00000	00000	00000	00000	00000
20	00000	00010	00000	00000	10000	00010	00000	00000	01000	00000
25	00000	00100	00000	00000	10000	00010	00000	00000	01000	00000
26	00000	00000	00000	00000	10000	00010	00000	01000	01000	00001
27A	00001	00100	00101	01010	10000	00011	00000	10000	00000	00000
27B	00100	00100	00000	01000	10010	00010	00000	00000	10000	00000
27C	00000	00000	00000	00000	10000	00010	00000	00000	10000	00000
28A	00100	00100	00000	00000	10000	00010	00000	00000	10000	00000
29A	00000	00100	00000	00000	10000	00011	01000	00000	10000	00000
30A	00000	10100	00000	00000	10000	00010	00000	00000	10011	00100
30D	00100	00100	01000	10100	10100	00000	00001	00100	10000	00000
30E	00000	00100	00000	00000	10000	00010	00000	00000	00000	00000
30F	00100	00100	00000	00000	10001	01000	01001	11000	10000	00000
31	01100	00100	00000	01100	10000	00000	00000	00000	00000	00000
32B	00000	00100	00001	01000	10000	00000	00000	00000	10000	00000
36B	00000	00000	00000	00000	10000	00000	00010	00000	10000	00000
37A	00000	00000	00000	00000	10000	00011	10000	00000	10011	10000
37B	00100	00100	00000	01010	10001	00011	01000	00000	10001	00000
37C	00110	00100	00000	01000	10000	00000	00000	10011	00000	00000
39	00000	00000	00000	00000	10000	00011	10000	00000	10000	10000
41A	01100	00101	00001	00110	10000	00100	00000	00010	00000	00000
41B	00000	00000	00000	00000	10000	00011	00000	00000	01000	10000
41C	00000	00000	00000	00000	10000	00011	00000	00000	10000	10100
43	00000	00100	10000	00000	10000	00011	00001	00001	10000	00000
45A	01100	00100	01000	10010	00000	00100	00000	00001	00000	00000
45B	10100	01100	01000	00011	10000	00100	00000	00000	00000	00000
45C	00100	00100	00001	00110	10001	00100	00000	00001	00000	00000
46A	00000	00000	00000	00000	10000	00011	00000	00000	10000	10000
46B	00000	00100	01000	00010	10100	00011	00000	00000	10000	00000
46C	00000	00000	00000	00000	10000	00011	10000	00000	10000	10000
48A	00000	00000	10000	00000	10000	00011	00000	00000	10000	00000
48B	10100	00100	00000	01000	10000	00010	00001	00001	00000	00000
48C	00000	00000	00000	00000	10000	00011	10000	00000	10000	00000
48D	00000	00000	10000	00001	10000	00011	00000	00000	10000	00000
50A	00000	00000	00000	00000	10000	00011	00000	00000	10110	00000
50B	00000	00000	00000	00000	10100	00010	00000	00000	10000	00000

Fragments 301-350

50C	00000	00000	00000	00000	10100	00011	10000	00000	10000	00000
50D	00000	00100	00000	00000	10000	00011	00000	00000	01000	00000
52B	00000	00100	00000	00000	10000	00011	00000	00000	01000	00000
54A	00000	00000	00000	00000	10000	00000	00010	00000	10000	00000
55A	00000	00000	00000	00000	10000	00000	00010	00000	10000	00000
57B	00000	00100	00000	00000	10000	00000	00000	00000	00000	00000
58B	00000	00100	00000	00000	10000	00010	00000	00000	01000	00001
58D	00100	00000	00000	00100	10000	00011	00001	10100	10000	00000
61	00000	00100	00000	00000	10000	00000	00000	00000	00000	00000
62D	00100	00010	00000	00000	10000	00011	01000	00000	01000	00000
63B	00000	00100	00000	00000	10000	00010	00000	00000	00000	00000
65C	01100	00101	00001	00100	10000	00100	00001	00001	00000	00000
65D	00100	10100	00000	00000	10000	00011	00000	00000	10000	00000
66C	00000	00010	00000	00000	10000	00010	01000	00000	00000	00000
66D	00100	00010	00000	00000	10000	00010	00000	00000	10000	00001
67A	00000	00010	00000	00000	10000	00000	00001	11000	01000	00001
67B	00000	00010	00000	00000	10000	00000	00001	01000	00000	00001
68A	00100	00100	00000	00000	10000	00010	00000	00000	10000	00000
68B	00000	00100	00000	00000	10000	00011	00001	00000	10000	00000
68C	00000	00000	00000	00100	10000	00011	01000	00000	10001	00000
69A	00000	00010	00000	00000	10000	00010	00001	11000	00000	00100
69B	00000	00010	00000	00000	10000	00010	00001	00000	00000	00001
70A	00000	00100	00100	00000	10000	00011	00001	00000	10000	00000
70B	00100	00100	00000	00000	10000	00011	11001	00000	10000	00000
71A	00000	00010	00000	00000	10000	00010	00000	00000	00000	00001
72A	00000	00100	00000	00000	10000	00011	00000	00001	10000	00000
72B	00000	00100	00000	00100	10000	00011	00100	00000	10000	00000
74D	00000	00100	00000	00000	10000	00000	00010	00000	10000	00000
75A	01100	00100	00001	00100	00000	00000	00000	00001	00000	00000
76A	00000	00010	00000	00000	00000	00000	00000	01000	00000	00001
76E	00000	00010	00000	00000	10000	00000	00000	00000	00000	00001
77B	00100	00010	00000	00000	10000	00010	00000	00000	00000	00001
78A	00000	00000	00000	00000	10000	00000	00000	00000	00000	00000
79	00100	00010	00000	00000	10000	00000	00000	00000	00000	00001
80A	00100	00101	00000	00000	10000	00110	00001	00000	00000	00000
80C	00000	00100	00000	01000	10001	00011	01001	00001	00000	00000
80D	00100	00000	00001	00000	11000	00010	00100	00000	00000	00000
81A	00000	00000	00000	00000	10000	00000	00000	00000	00000	00001
82A	00000	00010	00000	00000	10000	00011	00000	00000	10000	00001
83A	00000	00010	00000	00000	10000	00010	00000	00000	10000	00000
84G	00100	00100	00000	00100	10000	00000	00001	00000	00000	00001
85A	00000	00000	00000	00000	00000	00000	00001	00000	10000	00000
85B	00000	00100	00000	00000	10000	00011	00101	00000	10000	00000
85D	00000	00100	00000	00000	10000	00011	00000	10010	10000	00000
85E	00100	00100	10000	00000	10001	00111	11000	00000	10000	00000
86A	00000	00000	00000	00000	10000	01010	00000	00000	10000	00000
86B	00100	00100	01010	00000	10000	00110	01000	00000	00000	00000

Fragments 301-350

87A	01000	00100	00000	00100	10000	00000	01001	10001	10000	00000
89A	01100	00000	00000	00000	10000	00000	00000	00001	00000	00010
89B	00000	00000	00000	00000	10000	00011	00000	00000	10000	10000
90A	00000	00100	00000	00000	10000	01011	01000	00001	10000	00000
90B	00000	00000	00000	00010	10000	00011	00000	00000	10000	00000
90C	01100	00100	00000	00110	10000	00000	00001	00001	00000	00000
92A	00100	00100	00000	00110	10000	01000	00000	00010	00000	00000
92B	01100	00100	00000	00100	10000	00100	00001	00010	00100	00000
93C	00000	00100	00000	00000	10000	00000	00000	00000	00000	00000
95A	01000	00100	00000	00100	10000	01111	01001	01000	10000	00010
96A	00100	00100	00010	10000	00000	00000	00000	00000	00000	00000
96B	00100	00100	00000	00100	10000	00000	00000	00000	00000	00000
98A	00000	00000	00000	00000	00000	00000	00000	00000	00000	00000
98B	00000	00100	01000	00000	10000	00011	00000	00000	11000	00000
98C	01100	00001	00000	00010	00000	00000	00000	00000	00000	00000
99A	00000	00100	00000	00000	10000	00011	00000	00000	10000	00000
99B	00000	00100	00000	00000	10000	00011	00000	00000	10000	00000
99C	00000	00100	00000	00010	10000	10011	00000	00001	10000	00000
100B	00000	00100	00100	00000	10000	00000	00011	00000	10000	00000
101A	00000	00100	00000	00000	10000	00011	00000	00000	10000	00000
102A	00000	00001	00100	00000	10000	00011	01000	00000	10011	00100
102B	00100	00100	00010	00100	10000	00111	00000	00001	10000	00000
102C	00000	00100	00000	00000	10100	00011	00000	00000	10000	00000
103A	00000	00100	00000	00011	10000	00111	00000	00000	10000	00000
103B	00000	00000	10000	00000	10001	00010	10000	00000	10000	00000
103C	00000	00000	00000	00000	10000	00011	00000	00000	10001	10000
104	00000	00100	00000	00011	10000	00010	00000	00000	00000	01000
105	00000	00000	00000	00000	10000	00011	00000	00000	10000	00000
106A	00000	01100	01000	00000	10000	00100	00000	00000	00001	00000
106B	00000	10000	00000	00000	10000	00011	00000	00000	10000	00000
106C	00000	10000	00000	00000	10000	00011	10010	00000	10000	00000
108A	00000	00000	00000	00000	10100	00011	00000	00000	11000	00100
108B	00001	00100	00000	00000	10000	00011	10000	00000	10000	00000
108C	00000	00100	00000	00000	10000	00011	00000	00000	10000	00000
110	00000	00000	00000	00000	10000	00010	10000	00000	10000	00000
111A	01100	00100	00001	00111	10000	00100	00001	10001	00000	00000
111B	00000	00000	00000	00000	10000	00111	00000	00000	10000	00000
111C	00000	00000	00000	00010	10000	00011	00000	00000	10000	00000
112	01100	00100	00000	00100	00000	00100	00000	00001	00000	00000
113A	01100	00100	00001	00110	00000	00100	00001	10001	00000	00000
114	00000	00000	00000	00000	10000	00011	00000	00000	10000	00000
116	01100	00100	00000	00110	00000	00100	00001	10001	00000	00000
118A	00100	00100	00000	00000	10000	00110	00001	00000	00000	00000
120C	01100	00101	00000	00100	00000	00100	00001	10001	00000	00000
121A	00000	00100	00000	00000	10000	00010	00000	00000	10000	01000
122A	00100	00100	00000	00100	10000	00100	00000	00000	00000	00000
123F	00100	10100	01000	00000	10000	00010	00001	10000	10000	00000

Fragments 301-350

123G	00000	00100	00000	00000	10000	00010	00101	00000	10000	00000
125A	01100	00100	00001	00110	10000	00100	00001	10001	00000	00000
126A	01100	00100	00010	00100	10000	00111	00001	00001	00000	01000
126E	00000	00100	00000	00000	10000	00010	00000	00000	00000	00000
127A	00100	00100	00010	00110	00000	01100	00001	00001	00000	00000
128A	01100	00100	00000	00110	00000	01100	00001	00001	00000	00000
130A	01000	00100	00001	00100	10000	00100	00001	10001	00000	00000
131	00100	00100	00000	00000	10000	00011	00000	00000	10000	00000
134A	01100	00101	00001	00100	10000	00100	00001	10001	00000	00000
HEN1	01000	00101	00000	00000	00001	00100	00001	01001	00000	00000
S3	00000	00100	00000	00000	10000	00010	00000	00000	10000	00000
S7	01100	00100	00001	00100	10000	00000	00001	00000	00000	00000

Fragments 351-376

1	10000	00000	00000	00000	11000	1
3	10000	00000	00000	00000	11000	0
5	00000	00000	00000	00000	01010	0
8	00000	00000	01000	00000	01010	1
10	00010	00000	00000	00000	01000	1
12	10000	00000	00000	00000	01010	1
13	00100	00000	00000	00000	01000	1
14	00010	00000	00000	00000	01000	1
16	10000	00000	00100	00010	01010	0
17	00000	00000	00000	00000	01000	0
18	00000	00000	00000	00000	00000	0
19	00000	00000	00000	00000	01000	0
20	00000	00000	00000	00000	01000	0
25	00000	00000	00000	00010	01000	0
26	00000	00000	00000	00000	11000	0
27A	11000	00010	00000	00000	01010	1
27B	11000	00000	00000	00000	01000	1
27C	10000	00000	00000	00000	01000	1
28A	10000	00000	00000	00000	01000	0
29A	00000	00000	00010	00000	01000	0
30A	00010	00000	00000	00000	01000	1
30D	00000	00000	00000	00000	01000	1
30E	00000	00000	00000	00000	01000	1
30F	00000	00000	00100	00000	01000	1
31	00000	00000	00000	00000	01000	1
32B	00100	00000	00000	00000	01000	1
36B	00000	10000	00000	00000	00000	0
37A	00000	00000	00000	00000	01100	0
37B	00010	00000	00000	00000	01000	0
37C	00000	00000	00000	00000	01000	0
39	00000	00000	00100	00000	01000	0
41A	00000	00000	00000	00000	01000	0
41B	00000	00000	00000	00010	01000	0
41C	00000	00000	00000	00010	01000	0
43	00000	00000	00000	00010	01000	0
45A	00000	00000	00000	00000	01000	0
45B	00000	00000	00000	00000	01000	0
45C	00000	00000	00000	00000	01000	0
46A	00000	00000	00000	00000	01000	0
46B	00000	00000	00000	00010	01100	0
46C	01000	00000	00000	00000	01000	0
48A	00000	00000	00000	00000	01000	0
48B	00000	00000	00000	00010	01000	0
48C	00000	00000	00000	00000	01000	0
48D	01000	00000	00000	00000	01000	0
50A	00000	00000	00000	00010	01100	0
50B	00000	00000	00000	00010	01100	0

Fragments 351-376

50C	00000	00000	00000	00000	01010	0
50D	00000	00100	00000	00010	01000	0
52B	00000	00000	00000	00000	01000	0
54A	00000	10000	00001	00000	00000	0
55A	00000	10000	00001	00000	00000	0
57B	00000	00000	00000	00000	01000	0
58B	10000	00000	00000	00000	01100	0
58D	10000	00000	00000	00000	01000	0
61	00000	00000	00000	00000	01000	0
62D	00000	00000	00000	00000	01000	0
63B	00000	00000	00000	00000	01000	0
65C	00000	00000	00000	00000	01000	0
65D	00000	00000	00000	00000	01000	0
66C	00000	00000	00000	00000	01000	0
66D	00000	00000	00000	00000	01000	0
67A	00001	00000	00000	00000	11000	0
67B	00000	00000	00000	10001	11000	0
68A	01000	00000	00000	00000	01000	1
68B	11000	00000	00000	00000	11000	1
68C	00100	00000	00000	00000	01000	1
69A	00000	00000	00000	10001	01010	0
69B	00001	00000	00000	00000	01000	0
70A	10000	00000	00000	00000	01000	1
70B	00000	00000	00000	00000	01000	1
71A	00000	00000	00000	00000	11001	0
72A	00000	00000	00000	00000	01000	1
72B	00000	00000	00000	00000	01000	1
74D	00000	10000	00001	00000	01000	0
75A	00000	00000	00000	00000	01000	0
76A	00000	00000	00000	10000	01000	0
76E	00000	00000	00000	00000	01000	0
77B	00000	00000	00000	00000	01001	0
78A	00000	00000	00000	00000	01000	0
79	00000	00000	00000	00000	11000	0
80A	10000	00000	00000	00000	01000	1
80C	10000	10010	00000	00000	11000	1
80D	00000	00000	00100	00000	11000	0
81A	00000	00000	00000	00000	01000	0
82A	00000	00000	00000	00000	01000	0
83A	00000	00000	00000	00000	01000	0
84G	00000	00000	00000	00000	01001	0
85A	00000	00000	00000	00000	01000	0
85B	00000	00000	00000	00000	01010	0
85D	00000	01000	00000	00000	01000	0
85E	00000	00000	00000	00000	01010	0
86A	10100	00000	00000	00000	01010	1
86B	00000	00000	00000	00000	01010	1

Fragments 351-376

87A	10000	00000	00000	00000	01000	1
89A	10100	00000	00000	00000	01010	1
89B	00100	00000	10000	00000	01000	1
90A	00101	00000	00000	00000	01000	1
90B	00100	00000	00000	00000	01000	1
90C	00000	00000	00000	00000	01000	0
92A	00000	00100	00000	00000	01000	0
92B	00000	00000	00000	00000	01000	0
93C	00000	00000	00000	00000	01000	0
95A	00000	00000	00100	00000	01000	1
96A	00000	00000	00000	00000	11000	0
96B	00000	00000	00000	00000	01000	0
98A	00000	00000	00000	00000	01000	0
98B	01000	00000	00000	00000	01000	0
98C	00000	00100	00000	00000	01000	0
99A	00000	00000	00000	00000	01000	0
99B	00001	00000	00000	00000	01000	0
99C	00000	00000	00000	00000	01000	0
100B	00000	00010	00000	00000	01000	0
101A	00000	00000	00100	00000	01000	0
102A	00000	00000	01000	00010	01000	0
102B	00000	00100	00000	00000	01000	0
102C	00000	00000	00000	00000	01000	0
103A	00000	00000	00000	00010	01000	0
103B	00000	00000	00000	00010	01000	0
103C	00000	00000	00000	00010	01000	0
104	00000	00000	00000	00010	01000	0
105	00000	00000	00000	00000	01100	0
106A	00000	00000	00000	00000	01000	0
106B	00000	00010	00000	00010	01000	0
106C	00000	00010	00101	00010	01000	0
108A	00010	00000	00000	00010	01000	0
108B	00000	00000	00000	00000	01000	0
108C	00001	00000	00000	00000	01000	0
110	00000	00000	00000	00000	01000	0
111A	00000	00000	00000	00000	01000	0
111B	00000	00000	00000	00000	01000	0
111C	00000	00000	00100	00000	01000	0
112	00000	00000	00000	00000	01000	0
113A	00000	00000	00000	00000	01000	0
114	00000	00000	00000	00000	01000	0
116	00000	00000	00000	00000	01000	0
118A	00000	00000	00000	00000	01000	0
120C	00000	00000	00000	00000	01000	0
121A	00000	00000	00000	00000	01000	0
122A	00000	00000	00000	00000	01000	0
123F	00000	00000	00000	00000	01000	0

Fragments 351-376

123G	00000	00000	00000	00000	01000	0
125A	00000	00000	00000	00000	01000	0
126A	00000	00000	00000	00000	01000	0
126E	00000	00000	00000	00000	01000	0
127A	00000	00000	00000	00000	01000	0
128A	00000	00000	00000	00000	01000	0
130A	00000	00000	00000	00000	01000	0
131	10000	00000	00000	00000	01000	0
134A	00000	00000	00000	00000	01000	0
HEN1	00000	00000	00000	00000	01000	0
S3	10000	00000	00000	00000	01000	1
S7	10000	00000	00000	00000	01010	0

Appendix C. AFLP fragment sizes

This appendix gives the size in base pairs (bp) for the AFLP fragments listed in Appendix B. AFLP fragments 1 -135 were generated with the selective primer combination having extensions -CCC/-AGC, 136-269 were generated with the selective primer combination having extensions -CCC/-ATA, and 270-376 were generated with the selective primer combination having the extensions -CGG/-AGC. The extensions refer to the EcoRI and MseI adapters, respectively.

AFLP Fragment	Size (bp)	AFLP Fragment	Size (bp)	AFLP Fragment	Size (bp)
1	38	46	112	91	217
2	42	47	113	92	221
3	47	48	115	93	223
4	49	49	117	94	224
5	52	50	120	95	231
6	53	51	122	96	239
7	54	52	123	97	242
8	55	53	125	98	245
9	56	54	126	99	247
10	61	55	128	100	261
11	64	56	129	101	265
12	65	57	130	102	269
13	67	58	131	103	271
14	68	59	132	104	285
15	69	60	133	105	292
16	70	61	135	106	297
17	71	62	136	107	299
18	72	63	138	108	302
19	73	64	139	109	305
20	74	65	141	110	307
21	76	66	143	111	309
22	77	67	145	112	313
23	78	68	148	113	321
24	81	69	152	114	332
25	84	70	155	115	363
26	86	71	156	116	367
27	87	72	161	117	370
28	88	73	165	118	372
29	89	74	166	119	373
30	91	75	167	120	374
31	92	76	170	121	375
32	93	77	173	122	389
33	94	78	174	123	396
34	95	79	176	124	403
35	96	80	178	125	409
36	97	81	182	126	413
37	98	82	188	127	416
38	100	83	190	128	420
39	101	84	192	129	431
40	103	85	196	130	444
41	105	86	199	131	496
42	107	87	204	132	40
43	108	88	208	133	43
44	109	89	210	134	45
45	111	90	214	135	47

AFLP Fragment	Size (bp)	AFLP Fragment	Size (bp)	AFLP Fragment	Size (bp)
136	50	181	137	226	225
137	52	182	140	227	227
138	55	183	144	228	233
139	56	184	148	229	234
140	60	185	150	230	238
141	63	186	151	231	245
142	64	187	153	232	248
143	65	188	154	233	251
144	67	189	155	234	254
14	69	190	156	235	258
146	71	191	159	236	261
147	73	192	160	237	262
148	74	193	162	238	263
149	75	194	164	239	270
150	77	195	167	240	276
151	80	196	169	241	280
152	82	197	170	242	282
153	84	198	171	243	286
154	88	199	173	244	290
155	90	200	175	245	327
156	94	201	177	246	330
157	95	202	179	247	332
158	96	203	180	248	342
159	99	204	182	249	345
160	100	205	183	250	348
161	101	206	184	251	349
162	103	207	185	252	350
163	104	208	187	253	352
164	105	209	188	254	359
165	106	210	190	255	361
166	108	211	194	256	368
167	110	212	195	257	380
168	113	213	198	258	382
169	114	214	200	259	411
170	116	215	203	260	487
171	117	216	205	261	490
172	120	217	206	262	38
173	121	218	208	263	39
174	122	219	211	264	40
175	125	220	213	265	42
176	127	221	214	266	43
177	128	222	215	267	46
178	130	223	217	268	52
179	132	224	220	269	56
180	135	225	223	270	57

AFLP Fragment	Size (bp)	AFLP Fragment	Size (bp)	AFLP Fragment	Size (bp)
271	58	316	135	361	259
272	59	317	136	362	261
273	60	318	138	363	266
274	64	319	142	364	268
275	66	320	144	365	284
276	67	321	146	366	286
277	68	322	147	367	332
278	69	323	148	368	335
279	71	324	152	369	343
280	72	325	156	370	349
281	73	326	158	371	357
282	74	327	161	372	360
283	75	328	168	373	362
284	76	329	172	374	363
285	77	330	175	375	364
286	78	331	177	376	448
287	79	332	180		
288	81	333	184		
289	83	334	185		
290	86	335	188		
291	88	336	190		
292	91	337	192		
293	93	338	195		
294	95	339	198		
295	99	340	199		
296	101	341	202		
297	103	342	203		
298	104	343	209		
299	105	344	210		
300	107	345	212		
301	108	346	214		
302	109	347	216		
303	112	348	218		
304	113	349	219		
305	114	350	221		
306	116	351	222		
307	117	352	223		
308	118	353	225		
309	119	354	227		
310	120	355	228		
311	122	356	234		
312	124	357	236		
313	126	358	240		
314	131	359	242		
315	132	360	248		

Appendix D. ETS type 1 sequence data

This dataset contains 977 aligned base pairs for 55 external transcribed spacer (ETS) sequences used to reconstruct phylogenetic relationships in Chapter 3. ID codes for the sequences correspond to those found in Table 3.1. Gaps in the sequence are indicated by a dash (-). Uncertainties are denoted by standard IUPAC/IUB codes as follows: N=unknown; R=A/G; Y=C/T; M=A/C; K=G/T; S=C/G; W=A/T.

Bases 1-60

1	TCTC-AATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
10	TSTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
14	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
15	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
12	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
17	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
25.2	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGTT
26.1	TCTCCAATAT	TCTTACGTGT	TCTTGAGTGT	TCATCTACAG	TCCTTGCA-T	A-TTTGAGCT
27C.1	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
27C.10	-CTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
28A	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
3	TCTC-AATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
41B	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
45C	TCTCCAATAT	TCTTACGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
5.11	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
101A	TCTCCAATAT	TCTTACGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
102B.39	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
104	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
108C	-----	----ACGTGT	TCTTGAGTGT	TCATCTACAG	TTCTCACACT	ACTTTGAGCT
111A.4	-----	----CGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
111A.5	-----	----CGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
120C	-----	-----	---TGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
123G	-----	-----	-----	-----	-----	-----
128C	-----	-----	-----	-----	-----T	A-TT-GAGCT
131	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
30A	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
57B.2	-----	----GCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGTT
58B	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
62D.2	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
62D.3	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
65D.7	-CTGCA-TAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAA	TTCTTGCA-T	A-TTTGAGCT
65D.3	-CTGCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAA	TTCTTGCA-T	A-TTTGAGCT
67B.3	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
69A.2	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
69B.6	-----	----CGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
76A.1	TC--CAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
76E.2	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
78A.10	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
78A.5	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
8	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
80A	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
82A.4	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
85A	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
86B	-----	----CGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
90D.11	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
93C.2	TC-C--ATAT	TCTTGCGCGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
93C.3	TCTCCAACAT	TCTTGCGCGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
98A.12	-----	----GCGCAT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
126E.5	-----	-----	-----TGT	TCATCTACAA	TTCTTGCA-T	A-TTTGAGCT
103A	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTGCAG	TTCTTGCA-T	A-TTTGAGCT
50D	T-----	-----	---TGAGTGT	TCATCTACAG	TTCTTGCA-T	ATTTAGAGCT
75A	TCTCCAATAT	TCTTGCGTGT	TCTTGAGTGT	TCATCTACAG	TTCTTGCA-T	A-TTTGAGCT
96A	-----	-----	-----	-----	-----	-----
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 61-120

1	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
10	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
14	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
15	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
12	G-TCTAATCT	ACTT-TGGCA	GGCR-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
17	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
25	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTAACTTAGC	TTGTTCTTGG	AGGTGATTA
26	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
27C.1	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
27C.10	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
28A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
3	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGG	TTGTTCTTGG	AGGTGATCAA
41B	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTAACTTAGC	TTGTTCTTGG	AGGTGATTA
45C	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
5	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
101A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
102B	G-GCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
104	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
108C	ACTCTGATCT	A-TT-TGGCA	GGCACTTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
111A.4	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGATTTAGC	TTGTTCTTGG	AGGTGATCAA
111A.5	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGATTTAGC	TTGTTCTTGG	AGGTGATCAA
120C	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
123G	-----	-----	-----	-----	-----	-----
128C	G-TCTAATTT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
131	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
30A	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
57B	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTAG	CTTACTTAGC	TTGTTCTTGG	AGGTGATCAA
58B	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
62D.2	G-TCTAATCT	ACTT-TGGCA	AGCG-TTTAG	CTAACTTAGC	TTGTTTTTGG	AGGTGATCAA
62D.3	G-TCTAATCT	ACTT-TGGCA	AGCG-TTTAG	CTAACTTAGC	TTGTTCTTGG	AGGTGATCAA
65D.7	G-TCTAATTT	ACTT-TGGCA	GGCG-TTTGG	ATGACTTAGC	TTGTTCTTGG	AGGTGATCAA
65D.3	G-TCTAATTT	ACTT-TGGCA	GGCG-TTTGG	ATGACTTAGC	TTGTTCTTGG	AGGTGATCAA
67B	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
69A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
69B	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
76A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	ATGTTCTTGG	AGGTGATCAA
76E	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
78A.10	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	ATGTTCTTGG	AGGTGATCAA
78A.5	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	ATGTTCTTGG	AGGTGATCAA
8	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
80A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTARC	TTGTTCTTGG	AGGTGATCAA
82A	G-TCTAATCT	ACTT-TGGCA	GGCA-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
85A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
86B	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
90D	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
93C.2	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTAG	CTAACTTAGC	TTGTTCTTGG	AGGTGATCAA
93C.3	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTAG	CTAACTTAGC	TTGTTCTTGG	AGGTGATCAA
98A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTAG	CTAACTTAGC	TTGTTCTTGG	AGGTGATCAA
126E	G-TCTAATTT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
103A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
50D	G-GCTGATCT	ACYTCTGGCA	GGCG-TTTGG	CTGACTTAGC	TTGTTCTTGG	AGGTGATCAA
75A	G-TCTAATCT	ACTT-TGGCA	GGCG-TTTGG	CTGACTTAGS	TTGTTCTTGG	AGGTGATCAA
96A	-----	-----	-----	-----	-----	-----
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 121-180

1	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
10	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAATCATC	CCATGACAGT	TGGTATCAG-
14	GTCTAAGGTC	ACACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
15	GTCTAAGGTC	AYACAGGTAG	AGGACTTCAA	AACAATCATC	CCATGACAGT	TGGTATCAG-
12	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
17	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAS-
25	GTCTAAGGTC	GCACGGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATTAG-
26	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
27C.1	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
27C.10	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAT	AACAGTCATC	CCATGACAGT	TGGTATCAG-
28A	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
3	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
41B	GTCTAAGGTC	GCACGGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
45C	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
5	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
101A	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
102B	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
104	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
108C	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
111A.4	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
111A.5	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
120C	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAATCATC	CCATGACAGT	TGGTATCAG-
123G	-TCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
128C	GTCTAAGATC	GCATGGGTAG	AGGACTTCAA	AATAGTCATC	CCATGACAGT	TGGTATCAGT
131	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
30A	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAATCATC	CCATGACAGT	TGGTATCAG-
57B	ATCTAAGGTT	GCACGGGTAG	AGGACTTCAA	AATAGTTATC	CCATGACAGT	TGGTATCAG-
58B	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
62D.2	GTCTAAGATC	GCATGGGTAG	AGGACTTCAA	AATAGTCATC	CCATGACAGT	TGGTATCAG-
62D.3	GTCTAAGGTC	GCACGGGTAG	AGGACTTCAA	AATAGTTATC	CCATGACAGT	TGATATCAG-
65D.7	GTCTAAGATC	GCATGGGTAG	AGGACTTCAA	AATAGTCATC	CCATGACAGT	TGGTATCAG-
65D.3	GTCTAAGATC	GCATGGGTAG	AGGACTTCAA	AATAGTCATC	CCATGACAGT	TGGTATCAG-
67B	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AATAATCATC	CCATGACAGT	TGGTATCAG-
69A	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
69B	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
76A	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
76E	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
78A.10	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTTATC	CCATGACAGT	TGGTATCAG-
78A.5	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTTATC	CCATGACAGT	TGGTATCAG-
8	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
80A	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAA-
82A	GTCTAAGGTC	ATACAGGTAG	AGGACTTCAA	AACAATCATC	CCATGACAGT	TGGTATCAG-
85A	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
86B	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AATAGTCATC	CCATGACAGT	TGGTATCAG-
90D	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AATAGTCATC	CCATGACAGT	TGGTATCAG-
93C.2	ATCTAAGGTC	GCACGGGTAG	AGGACTTCAA	AATAGTTATC	CCATGACAGT	TGGTATCAG-
93C.3	ATCTAAGGTC	GCACGGGTAG	AGGACTTCAA	AATAGTTATC	CCATGACAGT	TGGTATCAG-
98A	ATCTAAGGTC	GCACGGGTAG	AGGACTTCAA	AATAGTTATC	CCATGACAGT	TGGTATCAG-
126E	GTCTAAGATC	GCATGGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATTAG-
103A	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
50D	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
75A	GTCTAAGGTC	GCACAGGTAG	AGGACTTCAA	AACAGTCATC	CCATGACAGT	TGGTATCAG-
96A	-----	-----	-----	-----TCATC	CCATGACAGT	TGGTCTCAC-
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 181-240

1	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
10	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
14	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
15	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
12	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
17	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATAGTGG	ACATGACAAA	TGTCACCGTG
25	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
26	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATAACAAA	TGTCACCGTG
27C.1	AACAG--TCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
27C.10	AGTAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
28A	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
3	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
41B	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTAG	ACATGACAAA	TGTCACCGTG
45C	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATAACAAA	TGTCACCGTG
5	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGCGG	ACATGACAAA	TGTCACCGTG
101A	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
102B	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	CGTCACCGTG
104	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
108C	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
111A.4	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACCATGGTGG	ACATGACAAA	TGTCACCGTG
111A.5	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
120C	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
123G	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
128C	AGTAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGAAAAA	TGTCACCGTG
131	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
30A	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
57B	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
58B	AGTAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
62D.2	AGTAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGAAAAA	TGTCACCGTG
62D.3	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
65D.7	AGTAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGAAAAA	TGTCACCGTG
65D.3	AGTAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGAAAAA	TGTCACCGTG
67B	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
69A	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
69B	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
76A	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGCTGG	ACATGACAAA	TGTCACCGTG
76E	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
78A.10	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGCTGG	ACATGACAAA	TGTCACCGTG
78A.5	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGCTGG	ACATGACAAA	TGTCACCGTG
8	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
80A	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
82A	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
85A	AGTAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
86B	AGCAGGGTCC	ATATGTGAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
90D	AGCAGGATCG	ATATGTGAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
93C.2	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
93C.3	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
98A	AGCAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
126E	AGTAGGATCG	ATATGTCGGA	CGAGGTGGTA	ACTATGGTGG	ACATGAAAAA	TGTCACCGTG
103A	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
50D	AGCAGGATCG	ATATGTCGGA	CAAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
75A	AGCAGGATCG	ATATGTCAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
96A	AGCAGGATCG	ATATGTGAGA	CGAGGTGGTA	ACTATGGTGG	ACATGACAAA	TGTCACCGTG
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 241-300

1	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
10	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
14	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
15	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
12	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
17	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGA	ATATTGTCCG	TGCGATGAAA
25	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
26	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TACGATGAAA
27C.1	GAGCGTGGTG	GCGAAAAGTC	CAAGAGAGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
27C.10	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
28A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
3	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
41B	GAGCGTGGTG	GYGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
45C	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TACGATGAAA
5	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
101A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TACGATGAAA
102B	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	GGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
104	GAGCGTGGTG	GCGAAAAGTC	CAAGAGTGGG	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
108C	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TACGATGAAA
111A.4	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
111A.5	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGTGATGAAA
120C	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
123G	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
128C	GAGCATGGTG	GCGAAAAGTT	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
131	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
30A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
57B	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	AAATTGTCCG	TGCGATGAAA
58B	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
62D.2	GAGCATGGTG	GCGAAAAGTT	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
62D.3	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
65D.7	GAGCATGGTG	GCGAAAAGTT	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
65D.3	GAGCGTGGTG	GCGAAAAGTT	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
67B	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
69A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
69B	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
76A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
76E	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
78A.10	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
78A.5	GAGCGTGGTG	GCGAAAAGTC	CTAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
8	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
80A	GASCCTGGTG	GCTAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
82A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
85A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
86B	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
90D	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
93C.2	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	AAATTGTCCG	TGCGATGAAA
93C.3	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	AAATTGTCCG	TGCGATGAAA
98A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	AAATTGTCCG	TGCGATGAAA
126E	GAGCATGGTG	GCGAAAAGTT	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
103A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
50D	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
75A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
96A	GAGCGTGGTG	GCGAAAAGTC	CAAGAGGGGA	AGGTCAAAGG	ATATTGTCCG	TGCGATGAAA
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 301-360

1	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
10	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
14	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
15	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
12	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
17	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TAAGGAAGAT
25	GGACGGCTTT	TGAGGATGGA	GAAGGCCATG	GCCTATTTTG	AGACTCAGCT	TGAGGAAGAT
26	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
27C.1	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
27C.10	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
28A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
3	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
41B	GGACGACTTT	TGAGGATGGA	GAAGGCCATG	GCCTATTTTG	AGACTCAGCT	TGAGGAAGAT
45C	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
5	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
101A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
102B	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
104	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
108C	GGGTGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
111A.4	GGACGGCTTT	TGAGGATGGA	GAAGGCCATG	GCCTATTTTG	AGACTCAGCT	TGAGGAAGAT
111A.5	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
120C	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
123G	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
128C	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TTAGGAAGAT
131	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
30A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
57B	GGATGACTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TAAGAAAGAT
58B	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
62D.2	GGACGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TTAGGAAGAT
62D.3	GGACGGCTTT	CAAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TAAGAAAGAT
65D.7	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TTAGGAAGAT
65D.3	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TTAGGAAGAT
67B	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
69A	GGATTGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
69B	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
76A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
76E	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
78A.10	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
78A.5	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
8	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
80A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
82A	GGACGGCTTT	TGAGGATGGA	GAAGGCCATG	GCCTATTTTG	AGACTCAGCT	TGAGGAAGAT
85A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
86B	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCTATTTTG	AGACTCAGCT	TGAGGAAGAT
90D	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
93C.2	GGACGACTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TAAGAAAGAT
93C.3	GGACGACTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TAAGAAAGAT
98A	GGACGACTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TAAGAAAGAT
126E	GGACGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAACT	TTAGGAAGAT
103A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
50D	GGACGGCTTT	TGAGGATGGA	GAAGGCCATG	GCCTATTTTG	AGACTCAGCT	TGAGGAAGAT
75A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
96A	GGATGGCTTT	CGAGGATGGA	GAAGGCCATG	GCCGATTTTG	AGACTCAGCT	TGAGGAAGAT
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 421-480

1	GCACTCACCT	TTCTGGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
10	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
14	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
15	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
12	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
17	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
25	GCACTCACCT	CTCTCGTAGA	CACAATCCAA	ATGGTAGTTA	AGGAGTAGCT	TGACCAGTAC
26	GTACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTG	AGGAGCAGCT	TGACCAGTAC
27C.1	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
27C.10	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
28A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
3	GCACTCACCT	TTCTGGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
41B	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
45C	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
5	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
101A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
102B	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
104	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
108C	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
111A.4	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
111A.5	-----	-----	-----	-----TA	AGGAGCAGCT	TGACCAGTAC
120C	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
123G	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
128C	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
131	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
30A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
57B	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGTT	TGACCAGTAC
58B	GCACTCACCT	TTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
62D.2	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
62D.3	GCACTCACCA	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGTT	TGACCAGTAC
65D.7	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
65D.3	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
67B	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
69A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGGCAGCT	TGACCAGTAC
69B	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
76A	GCACTCACCT	TTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
76E	GCACTCACCT	TTCTGGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
78A.10	GCACTCACCT	TTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
78A.5	GCACTCACCT	TTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
8	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
80A	GCACTCACCT	TTCTGGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
82A	GCACTCGCCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
85A	GCACTCACCT	TTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
86B	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
90D	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
93C.2	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGTT	TGACCAGTAC
93C.3	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGTT	TGATCAGTAC
98A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGTT	TGACCAGTAC
126E	GCACTCACCT	CTGTCCGAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
103A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
50D	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
75A	GCACTCACCT	TTCTGGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
96A	GCACTCACCT	CTCTCGCAGA	CACAATCCAA	ATGGAAGTTA	AGGAGCAGCT	TGACCAGTAC
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 481-540

1	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
10	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
14	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
15	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
12	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
17	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
25	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
26	CTTGCTGAGT	TCACCATAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
27C.1	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
27C.10	CTTGCTGAGT	TCACCACAAT	TCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
28A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
3	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGG	GATTGGGCCT
41B	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
45C	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
5	CTTGCTGAGT	TCACCACAAT	CCGTTGAGGA	GATTGAGTGA	CATGAAAGGA	GATTGGGCCT
101A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGAGCCT
102B	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
104	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
108C	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
111A.4	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
111A.5	CTTTCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGAGCCT
120C	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
123G	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
128C	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
131	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
30A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
57B	CTTGCTGAGT	TCACCATAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
58B	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
62D.2	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
62D.3	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
65D.7	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
65D.3	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
67B	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
69A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
69B	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
76A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
76E	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
78A.10	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
78A.5	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
8	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
80A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
82A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
85A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
86B	CTTGCTGAGT	TCACCACAAT	CTGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
90D	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
93C.2	CTTGCTGAGT	TCACCATAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
93C.3	CTTGCTGAGT	TCACCATAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
98A	CTTGCTGAGT	TCACCATAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
126E	CTTGCTGAGT	TCACCACAAT	CAGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
103A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
50D	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
75A	CTTGCTGAGT	TCACCACAAT	YCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
96A	CTTGCTGAGT	TCACCACAAT	CCGT-GAGGA	GAT-GAGTGA	CATGAAAGGA	GATTGGGCCT
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 541-600

1	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
10	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
14	TATGCAAAAA	GGCAGTGCTC	AACATCTTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
15	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
12	TATGCAAAAA	GGCAGTGCTC	AACA-CTTCC	CATGGGCCTA	AGGAGCC-GA	AGATGGATGC
17	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
25	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGACCTA	GGGAGCC-GA	AGATGGATGC
26	CATGCAAAAA	GGCAGTGCTC	GACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
27C.1	TATGCAAAAA	GGCAGTGCTC	AAAAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
27C.10	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GAGAGCC-GA	AGATGGATGC
28A	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
3	TATGCAAAAA	GGCAGTGAT-	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
41B	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGG-CCTA	GGGAACC-GA	AGATGGATGC
45C	CATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
5	TATGCAAAAA	GGCAGTGCTC	AA---ATTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
101A	CATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
102B	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	AGGAGCT-GA	AGATGGATGC
104	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
108C	CATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
111A.4	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATAGATGC
111A.5	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
120C	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	AGGAGCT-GA	AGATGGATGC
123G	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
128C	TATGCAAAAAC	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
131	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
30A	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
57B	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
58B	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
62D.2	TATGCAAAAAC	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
62D.3	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCT-GA	AGATGGATGC
65D.7	TATGCAAAAAC	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCCCGA	AGATGGATGC
65D.3	TATGCAAAAAC	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
67B	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GAGAGCC-GA	AGATGGATGC
69A	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
69B	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
76A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
76E	TATGTAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
78A.10	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
78A.5	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
8	TATGCAAAAA	GGCAGTGCTC	AA----TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
80A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----GC--GG	AGATGGATGC
82A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
85A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
86B	TATGCAAAAA	GGCAGTGCTC	ACCAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
90D	TATGCAAAAA	GGCAGTGCTC	ACCAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
93C.2	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
93C.3	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
98A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
126E	TATGCAAAAAC	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
103A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
50D	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
75A	TATGCAAAAA	GGCAGTGCTC	AACAA-TTCC	CATGGGC---	----C----	AGATGGATGC
96A	TATGCAAAAA	GGCAGTGCTC	ACCAA-TTCC	CATGGGCCTA	GGGAGCC-GA	AGATGGATGC
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 601-660

1	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGYTG
10	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
14	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
15	TTT--CAAGC	CYA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
12	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
17	TTT--CAAGC	CTA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
25	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
26	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
27C.1	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
27C.10	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
28A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
3	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGKGGAGTTG
41B	TTTTTCAAGC	CCAACCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
45C	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
5	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
101A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
102B	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CAAGGAGCTG
104	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
108C	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
111A.4	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
111A.5	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAAGAGCTG
120C	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CAAGGAGCTG
123G	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGGC	GATGGCACCT	CGAGGAGCTG
128C	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGACACCC	CGAGGAGCTG
131	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
30A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
57B	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
58B	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATAGCACCC	CGAGGAGCTG
62D.2	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGACACCC	CGAGGAGCTG
62D.3	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
65D.7	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGACACCC	CGAGGAGCTG
65D.3	TCT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGACACCC	CGAGGAGCTG
67B	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGCC	GATGGCACCC	CGAGGAGTTG
69A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
69B	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GACGGCACCC	CGAGGAGCTG
76A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
76E	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
78A.10	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
78A.5	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
8	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
80A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	ACTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
82A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAT	GATGGCACCC	CGAGGAGCTG
85A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATAGCACCC	CGAGGAGCTG
86B	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGTTG
90D	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
93C.2	TTT--CAAGC	CCA-CCTATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
93C.3	TTT--CAAGC	CCA-CCTATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
98A	TTT--CAAGC	CCA-CCTATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
126E	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGTTG
103A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
50D	TTT--CAAGC	CCA-CCCATG	CAATGTACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTA
75A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
96A	TTT--CAAGC	CCA-CCCATG	CAATGCACAA	GCTTGTTCGAC	GATGGCACCC	CGAGGAGCTG
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----	-----	-----	-----

Bases 661-720

1	CGATGATTTC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
10	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	T-GAAATGGT	GGATGAGAGA
14	CGATGATTCC	CAAATGTTGC	CTTTGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
15	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
12	CGATGATTTC	CAAATGTTGC	CTTTGMAATTT	GAW--GCCTT	C-GAAATGGT	GGATGAGAGA
17	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
25	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
26	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
27C.1	CGATGATTCC	CAAATGTTGC	CTTCGAA---	-----	-----ATGGT	GGATGAGAGA
27C.10	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
28A	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	T-GAAATGGT	GGATGAGAGA
3	CGATGATTTC	CAAATGTTGC	CTTAGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
41B	TGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
45C	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
5.11	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
101A	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
102B	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
104	CAATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
108C	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGG
111A.4	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
111A.5	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATTAGAGA
120C	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
123G	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
128C	CGATGATTCC	TAAATGTTTC	CTTCGGATTT	CTT--GCCTT	C-AAAATGGT	GGATGAGCGG
131	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	T-GAAATGGT	GGATGAGAGA
30A	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
57B	TGATGATTCC	CAAATGTTGC	CTTCAGATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGG
58B	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
62D.2	CGATGTTTCC	CAAATGTTTC	CTTCAGATTT	CTT--GCCTT	C-AAAATGGT	GGATGAGCGG
62D.3	CGATGATTCC	CAAATGTGGC	CTTCAGATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGG
65D.7	CGATGATTCC	TAAATATTTT	CTTCGGATTT	CTT--GCCTT	C-AAAATGGT	GGATGAGCGG
65D.3	CGATGATTCC	TAAATATTTT	CTTCGGATTT	CTT--GCCTT	C-AAAATGGT	GGATGAGCGG
67B	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTCTCGCCTT	CCGAAATGGT	GCATGAGAGA
69A	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
69B	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
76A	CGATGATTCC	CAAATGTTGC	CTTCAAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
76E	CGATGATTCC	CAAATGTTGC	CTTCAAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
78A.10	CGATGATTCC	CAAATGTTGC	CTTCAAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
78A.5	CGATGATTCC	CAAATGTTGC	CTTCAAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
8	CAATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
80A	CGATGATTTC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
82A	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
85A	CGATGATTCC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
86B	CGATGATTCC	CAAATGTTGC	CTTTGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
90D	CGATGATTCC	CAAATGTTGC	CTTTGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
93C.2	TGATGATTCC	CAAATGTTGC	CTTCAGATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGG
93C.3	TGATGATTCC	CAAATGTTGC	CTTCAGATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGG
98A	TGATGATTCC	CAAATGTTGC	CTTCAGATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGG
126E	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	CTT--GCCTT	C-AAAATGGT	GGATGAGCGG
103A	CAATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
50D	CGATGATTCC	CAAATGTTGC	CTTCGGATTT	GTT--GCCTT	C-AAAATGGT	GGATAAGCGG
75A	CGATGATTYC	CAAATGTTGC	CTTCGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
96A	CGATGATTGC	CAAATGTTGC	CTTTGAATTT	GTT--GCCTT	C-GAAATGGT	GGATGAGAGA
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	-----	-----	-----TG-	GTT--GCCTT	C-AGAATGGT	GGATGTGCGG

Bases 721-780

1	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
10	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
14	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
15	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
12	CGAATTGGAT	TGCCTATGCC	TGMTTCTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
17	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
25	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GAGCAATGCA	CAAGCTTGTC
26	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
27C.1	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	AAGCCCTTGC	GGGCAGTGCA	CAAGCTTGTC
27C.10	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
28A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
3	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
41B	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
45C	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
5	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
101A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
102B	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
104	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
108C	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
111A.4	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	AAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
111A.5	CGAATTGGAT	TGCGTATGGT	TGGTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
120C	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
123G	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
128C	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
131	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
30A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAT	GGGCAGTGCA	CAAGCTTGTT
57B	CAAATTGGAT	TGCCTATGCT	TGCCTATGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
58B	TGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
62D.2	CGAATTGGAT	TGCCTATGCT	TGCTTATGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
62D.3	CAAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
65D.7	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
65D.3	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
67B	CGAATTGGAT	TGCCTATGCT	TGGTGTGCG	GAGCCATTAC	GGGCAGTGCA	CAAGCTTGTC
69A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGTA	CAAGCTTGTT
69B	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGTA	CAAGCTTGTT
76A	CGAATTGGAT	TGCCTATGCT	TGCTTGTACG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
76E	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
78A.10	CGAATTGTAT	TGCCTATGCT	TGCTTGTACG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
78A.5	CGAATTGTAT	TGCCTATGCT	TGGTGTACG	GAGCCCTTAC	TGGCAGTGCA	CAAGCTTGTC
8	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTT
80A	CGAATTGGAT	TGCCTATGCT	TGCTTGTACG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
82A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAACGCG	CAAGCTTGTC
85A	TGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
86B	CCAATTGGAT	TGCCTATGTT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
90D	CCAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
93C.2	CAAATTGGAT	TGCCTATGCT	TGCTTATGCA	GAGCCCTTAT	GGGTAATGCA	CAAGCTTGTC
93C.3	CAAATTGGAT	TGCCTATGCT	TGCTTATGCA	GAGCCCTTAC	GGGTAATGCA	CAAGCTTGTC
98A	CAAATTGGAT	TGCCTATGCT	TGCTTATGCA	GAGCCCTTAC	GGGTAATGCA	CAAGCTTGTC
126E	CAAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
103A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
50D	CGAATTGGAT	TGCCTATGCA	TGCTTGTGCG	GAGCCCTTAC	GGGCAATGCA	CAAGCTTGTC
75A	CGAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
96A	CCAATTGGAT	TGCCTATGCT	TGCTTGTGCG	GAGCCCTTAC	GGGCAGTGCA	CAAGCTTGTC
Pur-132	-----	-----	-----	-----	-----	-----
Pur-34	CGAATCGGAT	TGCTTATGCT	TGCTTGGCG	GAGCCTTGT	GGGCAATGCA	CAAGCATGTC

Bases 781-840

1	GACGATCGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CRGGTGCTTG
10	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
14	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
15	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTRCTTG
12	GACGATGGCA	CCCCGAGGAG	TTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
17	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGCMTTGG
25	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
26	GACAATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
27C.1	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
27C.10	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
28A	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
3	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
41B	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
45C	GACAATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
5	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTGT	CGGGTGCTTG
101A	GACAATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
102B	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
104	GACAATGGCA	CCCTGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
108C	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
111A.4	GACAATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
111A.5	GACGATGGCA	CCCCGAGGAG	CTGCGACGAT	TCCCAAATGA	CCCTCACTCT	TGGGTGCTTG
120C	GCCCATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
123G	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
128C	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
131	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
30A	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
57B	GACGATGGCA	CCCCGAGGAG	CTGTGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
58B	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
62D.2	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
62D.3	GATGATGGCA	CCCCGAGGAG	CTACGATGAT	TCTCAAATGA	CCCTCACTCT	CGGGTGCTTG
65D.7	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
65D.3	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
67B	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
69A	GACGATGGCG	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
69B	GACGATGGCG	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
76A	GACGATCGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
76E	GACGATCGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
78A.10	GACGATCGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
78A.5	GACGATCGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
8	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
80A	GACGATCGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
82A	GACAATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
85A	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
86B	GACGATGGCA	CCCCGAAGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
90D	GACGATGGCA	CCCCGAAGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
93C.2	GACGATGGCA	CCCCGAGGAG	CTACGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
93C.3	GACGGTGGCA	CCCCGAGGAG	CTACGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
98A	GACGATGGCA	CCCCGAGGAG	CTACGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
126E	GACTATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
103A	GACAATGGCA	CCCTGAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
50D	GACAATGGCA	CCCCAAGGAG	CTGCGATGAT	TCCTAAATGA	CCCTCACTCT	TGGGTGCTTG
75A	GACGATGGCA	CCCCGAGGAG	CTGCGATGAT	TCCCAAATGA	CCCTCACTCT	CGGGTGCTTG
96A	GACGATGGCA	CCCCGAAGAG	-----	-----	-----	-----
Pur-132	-----	-----	-----	----A---GA	CTAGCATTGT	TAGGTTTTTG
Pur-34	GACGATGGCA	CCTCGAGCAG	CCGCGATGAC	ATCCA---GA	CCTGCATTGT	TGGGTTTTTG

Bases 841-900

1	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
10	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
14	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
15	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
12	TGAAGGTGCA	ATGGGTGCTA	ATCGATGTCG	TATATGCRGA	ACATACAAAG	GGCGCGGGGG
17	TGAAGGTGCA	TTGGKTGCTA	CTCGATMTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
25	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCAGA	ACATACAAAG	GGTGCAGGGGG
26	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
27C.1	TGAAGGTGCA	TTGGTTGCTA	CTTGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
27C.10	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
28A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
3	TGAAGGTGCA	TTGGTTGCTA	ATCGTCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
41B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TGTATGTAGA	ACATACAAAG	GGTGCAGGGGG
45C	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
5	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
101A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
102B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
104	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
108C	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
111A.4	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
111A.5	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CATATGCAGA	ACATACAAAG	GGTACGGGGG
120C	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
123G	TGAAGGTGCA	TTGGTTGGTA	CTCGATCTTT	TTTATGCGGA	ACATACAAAG	GGSGGGGGGG
128C	TGAAGGTGCA	TTGGTTCCTA	CTCGATCTCT	CGTATGCGGA	ACATACA---	-----
131	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
30A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
57B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGAGGGGG
58B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
62D.2	TGAAGGTGCA	TTGGTTCCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGCGGGGG
62D.3	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCCGA	ACATACAAAG	GGCGCGGGGG
65D.7	TGAAGGTGCA	TTGGTTCCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGCGGGGG
65D.3	TGAAGGTGCA	TTGGTTCCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGCGGGGG
67B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
69A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
69B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
76A	TGAAGGTGCA	TTGGTTGCCA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
76E	TGAAGGTGCA	TTAGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
78A.10	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
78A.5	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
8	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
80A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTTT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
82A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCAGA	ACATACAAAG	GGTGCAGGGGG
85A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
86B	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCAGA	ACATACAAAG	GGCGCGGGGG
90D	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	TCTATGCAGA	ACATACAAAG	GGCGCGGGGG
93C.2	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGAGGGGG
93C.3	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGAGGGGG
98A	TGAAGGTGCA	TTGGTTGCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGAGGGGG
126E	TGAAGGTGCA	TTGGTTCCTA	CTCGATCTCT	CGTATGCGGA	ACATACAAAG	GGCGCGAGGG
103A	TGAAGGTGCA	TTGGTTGCTA	CTCAATCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
50D	TGAAGGTGCA	TTGGTTGCTA	YTGATCTCT	TATATGCGGA	ACATACAAAG	GGTGCAGGGGG
75A	TGAAGGTGCA	TTGGTTGCTA	CTCGRTCTCT	TCTATGCGGA	ACATACAAAG	GGCGCGGGGG
96A	-----	-----	-----	-----	-----	-----
Pur-132	TGTCGGTCCA	TGTGTTGTCA	CTCGATCTCT	CGGATGCGGA	ACATGTTGCG	GGTGT-AGGT
Pur-34	TGTCGGTCCA	CGTGTGTTCA	CTCGATCTCT	CGGATGCAGA	ACATGTTGCG	GGCGC-AGGG

Bases 901-960

1	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTT-GT	CCCTATACAC	GAACGACAGT
10	TCTTCAGCCC	CY-GAMGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
14	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
15	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
12	TTTTTCAGCCC	CC-GACGTC-	AGATGCAAAW	-GCATTTTGT	CCCTAKACAC	GAACGACAGT
17	TCTTTAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGGCAGT
25	TCTTCAGCCC	TC-GACGTCC	-GATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
26	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
27C.1	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATAC--	-----
27C.10	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
28A	TCTTCAGCCC	CT-GAMGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
3	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
41B	TCTTCAGCCC	CC-GACGTCC	-AATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
45C	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
5	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
101A	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
102B	TCCTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
104	TCTTCAGCCC	CT-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
108C	TCTTCAGCCC	CC-GACGTC-	AGAAGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
111A.4	TCTTCCGCCC	CC-GACGTC-	AGACGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
111A.5	TCTTCAGCCC	CC-GACGTCC	-GATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
120C	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
123G	T-----	-----	-----	-----	-----	-----
128C	-----	-----	-----	-----	-----	-----
131	TCTTCAGCCC	CT-GAAGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
30A	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
57B	TCTTCAGCCC	TC-GATGTCC	-GATGCAAAA	C-CATTTTGT	CCCTATACAC	GAATGATAGT
58B	TCTTCAGCCC	CC-GACGTC-	AGATGTA AAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
62D.2	TCTTCAGCCC	CC-GACGTCC	-GATGCAAAA	C-CATTTTGT	CCCTACACAA	GAACGACAGT
62D.3	TTTTTCAGCCC	CC-GACGTCC	-GATGCAAAA	C-CATTTTGT	CCCTATACAC	GAATGACAGT
65D.7	TCTTCAGCCC	CC-GACGTCC	-GATGCAAAA	C-CATTTTGT	CCCTACACAA	GAACGACAGT
65D.3	TCTTCAGCCC	CC-GACGTCC	-GATGCGAAA	C-CATTTTGT	CCCTACACAA	GAACGACAGT
67B	TCTTCAGCCC	CA-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
69A	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
69B	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
76A	TCTTCAGCCC	CN-TACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
76E	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
78A.10	TCTTCAGCCC	CC-TACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
78A.5	TCTTCAGCCC	CC-TACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
8	TCTTCAGCCC	CC-GACGTC-	AGAT-CAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
80A	TCTTCAGCCT	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
82A	TCTTCAGCCC	TC-GACGTCC	-GATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
85A	TCTTCAGCCC	CC-GACGTC-	AGATGTA AAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
86B	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
90D	TCTTCAGCCC	CC-GACGTC-	AAATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
93C.2	TCTTCAGCCC	TC-GATGTCC	-GATGCAAAA	C-CATTTTGT	CCCTATACAC	GAATGACAGT
93C.3	TCTTCAGCCC	TC-GATGTCC	-GATGCAAAA	C-CATTTTGT	CCCTATACAC	GAATGACAGT
98A	TCTTCAGCCC	TC-GATGTCC	-GATGCAAAA	C-CATTTTGT	CCCTATACAC	GAATGACAGT
126E	TCTTCAGCCC	CC-GACGTCC	-GATGCAAAA	AGCATTTTGT	CCCTATACAC	GAACGACAGT
103A	TCTTCAGCCC	CT-GACGTCC	AGATGCAAAA	-GCATTTTGT	CCCTATACAC	GAACGACAGT
50D	TCTTCAGCCC	CM-GACGTC-	AGAWGCAAGA	-GCATTTTGT	CCCTATACA-	-AACA----T
75A	TCTTCAGCCC	CC-GACGTC-	AGATGCAAAA	-GCATTTTGT	CCCTATMCAC	GAACGACAGT
96A	-----	-----	-----	-----	-----	-----
Pur-132	TCTTCGGCCC	TTTGACGTCC	AAA-ACAAAG	CGTTTTTCG-	--CTAGACAC	GAATGACTGT
Pur-34	TCTTCGGCCC	CTTGACGTCC	AAA-ACAAAG	CGTTTTTYCG-	--CTAGACAC	GAACGACAGT

Bases 961-977

1 CGCGCGC-GT TGCAGGC
 10 CGCGCGC-GT TGCAGGC
 14 CGCGCGC-GT TGCAGGC
 15 CGCGCGC-GT TGCAGGC
 12 TGCGCGCTGT TGCACGG
 17 CGCGCGC-GT TGCAGGC
 25 CGCGCGC-GT TGCAGGC
 26 CGCGCGC-GT TGCAGGC
 27C.1 -----
 27C.10 CGCGCGC-GT TGCAGGC
 28A SGCGCGC-GT TGCAGGC
 3 CGCGCGC-GT TGCAGGC
 41B CGCGCGC-GT TGCAGGC
 45C CGCGCGC-GT TGCAGGC
 5 CGCGCGC-GT TGCAGGC
 101A CGCGCGC-GT TGCAGGC
 102B CGCGCGC-GT TGCAGGC
 104 CGCGCGC-GT TGCAGGC
 108C CGCGCGC-GT TGCAGGC
 111A.4 CGCGCGC-GT TGCAGGC
 111A.5 CGCGCGC-GT TGCAGGC
 120C CGCGCGC-GT TGCAGGC
 123G -----
 128C -----
 131 GGCGCGC-GT TGCAGGC
 30A CGCGCGC-GT TGCAGGC
 57B TGTGCGC-GT TGCAGGC
 58B CGCGCGC-GT TGCAGGC
 62D.2 CGCGCGC-GT TGCAGGC
 62D.3 CGTGCGC-GC TGTAGGC
 65D.7 CGCGCGC-GT TGCAGGC
 65D.3 CGCGCGC-GT TGCAGGC
 67B CGCGCGC-GT TGCAGGC
 69A CGCGCGC-GT TGCAGGC
 69B CGCGCGC-GT TGCAGGC
 76A CGCGCGC-GT TGCAGGC
 76E CGCGCGC-GT TGCAGGC
 78A.10 CGCGCGC-GT TGCAGGC
 78A.5 CGCGCGC-GT TGCAGGC
 8 CGCGCGC-GT TGCAGGC
 80A CGCGCGC-GT TGCAGGC
 82A CGCGCGC-GT TGCAGGC
 85A CGCGCGC-GT TGCAGGC
 86B CGCGCGC-GT TGCAGGT
 90D CGCGCGC-GT TGCAGGT
 93C.2 TGTGCAC-GT TGCAGGC
 93C.3 TGTGCAC-GT TGCAGGC
 98A TGTGCAC-GT TGCAGGC
 126E CGCGCGC-GT TGCAGGC
 103A CGCGCGC-A- -----
 50D -A-----T T-----
 75A CGCGCGC-GT TGCAGGC
 96A -----
 Pur-132 CGCGTAC-GT TGCAGGC
 Pur-34 CGCGTAC-GT TGCAGGC

Appendix E. Chloroplast DNA sequence data

This dataset contains the aligned sequences for the *trnS* and *trnG* intergenic spacer and the *rp120* and *rps12* intergenic spacer of the chloroplast genome used in Chapter 4. The sequences of the two regions have been combined for a total of 1337 bp. Sequence positions 1 - 580 bp represent the *trnS* and *trnG* intergenic spacer and positions 581 - 1337 represents the *rps12-rp120* intergenic spacer. 14 indels of 2 bp or more are coded in positions 1338-1351. ID codes for the sequences correspond to those found in Table 4.1. Gaps in the sequence are indicated by a dash (-). Uncertainties are denoted by standard IUPAC/IUB codes as follows: N=unknown; R=A/G; Y=C/T; M=A/C; K=G/T; S=C/G; W=A/T.

Bases 1-60

1	-----	-----	-----	-----	CATT	CATTTCAATTT	T--AAAAACT
10	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
101A	TGG-AACTTT	TGTCGAACAA	GGAGATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
102B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
103A	-----TTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
104	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
108C	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
111A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	TT-AAAAACT
12	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
120C	TGG-AACTTT	TGTCGAACAA	GGAGATAGGA	AAAGA-CATT		CATTTCAATTT	T--AAAAACT
121A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	---AAAAACT
123E	TGGCAACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGA-CATT		CATTTCAATTT	T--AAAAACT
123G	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
126E	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
128C	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
129	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTAATTT	A--ATAACT
131	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
132	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTAATTT	A--AAAAACT
134A	T-----	-----	-----	AGGA	AAAGAGCATT	CATTTCAATTT	T--AAAAACT
14	TGGCAACTTT	TGTCGAACAA	GAA-ATAGGG	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
15	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
25	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTAATTT	T--AAAAACT
26	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--ATAACT
27C	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
28A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	A--AAAAACT
3	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T-AAAAAACT
30A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
36C	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTAATTT	A--AAAAACT
41B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
45C	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	TTAAAAAACT
5	-----TTT	TGTCGAACAA	GAA-ATAGGA	AA-GAGCAT-		CATTTCAATTT	T--AAAAACT
50D	-----TTT	TGTCGAACAA	GAG-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
57B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T-AAAAAACT
58B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	TTAAAAAACT
62D	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
65D	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
67B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
69A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T-AAAAAACT
69B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
75A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
76A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
76E	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--ATAACT
77B	-----TTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
8	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
80A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
81A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
82A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
84F	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
86B	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
93C	TGG-AACTTT	TGTCGAACAA	GAA--TAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
98A	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
C.pri	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
S.3	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT
S.7	TGG-AACTTT	TGTCGAACAA	GAA-ATAGGA	AAAGAGCATT		CATTTCAATTT	T--AAAAACT

Bases 181-240

1	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
10	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
101A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
102B	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
103A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
104	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
108C	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
111A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
12	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
120C	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
121A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
123E	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
123G	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
126E	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AAAGAATAAA	CTAAATAAAT
128C	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAAA	CTAAATAAAT
129	TT-AC(TTTT	TACTTT----	-----A	CTGGAATAAA	AAA-AATAAA	CTAAATTAAT
131	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
132	TTTACTTTT	-ACTTC----	-----A	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
134A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAA--
14	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
15	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
25	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
26	TTTA----TT	TACTTT----	-----A	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
27C	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
28A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
3	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
30A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
36C	TTTACTTTT	TACTTTTAT	TTCACTGGAA	TAGGAATAAA	AA-GAATAA	TTAAATAAAT
41B	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
45C	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
5	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
50D	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
57B	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
58B	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
62D	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
65D	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
67B	TTTA----TT	TACTTT----	-----A-	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
69A	TTTA----TT	TACTTT----	-----A-	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
69B	TTTA----TT	TACTTT----	-----A-	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
75A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
76A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAA--
76E	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAA--
77B	TTTA----TT	TACTTT----	-----A-	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
8	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
80A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
81A	TTTA----TT	TACTTT----	-----AA	CTGGAATAA	AAAGAATAA	CTAAATAA--
82A	TTTA----TT	TACTTT----	-----A-	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
84F	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
86B	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
93C	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
98A	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
C.pri	TTTA----TT	TACTTT----	-----A-	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
S.3	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT
S.7	TTTA----TT	TACTTT----	-----AA	CTGGAATAAA	AA-GAATAA	CTAAATAAAT

Bases 361-420

```
1      GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
10     GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
101A   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
102B   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
103A   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
104     GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
108C   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
111A   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
12      GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGC---A
120C   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
121A   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
123E   GAGTGTTC  GTTAAATTAT TAAATTAAT AAATCATTTG ATTTACGATT CGTTGG---A
123G   GAGTGTTC  GTTAAATTAT TAAATTAAT AAATCATTTG ATTTACGATT CGTTGG---A
126E   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
128C   GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT C-TTGG---A
129     TAGTTTTT  CA-----ATTA- -----AAT AAATCATTTA ATTTATGATT CGTTGGAACA
131     GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CCTTGG---A
132     TAGTGTTC  CAATTAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
134A   GAGTGTTC  GTTAAATTAA -ATATTAAT AAATCATTTT ATTTATGATT CGTTGG---A
14      GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
15      GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CCTTGG---A
25      GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
26      GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
27C    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGC--CA
28A    GAGTGTTC  GTTAAATT-- -----AAAT AAATCATTTG ATTTACGATT CGTTGG---A
3       GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
30A    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
36C    TAGTGTTC  CAATTAATTA- -----AAA AAATCATTTA ATTTACGATT CGTTGG---A
41B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
45C    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
5       GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
50D    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
57B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CCCTCC---A
58B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
62D    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
65D    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
67B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
69A    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
69B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
75A    GAGTGTTC  GTTAAATTAA -ATATTAAT AAATCATTTT ATTTACGATT CGTTGG---A
76A    GAGTGTTC  GTTAAATTAA -ATATTAAT AAATCATTTT ATTTACGATT CGTTGG---A
76E    GAGTGTTC  GTTAAATTAA -ATATTAAT AAATCATTTT ATTTACGATT CGTTGG---A
77B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
8       GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
80A    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
81A    GAGTGTTC  GTTAAATTAA -ATATTAAT AAATCATTTT ATTTACGATT CGTTGG---A
82A    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
84F    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
86B    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
93C    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
98A    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
C.pri  GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTT ATTTACGATT CGTTGG---A
S.3    GAGTGTTC  GGTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
S.7    GAGTGTTC  GTTAAATTA- -----AAT AAATCATTTG ATTTACGATT CGTTGG---A
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Bases 421-480

1	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
10	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
101A	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
102B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
103A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTCC	AGTACTTAGG	CCGGGCCGTG	GAACTAAAAA
104	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
108C	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
111A	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
12	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
120C	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
121A	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
123E	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
123G	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
126E	ACAAAAAGGG	CCGGTCCCG	GCC-TGGTCC	AGTATTTAG-	-CCGGCCGTG	GAACTAAAAA
128C	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
129	AAAAAAAAAG	GGCGGGCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
131	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	CAAGTAAAAA
132	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTATTTAG-	CCGGGCCGTG	GAACTAAAAA
134A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
14	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
15	ACAAAAAGGG	C-GG-GCCCG	CCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
25	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
26	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
27C	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
28A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
3	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
30A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
36C	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTATTTAG-	CCGGGCCGTG	GAACTAAAAA
41B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
45C	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
5	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
50D	ACAAAAAGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAAG
57B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
58B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
62D	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
65D	ACAAAAAGGG	C-GG-TCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
67B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
69A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
69B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
75A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
76A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
76E	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
77B	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
8	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
80A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
81A	ACAAAAAGGG	CCGGGCCCGG	GCCCTGGTCC	AGTATTTAGC	CCGGGCCGTG	GAACTAAAAA
82A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
84F	ACAAAAAGGC	G-GG--CCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
86B	ACAAAAAGGG	C-GG--CCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
93C	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
98A	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
C.pri	ACAAAAAGGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
S.3	ACAAAAAGGG	C-GG--CCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA
S.7	ACAAAAAGG	C-GG-GCCCG	GCC-TGGTC-	AGTACTTAG-	CCGGGCCGTG	GAACTAAAAA

Bases 481-540

1	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
10	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTG	TCAAGCCTTA
101A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
102B	GCCCCCTTCG	GACGAAATCC	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
103A	GCCCCCTTCG	GACGAAATCC	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
104	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
108C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
111A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
12	GCCCCCTTCG	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
120C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
121A	GCCCCCTTC-G	-ACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
123E	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
123G	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
126E	GCCCCCTTCG	GACGAAATCC	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
128C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
129	GCCCCCTTC-G	GACGAAATCA	AAAAA---G	AGTGTATGA	AGGG-CTTTT	TCAAGCCTTA
131	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
132	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
134A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
14	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTG	TCAAGCCTTA
15	GCCCCCTTCG	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGGCTTTT	TCCACCCTTA
25	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAG-CTTA
26	GCCC-TTC-G	GGCGAAATCA	AAAAA---G	AATATTAT-A	AGGG-CTTTT	TCAAGC-TTA
27C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
28A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
3	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
30A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
36C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
41B	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTT-	TCAAGCCTTA
45C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTT-	TCAAGCCTTA
5	GCCC-TTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGC-TTA
50D	GCCCCCTTC-G	-ACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
57B	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
58B	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
62D	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
65D	GCCC-TTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
67B	GTCCCTTC-G	GGCGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
69A	GTCCCTTC-G	GGCGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
69B	GTCC-TTC-G	GGCGAAATC-	ATAAAA---G	AATATTATGA	AGGG-CTTTT	TCCACCCTTA
75A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
76A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
76E	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
77B	GTCC-TTCG	GGCGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTGT	TCAAGCCTTA
8	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
80A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
81A	GCCCCCTTCG	GACGAAATCC	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
82A	GTCCCTTCG	GGCGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
84F	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
86B	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
93C	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
98A	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
C.pri	GTCCCTTC-G	GGCGAAATC-	AAAAAA---T	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
S.3	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA
S.7	GCCCCCTTC-G	GACGAAATC-	AAAAAA---G	AATATTATGA	AGGG-CTTTT	TCAAGCCTTA

Bases 541-580

```
1      TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
10     TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
101A   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTAT---
102B   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
103A   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
104     TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
108C   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATTTA
111A   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATTTA
12      TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
120C   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTAT---
121A   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
123E   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
123G   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
126E   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
128C   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
129     TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAG-----
131     TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
132     TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTCTT--
134A   TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
14      TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
15      TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
25      TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
26      TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
27C    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
28A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
3       TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
30A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
36C    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTCTT--
41B    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
45C    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
5       TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
50D    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
57B    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTAT---
58B    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
62D    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
65D    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
67B    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
69A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
69B    TTTTTTATAA TGTAACCT-A TAATGTAATC ATAG-----
75A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
76A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
76E    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
77B    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
8       TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
80A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
81A    TTTTTTATAA TGTAACCT-A TAATGTAA-C AT-----
82A    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
84F    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
86B    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
93C    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
98A    TTTTTTATAA TGTAACCTTA TAATGTAA-C ATAGTATT--
C.pri  TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAG-----
S.3    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
S.7    TTTTTTATAA TGTAACCT-A TAATGTAA-C ATAGTATT--
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Bases 581-640

1	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAG--	GAATAGGA-T	AGAATGGAAG
10	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
101A	AAAACAATTA	-ATCCA-TAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
102B	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
103A	-----	-----TAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
104	-----	-----TAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
108C	AAAACAATTG	-ATCCAGTAT	CAACGGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
111A	AAAACAATTA	-ATCCA-TAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
12	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
120C	AAAACAATTA	-ATCCAGTAT	CAACGGATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
121A	AAAACAATTA	-ATCCAGTAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
123E	AAAACAATTA	-ATCCAGTAT	CAACG-ATCA	G-TACCAG--	GAATAGGA-T	AGAATGGAAG
123G	AAAACAATTA	-ATCCA-TAT	CAA-GGATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG
126E	AAAACA-TTA	-ATCCAGTAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
128C	AAAACAATTA	-ATCCAGTAT	CAACGGATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG
129	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG
131	AAAACAATTG	TATCCAGTAT	CAACGGATCA	G-TACCAGTG	GAATAGGA-T	AGAA-GGAAG
132	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG
134A	----CAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
14	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
15	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGAAT	AGAATGGAAG
25	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
26	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAG--	GAATAGGA-T	AGAATGGAAG
27C	-----	-----	-----	-----	-----	-----
28A	AAAACAATTG	-ATCCA-TAT	CAA-GGATCA	G-TACCAG--	-AATAGGA-T	AGAA-GGAAG
3	-----	-----	-----	-----	-----	-----
30A	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG
36C	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
41B	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
45C	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
5	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAG--	GAATAGGA-T	AGAATGGAAG
50D	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
57B	AAAACAATTG	-ATCCA-TAT	CAA-GGATCA	G-TACCA---	GAATAGGA-T	AGAA-GGAAG
58B	AAAACAATTG	-ATCCA-TAT	CAA-GGATCA	GCTACCAGT-	GAATAGGA-T	AGAAGGGAAG
62D	AAAACAATTA	-ATCCAGTAT	CAACGGATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
65D	AAAACAATTA	-ATCCAGTAT	CAACGGATCA	G-TACCAG--	GAATAGGA-T	AGAAGGGAAG
67B	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCA-T-	GAATAGGA-T	AGAATGGAAG
69A	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
69B	AAAACAATTG	TATCCAGTAT	CAA--GATCA	G-TACCAG--	-AATAGGA-T	AGAA-GGAAG
75A	AAAACA-TTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
76A	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
76E	-----	-----	-----	-----	-----	-----
77B	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAG--	GAATAGGA-T	AGAATGGAAG
8	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
80A	AAAACAATTG	-ATCCAGTAT	CAACGGATCA	G-TACCAGTG	GAATAGGA-T	AGAAGGGAAG
81A	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
82A	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCAGT-	GAATAGGA-T	AGAA-GGAAG
84F	AAAACAATTG	-ATCCA-TAT	CAACG-ATCA	G-TACCAGT-	GAATAG--AT	AGAATGGAAG
86B	AAAACAATTG	-ATCCAGTAT	CAA-G-ATCA	G-TACCAG--	-AATAGGA-T	AGAA-GGAAG
93C	AAAACAATTG	-ATCCAGTAT	CAA-GGATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG
98A	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
C.pri	AAAACAATTG	-ATCCAGTAT	CAACG-ATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
S.3	AAAACAATTG	-ATCCAGTAT	CAAC-GATCA	G-TACCAGT-	GAATAGGA-T	AGAATGGAAG
S.7	AAAACAATTG	-ATCCAGTAT	CAA-G-ATCA	G-TACCAG--	GAATAGGA-T	AGAA-GGAAG

Bases 641-700

1	A	A	A	-	G	T
10	A	A	A	-	G	T
101A	A	A	A	-	G	T
102B	A	A	A	-	G	T
103A	A	A	A	-	G	T
104	A	A	A	-	G	T
108C	A	A	A	-	G	T
111A	A	A	A	-	G	T
12	A	A	A	-	G	T
120C	A	A	A	-	G	T
121A	A	A	A	-	G	T
123E	A	A	A	-	G	T
123G	A	A	A	-	G	T
126E	A	A	A	-	G	T
128C	A	A	A	-	G	T
129	A	A	A	G	T	T
131	A	A	A	-	G	T
132	A	A	A	G	T	T
134A	G	-	A	-	G	T
14	A	A	A	-	G	T
15	A	A	A	-	G	T
25	A	A	A	-	G	T
26	A	A	A	-	G	T
27C	-	-	A	-	G	T
28A	A	A	A	-	G	T
3	-	-	A	G	T	T
30A	A	A	A	-	G	T
36C	A	A	A	-	T	T
41B	A	A	A	-	G	T
45C	A	A	A	-	G	T
5	A	A	A	-	G	T
50D	A	A	A	-	G	T
57B	A	A	A	-	G	T
58B	A	A	A	-	G	T
62D	A	A	A	-	G	T
65D	A	A	A	-	G	T
67B	A	A	A	-	G	T
69A	A	A	A	G	T	T
69B	A	A	A	-	G	T
75A	A	A	A	G	T	T
76A	A	A	A	-	G	T
76E	-	-	A	-	G	T
77B	A	A	A	-	G	T
8	A	A	A	-	G	T
80A	A	A	A	-	G	T
81A	A	A	A	-	G	T
82A	A	A	A	-	G	T
84F	A	A	A	-	G	T
86B	A	A	A	-	G	T
93C	A	A	A	-	G	T
98A	A	A	A	-	G	T
C.pri	A	A	A	-	G	T
S.3	A	A	A	-	G	T
S.7	A	A	A	-	G	T

Bases 1001-1060

1	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTT-	AAGAAGTAAC	CATAGAAA-C
10	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
101A	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
102B	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCCCCGTTT	AAGAAGTAAC	CATAGAAA-C
103A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
104	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
108C	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
111A	GAAGAAGGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
12	GAAAGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
120C	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
121A	GAAG-AAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
123E	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
123G	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
126E	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
128C	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
129	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATATAAA-C
131	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
132	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
134A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
14	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
15	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
25	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
26	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
27C	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
28A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCCCCGTTT	AAGAAGTAAC	CATAGAAA-C
3	GAAG-AAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
30A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
36C	GAAAGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
41B	-AAGGAAGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
45C	GAAG-AAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
5	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
50D	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
57B	GAAG-AAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
58B	GAAG-AAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
62D	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAC-C
65D	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
67B	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
69A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
69B	GAAG-AA-GG	CTC-GGTGT	TAGAGAGGAC	-TCACCGTTT	AAGAAGTAAC	CATAGAAA-C
75A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
76A	GAAG-AA-GG	CTCCG-TGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
76E	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
77B	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
8	GAAAGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTACC	CATAGAAA-C
80A	GAAGGAAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACGGTTT	AAGAAGTAAC	-ATAGAAA-C
81A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
82A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
84F	GAAAGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
86B	GAAGAAGGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
93C	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
98A	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
C.pri	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
S.3	GAAG-AAGG-	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C
S.7	GAAGGAAGGG	CTCCGGTGT	TAGAGAGGAC	CTCACCGTTT	AAGAAGTAAC	CATAGAAA-C

Bases 1061-1120

1	GATGGAACCC	ACTATATCCA	TT-ATATTAA	CTTTACTACT	TTTTAATTTA	CT-AT-----
10	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
101A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
102B	GATGGAACCC	CCTATATCCC	TTTATATTTA	CTTTTCTACT	TTTT-ATTTA	CT-AT-----
103A	GATGGAACCC	CCTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
104	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
108C	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
111A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
12	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
120C	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
121A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
123E	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
123G	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
126E	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
128C	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
129	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTGTTATTTA	TT-AT-----
131	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
132	GATGTAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
134A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	CT-AT-----
14	GATGGAACCC	ACTATATCCA	TTTATATATA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
15	GATGGAACCC	ACTATATCCA	TT-ATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
25	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
26	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTGFACTACT	TTTTTATTTA	TTTATTTACT
27C	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
28A	GATGGAACCC	CCTATATCCC	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
3	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
30A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
36C	GATGTAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	CT-AT-----
41B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
45C	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
5	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
50D	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
57B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
58B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
62D	GATGGAACCC	ACTATATCCA	TTTATATT-A	CCTTACTACT	TTTT-ATTTA	ACTAT-----
65D	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
67B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	TTTATTTACT
69A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	TTTATTTACT
69B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	TTTATTTACT
75A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	CT-AT-----
76A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	CT-AT-----
76E	GATGGAACCC	ACTATATGCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	CT-AT-----
77B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	TTTATTTACT
8	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
80A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
81A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	CT-AT-----
82A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	TTTATTTACT
84F	GATGGA-CCC	ACTATATCCA	TTTATATTTA	CCT-ACTACT	TTTT-ATTTA	CT-AT-----
86B	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
93C	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
98A	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
C.pri	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTTTATTTA	TTTATTTACT
S.3	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----
S.7	GATGGAACCC	ACTATATCCA	TTTATATTTA	CTTTACTACT	TTTT-ATTTA	CT-AT-----

Bases 1121-1180

1	GGGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
10	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	TGAGAGGTGA	AATGCCTAGA
101A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAAA
102B	GTGT-TTTT-	GATACCTAGT	ATCCATA-CC	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
103A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
104	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
108C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAAGTGA	AATGCCTAGA
111A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
12	GTGT-TTTT-	GATACCTAGT	ATCCATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
120C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	-TTTTTGATT	--AGAGGTGA	AATGCCTAGA
121A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGGGA	AATGCCTAGA
123E	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
123G	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	A-TGCCTAGA
126E	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
128C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
129	GTAT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTT-ATT	T-AGAGGTGA	AATGCCTAGA
131	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
132	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGTATT	T-AGAGGTGA	AATACCTATC
134A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
14	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
15	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
25	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
26	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
27C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CC	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
28A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CC	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
3	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
30A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
36C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGTATT	T-AGAGGTGA	AATGCCTAGA
41B	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
45C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
5	GTGT-TCCT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	TGAGAGGTGA	AATGCCTAGA
50D	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
57B	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
58B	GTGT-TTTT-	GATACCTAGT	ATCAATT-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
62D	GTGT-TTTT-	GATATCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
65D	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
67B	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
69A	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
69B	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	TTTTTTGATT	T-AGAGGTGA	AATGC-TAGA
75A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
76A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAAGTGA	AATGCCTAGA
76E	GTGT-TTTT-	GATACCTAGT	ATCAATAGCA	ATTTTTGATT	T-AGAGGAGA	AATGCCTAGA
77B	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
8	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGTATT	T-AGAGGTGG	AATGCCTAGA
80A	GTGG-TTTT-	GATAC-TAG-	ATCA-TA-C-	ATTTTTGATT	--AAAGGTGA	A-TGCCTAGA
81A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
82A	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
84F	GTGT-TTTT-	GATACCTAGT	ATCAATA-CC	ATTTTTGATT	T-AGAG-TGA	AATGCCTAGA
86B	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
93C	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
98A	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
C.pri	ATGTGTTTTT	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
S.3	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA
S.7	GTGT-TTTT-	GATACCTAGT	ATCAATA-CA	ATTTTTGATT	T-AGAGGTGA	AATGCCTAGA

Bases 1181-1240

1	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
10	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
101A	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
102B	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
103A	AAAAAAAAA-	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
104	AAAAAAAAA-	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
108C	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
111A	AAAAAAAAAA	AAA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
12	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
120C	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
121A	AAAAAAAAAA	AAAA----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
123E	AAAAAAAAAA	AAA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
123G	AAAAAAAAAA	AAAA--GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
126E	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
128C	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
129	AAAAAAAAAA	AAAA---CTA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
131	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
132	AAAAAAAAAA	AAAAAAAAGA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
134A	AAAAAAAAAA	AAAA----GA	AAAAATCGTG	GTTGGTAAGG	TTATAGTAGC	AAAAGCCATT
14	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
15	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
25	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
26	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
27C	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
28A	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
3	AAAAAAAAAA	AAA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
30A	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
36C	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
41B	AAAAAAAAAA	AA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
45C	AAAAAAAAAA	AAA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
5	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
50D	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
57B	AAAAAAAAAA-	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
58B	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
62D	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
65D	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
67B	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
69A	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
69B	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
75A	AAAAAAAAAA	AAA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
76A	AAAAAAAAAA	AAAAA--GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
76E	AAAAAAAAAA	AA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
77B	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
8	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
80A	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
81A	AAAAAAAAAA	A-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
82A	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
84F	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
86B	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
93C	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
98A	AAAAAAAAAA	AAA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
C.pri	AAAAAAA--	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
S.3	AAAAAAAAAA	AA-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT
S.7	AAAAAAAAAA	-----GA	AAAAATCGTG	GTTGGGAAGG	TTATAGTAGC	AAAAGCCATT

Bases 1301-1337 with the 14 presence/absence indel characters

1	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
10	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
101A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
102B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
103A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
104	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
108C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	1111101011	0010
111A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	1111101011	0010
12	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
120C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
121A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
123E	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101111	1010
123G	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101111	1010
126E	AAGGGAATAA	CTGAAAGAAA	GGAAATCA-T	TAGTTATT	0111101011	1010
128C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
129	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0110101001	1100
131	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
132	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0100101010	1010
134A	AACGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111111111	1010
14	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
15	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
25	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
26	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1011
27C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
28A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
3	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
30A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
36C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0100001010	1010
41B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
45C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
5	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
50D	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
57B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0011101011	1010
58B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
62D	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
65D	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
67B	AAGGGAATAA	CTGAAAGAAA	GGAA-TCAAT	TAGTTATT	0111101011	1011
69A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1011
69B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1111
75A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111111111	1010
76A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111111111	1010
76E	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111111111	1010
77B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1011
8	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
80A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
81A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0011111111	1110
82A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1011
84F	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
86B	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
93C	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
98A	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
C.pri	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111100011	1111
S.3	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010
S.7	AAGGGAATAA	CTGAAAGAAA	GGAAATCAAT	TAGTTATT	0111101011	1010

References

- Abrams, L. R. 1910. Studies on the Flora of Southern California III. Bulletin of the Torrey Botanical Club 37:149-153.
- Appels, R. and R. L. Honeycutt. 1986. rDNA: evolution over a billion years. *In* DNA Systematics, Vol. II, Plants, ed. S. K. Dutta, pp. 81-135. CRC Press, Boca Raton, Florida.
- Arnheim N., M. Krystal, R. Schmickel, G. Wilson, O. Ryder, and E. Zimmer. 1980. Molecular evidence for genetic exchanges among ribosomal genes on nonhomologous chromosomes in man and apes. Proceedings of the National Academy of Sciences U.S.A. 77:7323-7327.
- Arnheim, N. 1983. Concerted evolution of multigene families. *In* Evolution of Genes and Gene Proteins, eds. M. Nei and R. K. Koehn, pp. 38-61. Sinauer Associates, Sunderland, Massachusetts.
- Baker, D.D. and C.R. Schwintzer. 1990. Introduction *In* The Biology of Frankia and Actinorhizal Plants, eds. C.R. Schwintzer and J.D. Tjepkema, pp. 1-31. Academic Press, New York, New York.
- Baldwin, B. and S. Markos. 1998. Phylogenetic utility of the external transcribed spacer (ETS) of 18S-26S rDNA: Congruence of ETS and ITS trees of Calycadenia (Compositae). Molecular Phylogenetics and Evolution 10:449-463.
- Baldwin, B.G., M.J. Sanderson, J.M. Porter, M.F. Wojciechowski, C.S. Campbell, and M.J. Donoghue. 1995. The ITS region of nuclear ribosomal DNA: a valuable source of evidence on angiosperm phylogeny. Annals of the Missouri Botanical Garden 82:247-277.
- Bena, G., Jubier, M.-F., Olivieri, I., and B. LeJueune. 1998. Ribosomal external and internal transcribed spacers: Combined use in phylogenetic analysis of *Medicago* (Leguminosae). Journal of Molecular Evolution 46:299-306.
- Blauer, A.C., A.P. Plummer, E.D. McArthur, R. Stevens, and B.C. Giunta. 1975. Characteristics and hybridization of important Intermountain shrubs. I. Rose family. USDA Forest Service Research Paper INT-169. 36 pgs.

- Brayton, R. and H.A. Mooney. 1966. Population variability of *Cercocarpus* in the White Mountains of California as related to habitat. *Evolution* 20:383-391.
- Buckler, E. S. IV, A. Ippolito, and T. P. Holtsford. 1997. The evolution of Ribosomal DNA: Divergent Paralogues and phylogenetic implication. *Genetics* 145:821-832.
- Buckler, E. S., IV, and T. P. Holtsford. 1996. *Zea* Systematics: ribosomal ITS evidence. *Molecular Biology and Evolution* 13:612-622.
- Bull, J. J., J. P. Huelsenbeck, C. W. Cunningham, D. L. Swofford, and P. J. Waddell. 1993. Partitioning and combining data in phylogenetic analysis. *Systematic Biology* 42:384-397.
- Correll, D. S., and M. C. Johnston. 1970. Manual of the Vascular Plants of Texas. Texas Research Foundation, Renner, Texas.
- Campbell, C. S., M. F. Wojciechowski, B. G. Baldwin, L. A. Alice, and M. J. Donoghue. 1997. Persistent nuclear ribosomal DNA sequence polymorphism in *Amelanchier* agamic complex (Rosaceae). *Molecular Biology and Evolution* 14:81-90.
- Davis, J.N. and J.D. Brotherson. 1991. Ecological Relationships of Curlleaf Mountain Mahogany (*Cercocarpus ledifolius* Nutt.) Communities in Utah and implications for Management. *Great Basin Naturalist* 51:153-166.
- Dover, G. A. 1982. Molecular drive: a cohesive mode of species evolution. *Nature* 299:111-117.
- Doyle, J. J., and S. Doyle. 1987. A rapid DNA isolation procedure for small quantities of fresh leaf tissue. *Phytochemistry Bulletin* 19:11-15.
- Dvorak, J. 1990. Evolution of multigene families: the ribosomal RNA loci of wheat and related species. *In Plant Populational Genetics, Breeding, and Genetic Resources*, eds. A. H. D. Brown and M. T. Clegg, pp. 83-97. Sinauer Associates, Sunderland, Massachusetts.
- Eastwood, A. 1898. *Proceedings of the California Academy of Sciences* 3:136.
- Evans, R. C., L. A. Alice, C. S. Campbell, E. A. Kellogg, and T. A. Dickson. 2000. The granule-bound starch synthase (GBSSI) gene in the Rosaceae:

- Multiple loci and phylogenetic utility. *Molecular Phylogenetics and Evolution* 17:388-400.
- Faith, D. P. 1991. Cladistic permutation test for monophyly and nonmonophyly. *Systematic Zoology* 40:366-375.
- Farris, J. S., M. Källersjö, A. G., Kluge, and C. Bult. 1994. Testing significance of incongruence. *Cladistics* 10:315-319.
- Farris, J. S., M. Källersjö, A. G., Kluge, and C. Bult. 1995. Constructing a significant test for incongruence. *Systematic Biology* 44:570-572.
- Felsenstein, J. 1985. Confidence limits on phylogenies: an approach using the bootstrap. *Evolution* 39: 783-791.
- Gray, A. 1852. *Plantae wrightianae* part 1, pp. 54. Smithsonian Institution, G. P. Putnam, New York, New York.
- Hamby, K. R. and E. A. Zimmer. 1992. Ribosomal RNA as a phylogenetic tool in plant systematics. *In* *Molecular Systematics of Plants*, eds. P. Soltis, D. Soltis, and J. Doyle, pp. 50-91. Chapman and Hall, New York, New York.
- Hamilton, M.B. 1999. Four primer pairs for the amplification of chloroplast intergenic regions with interspecific variation. *Molecular Ecology* 8:521-523.
- Hedren, M., M. F. Fay, and M. W. Chase. 2001. Amplified fragment length polymorphisms (AFLP) reveal details of polyploid evolution in *Dactylorhiza* (Orchidaceae). *American Journal of Botany* 88:1868-1880.
- Henrickson, J. 1987. Two new species of *Cercocarpus* (Rosaceae) from Mexico. *Systematic Botany* 12:293-298.
- Hillis, D. M. 1987. Molecular versus morphological approaches to systematics. *Annual review of Ecology and Systematics* 18:23-42.
- Hillis, D. M. and M. T. Dixon. 1991. Ribosomal DNA: Molecular evolution and phylogenetic inference. *Quarterly Review Biology* 66:411-452.
- Hitchcock, C. L., and A. Cronquist. 1961. *Cercocarpus* *In* *Vascular Plants of the Pacific Northwest, Part 3: Saxifragaceae to Ericaceae*, pp. University of Washington Press, Seattle, Washington.

- Holmgren, N. H. 1987. *Cercocarpus ledifolius* var. *intermontanus* (Rosaceae), a new varietal name for the intermountain curl-leaf mountain-mahogany. *Brittonia* 39:423-427.
- Hooker, W. J. 1840. *Icones plantarum series 1*, vol. 4, pp. 332. Royal Botanic Society, Kew Gardens, London.
- Hooker, W. J., and G. A. W. Arnott. 1839. *The Botany of Captain Beechey's Voyage*, part 8, pp. 337. H. G. Bohn, London.
- Humboldt, F. W. H. A., A. Bonpland, and C. S. Kunth. 1832. *Cercocarpus In Nova Genera et Species Plantarum*, vol. 6, pp. 232. reprinted by Weldon, Wesley and Hafner, New York, New York.
- Jansen, R. K. and J. D. Palmer. 1987. Chloroplast DNA from lettuce and *Barnadesia* (Asteraceae): structure, gene location, and characterization of a large inversion. *Current Genetics* 11:553-564.
- Jones, M. E. 1891a. New plants from Arizona, Utah, and Nevada. *Zoe* 2:12-17.
- Jones, M. E. 1891b. New species and notes of Utah Plants. *Zoe* 2:236-252.
- Jourgensen, R. A., R. E. Cuellar, W. F. Thompson, and T. A. Kavanagh. 1987. Structure and Variation in Ribosomal RNA genes of pea: characterization of a cloned ribosomal DNA repeat and chromosomal ribosomal DNA variants. *Plant Molecular Biology* 8:3-12.
- Kearney, T. and R. Peebles. 1951. *Cercocarpus In Arizona Flora*, pp. 388. University of Arizona Press, Tucson, Arizona.
- Kellogg, E. A., R. Appels, and R. J. Mason-Gamer. 1996. When genes tell different stories: the diploid genera of Triticeae (Gramineae). *Systematic Botany* 21:321-347.
- Klemmedson, J.O. 1979. Ecological importance of actinomycete-nodulated plants in the western U.S. *Botanical Gazette* 140(suppl.):S91-S96.
- Koopman, W. J. M., M. J. Zevenbergen, and R. G. Van den Berg. 2001. Species relationships in *Lactuca* s.l. (Lactuceae, Asteraceae) inferred from AFLP fingerprints. *American Journal of Botany* 88:1881-1887.
- Linder, C. R., L. R. Goertzen, B. Vanden Heuvel, J. Francisco-Ortega, and R. K. Jansen. 2000. The complete External Transcribed Spacer of 18S - 26S

rDNA: Amplification and Phylogenetic utility at low taxonomic levels in the Asteraceae and closely allied families. *Molecular Phylogenetics and Evolution* 19:167-176.

- Lis, R. A. 1992. Leaf Architectural Survey of *Cercocarpus* (Rosaceae) and its systematic significance. *International Journal of Plant Science* 153:258-272.
- Lis, R. A. 1993. *Cercocarpus*. In *The Jepson Manual: Higher Plants of California*, ed. J. C. Hickman, pp. 948-949. University of California Press, Berkeley, California.
- Little, E. L. 1953. Five varietal transfers of United States trees. *Phytologia* 4:305-310.
- MacArthur, E. D., and S. C. Anderson. 1985. A Cytotaxonomic Contribution to the western North American Rosaceous flora. *Madrono* 32:24-28.
- Martin, F. L. 1950. A revision of *Cercocarpus*. *Brittonia* 7:91-111.
- Mayol, M., and J. A. Rossello. 2001. Why Nuclear Ribosomal DNA spacers (ITS) tell different stories in *Quercus*. *Molecular Phylogenetics and Evolution* 19:167-176.
- McVaugh, R. 1952. Remarks of the genus *Cercocarpus* in Texas. *Field and Laboratory* 10:35-40.
- Mindell, D. P. and R. L. Honeycutt. 1990. Ribosomal RNA in vertebrates: evolution and phylogenetic applications. *Annual Review of Ecology and Systematics* 21:541-566.
- Morgan, D. R., D. E. Soltis, and K. Robertson. 1994. Systematic and Evolutionary Implications of *rbcL* sequence variation in Rosaceae. *American Journal of Botany* 81:890-903.
- Morley, T. 1949. In Documented chromosome number of plants. *Madrono* 10:95.
- Mortenson, T.H. 1973. Ecological variation in the leaf anatomy of selected species of *Cercocarpus*. *Aliso* 8:19-48.
- Mueller, U. G., and L. L. Wolfenbarger. 1999. AFLP genotyping and fingerprinting. *Trends in Ecology and Evolution* 14:389-394.

- Munz, P. A. 1959. A California Flora. University of California Press, Berkeley, California.
- Murphy, W. J., E. Eizirik, W. E. Johnson, Y. P. Zhang, O. A. Ryder, and S. J. O'Brien. 2001. Molecular phylogenetics and the origins of placental mammals. *Nature* 409:614-618.
- Nickrent, D. L. and D. E. Soltis. 1995. A comparison of angiosperm phylogenies from nuclear 18S rDNA and *rbcL* sequences. *Annals of the Missouri Botanical Garden* 82:208-234.
- O'Kane, S. L., Schaal, B. A., and I. A. Al-Shehbaz. 1996. The origins of *Arabadopsis suecica* (Brassica).
- Palmer, J. D. 1986. Isolation and structural analysis of chloroplast DNA. *Methods in Enzymology* 118:167-186.
- Rafinesque, C. S. 1832. Twenty new genera of plants from the Oregon Mountains. *Atlantic Journal* 1:146.
- Rieseberg, L. H., J. Whitton, and C. R. Linder. 1996. Molecular marker incongruence in plant hybrid zones and phylogenetic trees. *Acta Botanica Neerlandica* 45:243-262.
- Rieseberg, L. H., and O. Gerber. 1994. Hybridization in the Catalina Island Mountain Mahogany (*Cercocarpus traskiae*):RAPD evidence. *Conservation Biology* 9:199-203.
- Rieseberg, L. H., S. Beckstromsternberg, and K. Doank. 1990. *Helianthus annuus* ssp *texanus* has chloroplast DNA and nuclear ribosomal RNA genes of *Helianthus debilis* ssp *cucumerifolius*. *Proceedings of the National Academy of Science USA* 87:593-597.
- Rieseberg, L. H., S. Zona, L. Aberbom, and T. D. Martin. 1989. Hybridization in the island endemic, Catalina Mahogany. *Conservation Biology* 3:52-58.
- Rosenberg, M. S., and S. D. Kumar. 2001. Incomplete taxon sampling is not a problem for phylogenetic inference. *Proceedings of the National Academy of Sciences USA* 98: 10751-10756.
- Russell S. K., and E. W. Schupp. 1998. Effects of microhabitat patchiness on patterns of seed dispersal and seed predation of *Cercocarpus ledifolius* (Rosaceae). *Oikos* 81:434-443.

- Russell, S. K. 1997. Thesis. Utah State University.
- Rydberg, P. A. 1913. *Cercocarpus*. North American Flora 22:418-424.
- Sanderson, M. J. and J. J. Doyle. 1992. Reconstruction of organismal gene phylogenies from data on multigene families: concerted evolution, homoplasy, and confidence. *Systematic Biology* 41:4-17.
- Sang, T., Crawford, D. J., and Stuessy, T. F. 1995. Documentation of reticulate evolution in peonies (*Paeonia*) using sequences of internal transcribed spacer of nuclear ribosomal DNA: Implications for biogeography and concerted evolution. *Proceedings of the National Academy of Sciences USA*. 92:6813-6817.
- Sang, T., D. J. Crawford, T. F. Stuessy, and O. Silva. 1995. ITS sequences and the phylogeny of the genus *Robinsonia* (Asteraceae). *Systematic Botany* 20:55-64.
- Saunders, M. A., and S. V. Edwards. 2001. Dynamics and phylogenetic implications of mtDNA control region sequences in new world jays (Aves:Corvidae). *Journal of Molecular Evolution* 51:97-109.
- Searcy, K. B. 1969. Variation in *Cercocarpus* in southern California. *New Phytologist* 68:829-839.
- Schneider, C. K. 1905a. *Illustriertes Handbuch der Laubholzkunde*, vol. 1, pp. 530. Verlag von Gustav, Fisher, Jena.
- Schneider, C. K. 1905b. Beitrag zur Kenntnis der Arten und Formen der Gattung *Cercocarpus* Kunth. *Mitteilungen der Deutschen Dendrologischen Gesellschaft* 14:125-129.
- Soltis, D.E. and P.S. Soltis. 1998. Introduction *In* *Molecular Systematics of Plants II: DNA sequencing*. Soltis, P.S., D.E. Soltis, and J.J. Doyle (eds). Kluwer Academic publishers, Boston, Massachusetts.
- Soltis, P. S., and D. E. Soltis. 1991. Multiple origins of the allotetraploid *Tragapogon mirus* (Compositae)-rDNA evidence. *Systematic Botany* 16:407-413.
- Soltis, P. S., D. E. Soltis, P. G. Wolf, D. L. Nickrent, S. -M. Chaw, and R. L. Chapman. 1999. The phylogeny of land plants inferred from 18S rDNA sequences: Pushing the limits of rDNA signal? *Molecular Biology and Evolution* 16:1774-1784.

- Steane D. A., G. E. McKinnon, R. E. Vaillancourt, and B. M. Potts. 1999. ITS sequence data resolve higher level relationships among the eucalypts. *Molecular Phylogenetics and Evolution* 12:215-223.
- Stutz, H.C. 1974. Rapid Evolution in Western Shrubs. *Utah Science* 35:16-20.
- Suh, Y., L. B. Thien, H. E. Reeve, and E. A. Zimmer. 1993. Molecular evolution and phylogenetic implications of internal transcribed spacer sequences of ribosomal DNA in Winteraceae. *American Journal of Botany* 80:1042-1055.
- Swofford, D. L. 1999. PAUP*: Phylogenetic Analysis Using Parsimony beta test version 4.08b. Sinauer, Sunderland, Massachusetts
- Templeton, A. R. 1983. Convergent evolution and non-parametric inference from restricted fragment and DNA sequence data *In* Statistical Analysis of DNA sequence Data, ed. B. Wier, pps. 151-179. Marcel Dekker, New York, New York.
- Thompson, W. F. and R. B. Flavell. 1988. DnaseI sensitivity of ribosomal RNA genes in chromatin and nucleolar dominance in wheat. *Journal of Molecular Biology* 204:535-548.
- Thorne, R. F. 1978. New Subspecific Combinations for Southern California Plants. *Aliso* 9:189-196.
- Torrey, J., and A. Grey. 1840. A Flora of North America. Vol. 1, part 3. Wiley and Putnam, New York, New York.
- Vargas P., B. G. Baldwin, and L. Constance. 1998. Nuclear ribosomal DNA evidence for a western North American origin of Hawaiian and South American species of *Sanicula* (Apiaceae). *Proceedings of the National Academy of Sciences USA* 95:235-240.
- Vos, P., R. Hogers, M. Bleeker, M. Reijans, T. Van De Lee, M. Hornes, A. Freijters, J. Pot, J. Peleman, M. Kuiper, and M. Zabeau. 1995. AFLP, a new technique for DNA fingerprinting. *Nucleic Acid Research* 23:4407-4414.
- Waters, E. R. and B. A. Schaal. 1996. Biased gene conversion is not occurring among rDNA repeats in the *Brassica* triangle. *Genome* 39:150-154.

- Watson, S. 1882. Proceedings of the American academy of Arts and Sciences 17:353.
- Watson, S. 1876. Botany of California, vol. 1, pp. 175. Welch, Bigelow and Co., Cambridge, Massachusetts.
- Watson, S. 1875. Revision of the Genus *Ceanothus*, and Descriptions of new plants. Proceedings of the American Academy of Arts and Sciences 10:333-350.
- Wendel, J. F. and J. J. Doyle. 1998. Phylogenetic incongruence: window into genome history and molecular evolution. In Molecular Systematics of Plants II: DNA sequencing, eds. D. E. Soltis, P. S. Soltis and J. J. Doyle, pp 265-296. Kluwer Academic Publishing, Boston, Massachusetts.
- Wendel, J. F., A. Schnabel, and T. Seelanan. 1995. Bidirectional interlocus concerted evolution following allopolyploid speciation in cotton (*Gossypium*). Proceedings of the National Academy of Sciences USA 92:280-284.
- Wendel, J. F., A. Schnabel, and T. Seelanan. 1995. An unusual ribosomal DNA sequence from *Gossypium gossypioides* reveals ancient, cryptic intergenomic introgression. Molecular Phylogenetics and Evolution 4:298:313
- Zhang, L. B., H. P. Comes, and J. W. Kadereit. 2001. Phylogeny and Quaternary history of the European Montane/Alpine endemic *Soldanella* (Primulaceae) based on ITS and AFLP variation. American Journal of Botany 88:2331-2345.
- Zimmer, E. A., S. L. Martin, S. M. Beverly, Y. M. Kan, and A. C. Wilson. 1980. Rapid duplication and loss of genes coding for the chains of hemoglobin. Proceedings of the National Academy of Sciences U.S.A. 77:2158-2162.

Vita

Brian David Vanden Heuvel was born in Greeley, Colorado on September 4, 1975. Joseph and Raedene Purma adopted him soon after his birth. After the death of Joseph Purma, his mother remarried Steve Vanden Heuvel, and both mother and son changed their last name to Vanden Heuvel. He grew up in Castle Rock, Colorado, and graduated from Douglas County High School in 1993. He attended Fort Lewis College in Durango, Colorado from 1993 to 1994. In 1994, he transferred to Colorado State University to pursue a Bachelor of Science degree, majoring in Botany. He graduated Phi Beta Kappa and cum laude in 1997. In the fall of 1997, he began his graduate career at The University of Texas at Austin.

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