North-Atlantic records of *Schizymenia dubyi* extend along the eastern shores of the North Atlantic from Morocco to southern Britain and Ireland, and the species is also recorded from Iceland. A study was undertaken to confirm the identity of the specimens from Iceland that were geographically separate from the main distribution of *S. dubyi* and in contrast to other species of the genus did not have gland cells. We analyzed *rbcL* and COI molecular sequence data from Icelandic specimens and compared the results with those for *Schizymenia* specimens available in GenBank. For both markers, *Schizymenia* was shown to be a monophyletic genus. The Icelandic specimens were clearly genetically distinct from *S. dubyi* and formed a well-supported clade with *Schizymenia* species from the Northern Pacific. Based on these results, we have described a new species, *Schizymenia jonssonii*, which can be distinguished by molecular phylogeny, its lack of gland cells and by being strictly intertidal. Crustose tetrasporophytes with identical COI and *rbcL* sequences were found at the same locations as foliose plants. *Schizymenia apoda* is reported for the first time in the UK, its identity confirmed by *rbcL* sequence data. In light of these findings, it is likely that by further molecular analysis of the genus *Schizymenia* in the north-eastern Atlantic and the Mediterranean, a higher diversity of *Schizymenia* spp. will be discovered in this region.

Key index words: COI; Iceland; molecular phylogeny; phenology; North Atlantic; *rbcL*; *Schizymenia jonssonii*

Abbreviations: BP, before present; COI, Cytochrome c oxidase I gene; dNTP, deoxyribonucleotide triphosphate nucleobases
Iceland might be genetically separated from the relations within the genus Schizymenia. In this study, we used COI of 30 phases (Saunders et al. 2015). All the species are lar evidence to have two dissimilar life history Saunders et al. 2015). Two species, Schizymenia dubyi and S. apoda are also known from the Atlantic and the Indian Oceans (Silva et al. 1996, Gabriel et al. 2011). are also known from the Atlantic and the Indian Oceans (Silva et al. 1996, Gabriel et al. 2011). In Iceland, Schizymenia was first registered by Caram and Jónsson (1972) in their Icelandic checklist as Schizymenia dubyi. Previously, H. Jónsson (1901) had recorded Dilsea edulis (=D. carnosa) from Iceland. However, this species has not been recorded in Iceland since and examination of H. Jónsson’s specimens, kept in the Botanical Museum in Copenhagen (C), showed them to be a Schizymenia sp. On a visit in August 2007 to the collecting site of H. Jónsson in the intertidal zone at Óndverðarnes, Schizymenia sp. was found to be common (K. Gunnarsson, pers. obs.). The Icelandic specimens of Schizymenia sp. are structurally and anatomically similar to other Schizymenia species in all aspects except for the total lack of gland cells and that they are only found in the intertidal zone. “Haematocelis”-like crusts have been found among the foliose Schizymenia at many sites in Iceland. The present-day distribution of Schizymenia in Iceland is from the southwest coast, where it is relatively common, becoming sparser along the West coast and into the middle part of the colder North coast. It is not found in the coldest, eastern part, of the coast. A similar pattern is encountered by species that have their main distribution area south of Iceland and their northern limit of distribution along the Icelandic coast (as e.g., Chondrus crispus, Corallina officinalis, and Pelvetia canaliculata). These species are common in the Faeroes, southern Scandinavia, and Scotland, whereas Schizymenia, except for unconfirmed records (due to a lack of specimens) from northern Scotland, is not found in these areas (Rueness 1977, Nielsen et al. 1995, Nielsen and Gunnarsson 2001, Hardy and Guiry 2003). The collecting sites for specimens of Schizymenia closest to Iceland in the North Atlantic are in southern England and southern Ireland. The disjunct distribution of Schizymenia in the northern North Atlantic, the lack of gland cells in the specimens collected in Iceland, and the observation that they have only been found in the intertidal zone evoked suspicion that the Schizymenia found in Iceland might be genetically separated from the more southern relatives. In this study, we used COI and rbcL molecular markers to analyze phylogenetic relations within the genus Schizymenia with the aim of resolving the identity and affinity of the species found in Iceland.

MATERIALS AND METHODS

Sampling. Schizymenia plants used in the molecular analysis were collected at the localities listed in Table S1 in the Supporting Information. A piece of each specimen, c. 1–2 cm², was cut off and placed in silica gel for drying. Voucher specimens were dried on herbarium sheets and deposited in the herbarium of the Icelandic Institute of Natural History (ICEL). Herbarium abbreviations follow Thiers (2018).

A sampling location at Stekkjarvikur, Reykjanes peninsula, southwestern Iceland (64°1.770’ N, 22°14.616’ W), was visited every month at low water, spring tide from April 2016 to May 2017 to monitor seasonal changes in morphology and reproduction in both the foliose and crustose plants of Schizymenia. Plants were collected to examine the presence of reproductive structures, and the lengths of the ten largest foliose specimens were measured. The substratum at the study site is of uneven, basaltic lava rocks. The mean tidal range at spring tide is c. 3.5 m and the sea surface temperatures vary on average from 1.5°C for March to 13.5°C in August (Icelandic Coast Guard 2019, Marine and Freshwater Research Institute 2019).

In the laboratory, the plants were photographed and hand-sectioned for microscopic observations using a Leitz RMZ light microscope equipped with a Leitz DFC320 digital microphotographic camera. Voucher specimens were prepared and deposited at the Natural History Institute in Reykjavik (ICEL).

DNA extraction and sequencing. Pieces c. 5–10 mm² of silica-dried Schizymenia blades and crusts were ground in a mortar with chemically pure sand (Merck, Darmstadt, Germany) to a fine powder. The powder was rehydrated in CTAB (cetyltrimethylammonium bromide), Sarkosyl, and proteinase K (500:50:10) and then DNA was dissolved with SEVAC (chloroform/isoamyl alcohol, 24:1) at 65°C and precipitated with isopropanol and ethanol. The DNA was then cleaned using GFX DNA cleaning kit (GE Healthcare, UK) following the manufacturer’s protocol. To each 1 μL DNA sample, we added 22 μL of PCR mixture contained, 2.5 μL buffer, 1.5 μL MgCl₂, 0.5 μL dNTP, 1 μL of each forward and reverse primers, and 0.5 μL Taq (Biotaq DNA Polymerase kit; Bioline Ltd, UK) for amplification. The plastid-encoded rbcL was amplified using F57 and R1442 primers (Freshwater and Rueness 1994). The mitochondrial COI gene was amplified using GAZFI and GAZRI primers (Saunders 2005) and M13IF and M13Rx (Saunders and Moore 2013). PCR amplifications were undertaken in a Techne thermal cycler (Cole-Parmer Ltd., St. Neots, UK). For rbcL amplification, we used an initial denaturation step at 95°C for 2 min, followed by 35 cycles of denaturation at 95°C for 1 min, annealing at 47°C for 1 min and extension at 72°C for 2 min, and then a final extension at 72°C for 2 min. For COI, we used an initial denaturation step at 94°C for 2 min, followed by five cycles of denaturation at 95°C for 30 s, annealing at 45°C for 30 s and 1 min extension at 72°C and then 35 cycles starting with denaturation at 94°C for 30 s and annealing at 46.5°C for 30 s, and an extension at 72°C finishing by a final extension at 72°C for 1 min (Saunders and Moore 2013). The amplification success was tested by agarose gel electrophoresis. The PCR amplifications were undertaken with 18 samples of foliose Schizymenia (17 from Iceland and 1 from UK) and 3 samples of “Haematocelis”-like crusts. The PCR products were sent for DNA sequencing at the sequencing laboratory of the Natural History Museum in London on an ABI 3730 XL capillary DNA analyser (Applied Biosystems).

Phylogenetic analysis. The resulting sequences were edited with BioEdit, version 7.2.6 (Hall 1999). Sequences from
GenBank of *rbcL* and COI for *Schizymenia* spp. were compared with our sequences and *Platoma cyclocolpum* and *Titanophora weberae* used as outgroup (Saunders et al. 2015). Information on GenBank accession numbers, origin, date of collection, etc., of the sequences used to construct the phylogenetic trees is shown in Table S1.

Phylogenetic analyses were performed with maximum likelihood and Bayesian methods. Maximum likelihood (ML) analyses were done using MEGA5 (Tamura et al. 2011). Support for ML analysis was assessed both with approximate likelihood ratio test and bootstrap values were created with 1,000 resampling replicates for both likelihood and bootstrap methods. Maximum likelihood and Bayesian methods. Maximum likelihood (ML) trees is shown in Table S1.

The first 25% of the samples from the cold chain were discarded as burn-in. Maximum likelihood and bootstrap values were created using MEGA5 (Tamura et al. 2011).

**RESULTS**

**Taxonomy.** *Schizymenia jonssonii* K.Gunnarsson & J.Brodie sp. nov. (Fig. 1)

**Diagnosis:** Gametophytic phase foliose with surface either flattened or with irregular ridges and depressions. Thallus lacking gland cells. Color dark red to brownish red. Tetrasporophytic phase crustose, with uneven surface having numerous small swellings. Can equally be distinguished by its nucleotide sequences of the mitochondrial COI and plastid *rbcL* genes.

**Description:** Species consisting of an upright foliose phase and a “*Haematocelis*” crustose phase. Upright fronds have a cylindrical stipe, 1-3 mm long, and are attached with a discoid or slightly conical holdfast. Thallus oblong, sometimes lobed or split, 50–350 mm in length, semi-gelatinous; surface either flattened or with irregular ridges and depressions. Color dark red to brownish red. Fronds are 250–600 μm thick (older blades in winter up to 900 μm).

In surface view, cells are rounded, 7–9 μm in diameter. The blade consists of an inner medulla and an outer cortex (Fig. 2a). In transverse section (TS), the medulla consists of loosely interwoven, branched filaments of elongated cells, 3–7 μm in diameter, and up to more than 150 μm long. The cortex consists of cell rows, typically 6–8 cells (–20) in length in TS with dichotomous branching. The innermost cortical cells are spherical, 30 μm in diameter in TS. Cells diminish in size toward the surface and the outermost cortical cells are elongate, c. 7.5 × 15 μm in TS. The medullary filaments are connected by pit connections to the cortical branches but no secondary pit connections were observed. Gland cells are absent.

Cystocarps are visible as small dark-red spots on the surface of the fertile blades and are most common in the distal end but are also occasionally found in the middle and the lower part of the thallus. Cystocarps are situated in the inner cortex or between the medullary filaments, close to the cortex, 130–300 μm across and are without pericarp. Ostioles are visible as depressions in the surface of the thallus by the cystocarps (Fig. 2b). Individual carposporangia are 15–40 μm in diameter. Carpogonia and spermatangia were not observed.

Crusts are dark red in color, irregularly shaped, 1–7 cm across, and cartilaginous. Crusts are up to 2.5 mm thick and layered with darker and lighter bands seen in cross section. Horizontal basal filaments curve up and give rise to upright, narrower filaments. Tetrasporangia are zonate, produced in the surface layer, 50–70 μm × 15–20 μm (Fig. 2c). Released tetracosperos are spherical, c. 25 μm in diameter (Fig. 2d). Surface of crust is uneven with numerous small swellings (Fig. 2e). Rhizoids, two to five cells long, are occasionally found on the underside of the crust (Fig. 2f).

**Holotype:** Gametophyte collected in the lower intertidal zone at Stekkjarvikur, Reykjanesessagi, Iceland (64°1.77′ N, 22°14.62′ W), April 21, 2016 (Fig. 1). GenBank accession numbers: *rbcL* = MN567259, COI = MN567252. Specimen deposited in the Cryptogamic Herbarium of the Natural History Museum, London (BM). BM museum number: BM013844101.

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**Fig. 1.** Holotype specimen of *Schizymenia jonssonii* sp. nov. (BM013844101), collected at Stekkjarvikur, SW Iceland on April 21, 2016. Scale = 2 cm. [Color figure can be viewed at wileyonlinelibrary.com]
Isotype: Crustose tetrasporophyte, collected in the lower intertidal zone at Stekkjarvikur, Reykjanesskagi, Iceland (64°1.77' N, 22°14.62' W), April 21, 2016. GenBank accession numbers: \textit{rbc}L = MN567260, COI = MN567253, deposited in the Cryptogamic Herbarium of the Natural History Museum, London (BM). BM Museum number: BM013844102.

Etymology: The species is named in honor of the Icelandic phycologist Sigurður Jónsson, for his contribution to the knowledge of Icelandic macroalgae and phycology in general.

Paratypes: Gametophytes collected at Sölvaflá, Vestmannaeyjar, August 12, 1999 (Voucher =ICEL586), Stekkjarvikur, Reykjanesskagi, April 21, 2016 (Voucher =ICEL13226), Ondverðarnes, August 8, 2007 (Voucher =ICEL6225), Flatey, Breidjördur, June 17, 1977 (Voucher =ICEL856), Skálavík, July 3, 2008 (Voucher =ICEL7748) and Laugatangi, Hrísey, June 14, 2006 (Voucher =ICEL5343). All paratype specimens are deposited in the algal herbarium of the Icelandic Institute of Natural History (ICEL).

Type Locality: Stekkjarvikur, Reykjanesskagi, Iceland (64°1.77' N, 22°14.62' W).

Distribution: \textit{Schizymenia} is found in Vestmannaeyjar Archipelago off the south coast of Iceland and is distributed more or less continuously from there, along the southwest coast. It is then found sporadically along the west and the northwest coast and at one locality on the north coast (Fig. 3).

The upright fronds started to appear in February, most often attached directly to the rock by a small (2–4 mm diameter), conical holdfast, but sometimes they grew up from a \textit{Haematocelis}-like crust. The fronds increased in size until July when they were at their maximum size. From May until August, the fronds were mostly yellow in color (Fig. 4a) and were often seen with bleached spots. During the period from June to August, the blades had epiphytes and were grazed by numerous gastropods. In September, the blades started to erode and many of the plants died. Only a very few, small plants were observed from December to February. Cystocarps were first observed in September and last seen in March in some of the small blades that had overwintered from the previous year.

Crusts were cartilaginous, with a firm, uneven surface and often with dark red swellings, 1–5 cm in diameter (Fig. 4b). The crusts were loosely attached to the substratum and were easily detached. During the winter, the crusts were without overgrowth of ephemeral algae and could be spotted by eye due to their slightly lighter color than other red algal crusts, \textit{Hildenbrandia} spp. and \textit{Haemesharia hennedyi} that were growing in the vicinity. In the summer, the crusts were overgrown by the red and green algae \textit{Cystoclonium purpureum}, \textit{Ceramium virgatum}, and \textit{Ulva fenestrata} (cf Hughey et al. 2019) and difficult to spot. Crusts were up to 7 cm in diameter. Tetrasporangia were first seen in December in the
inner layers of some older crusts. In January, tetrasporangia were observed in the surface layer on many of the crusts but often the sporangia contained only two spores. The portion of sporangia with four spores increased from January to March (Fig. 2c). In April, only a few sporangia were observed, all with two spores. The older crusts were layered, and often had holes both between layers and between the crust and the substratum (Fig. 2e). These holes were inhabited by a variety of invertebrates. The crustose plants grew along with the blade-phase on moderately exposed to exposed shores and were found throughout the year.

Phylogenetic analysis. 17 rbcL sequences of 1210 bp and 17 COI sequences of 680 bp were obtained from specimens of Schizymenia jonssonii collected in Iceland. An additional rbcL sequence was obtained for a specimen collected in Plymouth, UK. All the sequences from the Icelandic samples turned out to be identical for the respective markers. The ML tree for rbcL contained 23 Schizymenia sequences of which 6 were new sequences from Icelandic specimens and one from Plymouth.

Analysis of COI and rbcL sequences of Schizymenia jonssonii, and Schizymenia sequences available from GenBank showed a monophyletic tree and two distinct clades of species (Figs. 5 and 6). The new species S. jonssonii formed a highly supported cluster with the Pacific species S. pacifica, S. tenuis and the undescribed species “Schizymenia sp._1Cal” (Saunders et al. 2015), and was most closely related to “Schizymenia sp._1Cal” with c. 97–98% similarity for the two markers studied. In the phylogenetic tree, this clade was distinctly separated from a clade containing the two other species of Schizymenia found in the Atlantic, S. dubyi, and S. apoda. The rbcL sequences from two specimens of unidentified Schizymenia spp. from Japan, clustered with S. dubyi and S. apoda, but formed separate branches. The rbcL sequence of a sample of Schizymenia sp. from Plymouth turned out to be conspecific with S. apoda from Namibia and the Azores (Fig. 5). The differences in rbcL and COI sequences between the two clades ranged from 4.6% to 4.8% and 5.1% and 6.8%, respectively (Table 1).

Discussion

The results from the present analysis of COI and rbcL data reveal a new species of Schizymenia in the Atlantic, S. jonssonii K.Gunnarsson & J.Brodie, which adds a third species of Schizymenia to the North Atlantic seaweed flora. This species differs from other Schizymenia species by being strictly intertidal and lacking gland cells. It is only distantly related to the other species of the genus found in the Atlantic (i.e., S. dubyi and S. apoda) but clusters more closely with the Pacific species S. tenuis, S. pacifica and the undescribed species “Schizymenia sp._1Cal” (Saunders et al. 2015).

Analysis of the rbcL marker from a specimen collected in Plymouth proved to be conspecific with Schizymenia apoda from the Azores and Namibia. This is the first record of S. apoda in the UK and the first genetic sequence analysis of the genus Schizymenia from the UK.

The uncertainty of the identity of the species in relation to the type specimen of type of the genus, Schizymenia dubyi, which is from Cherbourg, France, cannot be resolved until sequence data of the type specimen is available. We attempted without success to obtain a sequence from the type specimen of the genus dated 1826 (material kindly given to us by the University of Strasbourg Herbarium [STR]) following the method and recommendations given by
Hughey and Gabrielson (2012). Although there is a DNA sequence from a specimen identified as *S. dubyi* for the west of Brittany, the presence of species identified as *S. apoda* nearby makes it possible that either of the two species or even another species will prove to be identical to the type specimen. This is further confounded by possible introgression of *S. apoda* and *S. dubyi* (Saunders et al. 2015). The clear genetic separation of the two clades in the phylogenetic tree (Figs. 5 and 6) and the high sequence differences between the two clades (Table 1) makes an argument for considering a revision of the concept of the genus *Schizymenia*, including the creation of a new genus, but until more species of the genus have been sequenced and the phylogenetic position of the type has been established, this remains a future goal.

*Schizymenia dubyi* has been recorded from Morocco (Benhissoune et al. 2003), in the Mediterranean sea (Sciuto et al. 1979, Alongi and Cormaci 1993) and along the Atlantic coasts of Portugal (Ardré 1970), Northern Spain (Rodriguez and Mollner 2010), and France (Gayral 1966) north to the UK (Hardy and Guiry 2003) as well as Iceland (Gunnarsson and Jónsson 2002). The Icelandic specimens have now been shown to belong to a separate species and apart from the *S. apoda* sequences from the Azores, only one *rbcL* sequence is available in GenBank of *Schizymenia* from the Atlantic coast of Europe. Discovering the new species *S. jonssonii* in Iceland and finding *S. apoda* in the Plymouth area in Britain emphasizes the need for further molecular studies on *Schizymenia* specimens from the area to establish the identity and distribution of *Schizymenia* spp. in the North Atlantic and the Mediterranean Sea.

Both *Schizymenia apoda* and *S. dubyi* have been found at several sites in the Pacific (Hughey and Miller 2009, Gabriel et al. 2011, Kim et al. 2012, D’Archino and Zuccarello 2013, Saunders et al. 2015), and *S. dubyi* has also been found on the Atlantic coast of Argentina (Ramirez et al. 2012). In some cases, they are thought to have been recently introduced (Ramirez et al. 2012, D’Archino and Zuccarello 2013).

As there are still five additional species of *Schizymenia* listed in Algaebase (Guiry and Guiry 2019) for which there are no sequence data yet, the question remains as to whether one of them might be synonymous with *S. jonssonii*. Two of them, *S. violacea* and *S. johnstonii* (Setchell and Gardner 1924) from the Gulf of California are considered to be synonymous and belong to the genus *Grateloupia*, as *G. violacea* (Dawson 1944, 1961, Gabrielson et al.
2019). *Schizymenia obliqua* was found on the island St Paul in the Indian Ocean and is only known from its original collection by Grunow (1868, Silva et al. 1996). This species was first described as a variety of *S. erosa*, differing "just" in the form of the blade that also resembled *Iridaea curvata* (Grunow 1868). Kylin (1932) found that the original specimen of *S. erosa* was a male plant belonging to the genus *Iridaea*. *Schizymenia ecuadoreana* is found in the Galapagos Islands. It is a deep-water species with distinctive gland cells (Taylor 1945, Abbott 1967). *Schizymenia binderi* is found on the Pacific coast of South America (Papenfuss 1964, Ramirez and Santelices 1991) and also possesses gland cells (Kylin 1932). Considering the strict subtidal habitat of the two previously

![Fig. 6. Maximum likelihood tree for *Schizymenia* spp. using *rbl* sequences with *Platoma cyclocolpum* and *Titanophora weberae* as outgroup. Values at nodes: ML bootstrap values/Bayesian posterior probability. Sequences produced in the present study are in bold. Scale bar = 0.01 substitutions per site. *Specimen has also been sequenced for COI and is shown in Fig. 5. *Tetrasporophyte. For details on the specimens, from which the sequences were obtained, see supplementary data in Table S1."

named species, with plants growing to a depth of 55 m (Taylor 1945), distinct gland cells and their distribution in the southern hemisphere, they are unlikely to be synonymous with, or the source of the *Schizymenia* population in Iceland.

The DNA sequence analysis shows a clear separation of two phylogenetically distinct species groups of the genus *Schizymenia*, one represented by the species *S. dubyi* and *S. apoda* and the other with the closely related species *S. pacifica*, *S. tenuis*, "Schizymenia. sp._1Cal," and the new species *S. jonssonii* (Figs. 5 and 6). This indicates a firm and extended geographical isolation of the two clades, perhaps, between the Atlantic and the Pacific and that *S. jonssonii* has arrived in the Atlantic after the
evolutionary separation of the clades and that S. apoda and S. dubyi are possibly inversely, relatively recent introductions into the Pacific.

The results raise the question of how old the introduction and the genetic separation of Schizymenia jonssonii from its Pacific relatives is likely to be. Schizymenia jonssonii possibly has an ancestor of Pacific origin and colonized the Atlantic after the opening of the Bering Strait, evolved in the Atlantic and is a relic from the last glacial maximum (LGM). But it is equally possible that it is a relatively recent introduction into the Atlantic. The following two scenarios could possibly explain the presence of the new species in Iceland. A: The species passed through the Bering Strait during ancient warm periods after 3.5 million years BP and colonized the North Atlantic. The opening of the Bering Strait is dated at about 5.5 million years BP (Gladenkov et al. 2002) while inflow of biota from the Pacific into the Atlantic possibly started much later or at about 3.5 million years BP (Vermeij 1991). The Icelandic coast is thought to have been completely covered by ice during the LGM in the North Atlantic, from 26,500 years BP to at least about 14,600 to 12,700 years BP (Ingólfsson et al. 2010). If the species is to have survived in the Atlantic for a more extended period, it would have been pushed southwards by the glaciation at least to the English Channel (Sejrup et al. 2005, Carr et al. 2006) and when the glacier started retreating it would have moved toward the Arctic in the wake of the deglaciation. If this was the case, it would be expected for there to be traces of the species left behind south of Iceland (e.g., in the Faroes, southern Scandinavia or the UK). However, since then it appears to have died out south of Iceland and in the Pacific (or the species is still living cryptically further south in Europe and/or in the Pacific). B: The species colonized Iceland from the Pacific after LGM. The source population in the Pacific died out (or a cryptic source population still exists in the Pacific). Only more molecular studies of Schizymenia from more locations in the North Pacific will help solve this puzzle.

Until now only Schizymenia dubyi had been recorded for Britain and Ireland. The finding of S. apoda in England raises the question of the identity of Schizymenia in the region. To resolve this problem, more sequence data are needed from contemporary and historical material. This has been proved to be valuable for the study of algal diversity (this study and Gunnarsson et al. 2016).

Considering the discovery of the new species in Iceland, the presence of Schizymenia apoda in England, and the extreme difficulty in distinguishing species within the genus, it is likely that there is greater diversity of Schizymenia spp. at the local level in the north-eastern Atlantic.

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest.
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Supporting Information

Additional Supporting Information may be found in the online version of this article at the publisher’s web site:

Table S1. GenBank accession-numbers, species, voucher identity, sampling localities, collectors and collecting dates for specimens used in molecular phylogenetic analysis. Sequences that were generated in the present study are in bold.