

# Species delimitation and DNA barcoding of Atlantic *Ensis* (Bivalvia, Pharidae)

Joaquín Vierna<sup>a,b</sup>, Joël Cuperus<sup>c</sup>, Andrés Martínez-Lage<sup>a</sup>, Jeroen M. Jansen<sup>c</sup>, Alejandra Perina<sup>a,b</sup>, Hilde Van Pelt<sup>c</sup>, Ana M. González-Tizón<sup>a</sup>

<sup>a</sup> Evolutionary Biology Group (GIBE), Department of Molecular and Cell Biology, Universidade da Coruña, A Fraga 10, A Coruña, E-15008, Spain

<sup>b</sup> AllGenetics, Edificio de Servicios Centrales de Investigación, Campus de Elviña s/n, A Coruña, E-15008, Spain

<sup>c</sup> IMARES Wageningen UR, Ambachtsweg 8a, Den Helder, NL-1785 AJ, The Netherlands

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## Abstract

*Ensis* Schumacher, 1817 razor shells occur at both sides of the Atlantic and along the Pacific coasts of tropical west America, Peru, and Chile. Many of them are marketed in various regions. However, the absence of clear autapomorphies in the shell and the sympatric distributions of some species often prevent a correct identification of specimens. As a consequence, populations cannot be properly managed, and edible species are almost always mislabelled along the production chain. In this work, we studied whether the currently accepted Atlantic *Ensis* morphospecies are different evolutionary lineages, to clarify their taxonomic status and enable molecular identifications through DNA barcoding. For this, we studied 109 specimens sampled at 27 sites, which were identified as belonging to nine of those morphospecies. We analysed nucleotide variation at four nuclear (18S, 5.8S, ITS1, and ITS2) and two mitochondrial (COI and 16S) regions, although the 18S and 5.8S regions were not informative at the species level and were not further considered. The phylogenetic trees and networks obtained supported all morphospecies as separately evolving lineages. Phylogenetic trees recovered *Ensis* at each side of the Atlantic as reciprocally monophyletic. Remarkably, we confirm the co-occurrence of the morphologically similar *E. minor* (Chenu, 1843) and *E. siliqua* (Linné, 1758) along the NW Iberian coast, a fact that has been often overlooked. In South America, a relevant divergence between *E. macha* (Molina, 1792) individuals from Chile and Argentina was unveiled and suggests incipient speciation. We also confirm the occurrence of the North American species *E. directus* (Conrad, 1843) as far south as north-eastern Florida. Among the genomic regions analysed, we suggest COI as the most suitable DNA barcode for Atlantic *Ensis*. Our results will contribute to the conservation and management of *Ensis* populations and will enable reliable identifications of the edible species, even in the absence of the valves. The name *Ensis coseli* Vierna nom. nov. is proposed to replace *E. minor* Dall, 1899 non (Chenu, 1843).

**Keywords:** Razor clam, COI, 16S rDNA, 18S rDNA, phylogeny

## Introduction

*Ensis* Schumacher, 1817 razor shells are a group of marine bivalve molluscs, characterised by their elongated shells, which are often found along the coasts at both sides of the Atlantic Ocean and along the Pacific coasts of tropical west America, Peru, and Chile. Several species are considered a delicacy and are marketed in some European, South American, and North American regions. For instance, in Chile alone, almost 6000 tons of *E. macha* (Molina, 1792) were landed in 1999 (Barón *et al.* 2004) and 1400 tons during 2005 (Ariz Abarca *et al.* 2007).

In the Atlantic, the genus is composed of 9–10 extant morphospecies that inhabit sandy and fine gravel substrata with limited exposure to wave action, from the intertidal to a depth of ca. 80 m (Cosel 2009). *Ensis goreensis* (Clessin, 1888) occurs in tropical West Africa, from the southern part of West Sahara to southern Angola (Lucira) and the Cape Verde Islands (Cosel 2009; and references therein). In Europe, there are four extant native *Ensis*, namely *E. magnus* Schumacher, 1817 [syn. *E. arcuatus* (Jeffreys, 1865)], *E. ensis* (Linné, 1758), *E. minor* (Chenu, 1843), and *E. siliqua* (Linné, 1758) (Cosel 2009). The American *E. directus* (Conrad, 1843) [syn. *E. americanus* (Gould, 1870)] is native to Atlantic North America, but was introduced to European coastal waters in 1978 (Cosel *et al.* 1982), where it now occurs from the North Sea to the Cantabrian Sea (Arias & Anadón 2012; Vierna *et al.* 2012). Another representative from Atlantic North America is the recently discovered *E. terranovensis* Vierna & Martínez-Lage, 2012, which was found off Newfoundland (Canada) and whose possible co-occurrence with *E. directus* is still unknown (Vierna *et al.* 2012). *Ensis macha* occurs along the southern coasts of Argentina, Peru, and Chile, and the remaining two taxa, *E. minor* Dall, 1899, and *E. minor megistus* Pilsbry & McGinty 1943, are native to the south-eastern coast of the USA (mainly Florida and the Gulf of Mexico).

The name *E. minor* Dall, 1899 is a junior homonym of the European *E. minor* (Chenu, 1843), but so far, without replacement name. Here, we propose the name *Ensis coseli* Vierna nom. nov. to replace *E. minor* Dall, 1899 non (Chenu, 1843). The new species name is in honour of Dr. Rudo von Cosel, for his contribution to the taxonomy and systematics of razor shells. *Ensis minor* Dall, 1899 has recently been renamed into *E. megistus* Pilsbry & McGinty, 1943, based on the synonymisation of *E. minor megistus* Pilsbry & McGinty, 1943 with the nominotypical subspecies (Huber 2010), which seems to make the proposition of a replacement name obsolete. However, after having studied the *E. minor megistus* type material and having compared it with specimens from some museum collections identified as *E. minor* sensu Dall (J. Vierna, A. M. González-Tizón, and A. Martínez-Lage, unpublished), we found a clear difference between both taxa in the position of the posterior adductor scar, which is a relevant taxonomic character in *Ensis* (see Cosel 2009; and Vierna *et al.* 2012). This scar is situated much more anterior to the pallial sinus in *E. minor megistus* than in the *E. minor* sensu Dall specimens examined. Therefore, we reject the synonymy in Huber (2010), which was provided without evidence, and we maintain *E. coseli coseli* and *E. coseli megistus* Pilsbry & McGinty, 1943 as subspecies, pending further studies, which may reveal species status for both. Molecular data of *E. coseli megistus* are currently unavailable, because to our best knowledge, there are no specimens available other than the valves that comprise the type material (see Pilsbry & McGinty 1943).

Despite their economic value, *Ensis* taxonomy is not well defined, due to the absence of clear autapomorphies in the shell and the sympatric distributions of some species. In fact, Cosel (2009) found some specimens that he considered as ‘intergrades’ of some European morphospecies, based on the analysis of the valves. This poorly defined taxonomy seems to be the cause of the mislabelling of specimens sold in European markets, where razor shells are often labelled as ‘*Ensis ensis*’ even though they usually are either *E. magnus*, *E. siliqua*, *E. minor*, or *E. directus* (pers. obs).

We have noticed that several studies (Arias *et al.* 2011; Arias-Pérez *et al.* 2012; Varela *et al.* 2012; Rufino *et al.* 2012) have overlooked the fact that the European morphologically similar *E. minor* and

*E. siliqua* occur sympatrically along Atlantic Europe. According to the review by Cosel (2009) based on shell morphology, *E. minor* occurs in both Atlantic and Mediterranean Europe, whereas *E. siliqua* is restricted to the Atlantic. However, the Galician Regional Government ('Xunta de Galicia', NW Spain), does not distinguish between these two taxa, even though fishing and commercialisation of these bivalves are strictly regulated in the area. Interestingly, González-Tizón *et al.* (2013) supported both taxa as separate species according to cytogenetics.

Phylogenetic analysis of DNA sequences is a common and useful approach for species delimitation (e.g. Fontaneto *et al.* 2011; Puillandre *et al.* 2012; Ornelas-Gatdula *et al.* 2012; Esselstyn *et al.* 2012) that should be considered in the light of other evidences such as morphology, biogeography, and ecology (integrative taxonomy, see Dayrat 2005). Even though the term 'species delimitation' is sometimes used in a similar way to 'DNA barcoding' (the use of standard genomic regions for identifying organisms to the species level, Hebert *et al.* 2003), for clarity reasons, in this paper, we make the distinction between these approaches.

Hebert *et al.* (2003) proposed to use a fragment of the so-called COI mitochondrial gene as the standard barcode for animals, but they recognised that supplemental analyses of one or more nuclear genes could be required when hybridisation or introgression occurs. Therefore, to delimit species and to select a suitable genomic region that can be used as a DNA barcode, in this paper, we analyse variation not only at COI, but also at other mitochondrial and nuclear regions, in more than one hundred specimens that were identified as belonging to the Atlantic morphospecies. If this work is taken into account by policymakers, it will contribute to the conservation and management of *Ensis* populations and will also enable a reliable labelling of the edible species, even in the absence of the valves (i.e. when processed as seafood), as well as a reliable identification of larvae and juvenile stages.

## **Materials and methods**

### Specimen identifications

We considered all currently known extant Atlantic *Ensis* morphospecies according to Cosel (2009) and Vierna *et al.* (2012). We studied a total of 109 specimens from 27 sampling sites (Table S1). Razor shell samples were obtained from museum collections and from colleagues, or they were directly sampled by us. Initial morphological identification of specimens was performed following Cosel (2009) and Vierna *et al.* (2012), and these were confirmed after phylogenetic analyses. Specimens were deposited in the collections of various natural history museums (see Table S1).

### DNA isolation, PCR, and sequencing

Razor shell specimens were preserved either frozen or in 100% ethanol, except the *E. goreensis* sample. In this case, dry tissue was obtained from the interior part of the shell and rehydrated in sterile milli-Q water.

DNA was extracted from muscle tissue using the NucleoSpin Tissue kit (Macherey-Nagel GmbH and Co. KG, Düren, Germany). The mitochondrial regions studied were fragments of the cytochrome oxidase subunit I gene (COI) and the 16S ribosomal RNA gene (16S). The nuclear ribosomal DNA genes and spacers considered were the internal transcribed spacers 1 and 2 (ITS1 and ITS2), the 5.8S ribosomal gene (5.8S), and a fragment of the 18S ribosomal gene (18S). The ITS1-5.8S-ITS2 region was amplified, cloned, and sequenced as a whole. The COI region was sequenced in all 109 specimens, whereas the five other regions were sequenced in a subset of specimens (Table S1).

Using the COI 'universal' primers LCO1490 and HCO2198 (Folmer *et al.* 1994), we obtained some sequences that were then input in GeneFisher (Giegerich *et al.* 1996) to design four additional species-

specific internal primer pairs (COI-Europe-1-F, 5' GGG ATT AGT TGG GAC TAG G; COI-Europe-1-R, 5' GTT AAA GCC CCT GCC AA; COI-Europe-2-F, 5' TAG AGT TAG CTC GTC CT; COI-Europe-2-R, 5' AAA TAG GGT CAC CAC CA; COI-*macha*-F, 5' TAG TTG GGA CTA GGT TGA GA; COI-*macha*-R, 5' TAG GAT CTC CTC CAC CTC T; COI-*coseli*-F, 5' GAT TCG GTT AGA GTT AGC TCG A; and COI-*coseli*-R, 5' GTT AAA GCA CCA GCT AGT ACA G). These new primers and the COI-*directus* ones (Vierna *et al.* 2012) were used in all COI amplification reactions. The primers used to amplify the 16S region were 16Sar and 16Sbr (Palumbi 1996). For the ITS1-5.8S-ITS2 region, we used the primers by Heath *et al.* (1995). Finally, for the 18S, we used the primers annealing at the 5' and 3' ends of the gene (Winnepeninckx *et al.* 1994). With the sequences obtained, we designed a pair of internal primers (18SintF, 5' GAT CGT ACA ATC CTA CTT GG; and 18SintR, 5' GCT CAT TAA CGG GAA CGA T) in GeneFisher. These new primers were employed in one of the *E. magnus* specimens analysed.

Each PCR (25  $\mu$ L) contained ~25 ng of genomic DNA, 0.625 U of Taq DNA polymerase (Roche Diagnostics, Switzerland), 5 nmol of each dNTP (Roche Diagnostics), 20 pmol of each primer and the buffer recommended by the polymerase supplier. The general reaction conditions were as follows: an initial denaturation step at 94 °C for 3 min followed by 35 cycles of denaturation at 94 °C for 20 s; annealing at the following temperatures (LCO1490/HCO2198, 45 °C; COI-Europe-1, 57 °C; COI-Europe-2, 49 °C; COI-*macha*, 55 °C; COI-*coseli*, 52 °C; COI-*directus*, 48 °C; 16Sar/16Sbr, 44 °C; ITS1-5.8S-ITS2 region, 59 °C; 18S (Winnepeninckx *et al.* 1994), 56 °C; and 18Sint, 55 °C) for 20 s; extension at 72 °C for 30–50 s; and a final extension at 72 °C for 5 min. PCR products were run on 1 % agarose gels, stained with either ethidium bromide or Real Safe (Real, Valencia, Spain) and imaged under UV light.

All PCRs yielded single-band patterns, and therefore, amplicons were directly sequenced (except the ITS1-5.8S-ITS2 region ones), after being purified with ExoSAP-IT (USB, Santa Clara, CA, USA). PCR primers were used to sequence amplicons in both directions. In the case of 18S amplicons, we designed internal sequencing primers (18SseqF, 5' CCC GTA ATT GGA ATG AGT AC; and 18SseqR, 5' CGA ATC AAG AAA GAG CTC TC) to cover the whole amplified region. Due to the occurrence of intragenomic variation (Vierna *et al.* 2010), ITS1-5.8S-ITS2 PCR products were cloned using the TOPO TA cloning kit (Invitrogen, Paisley, UK). Transformant colonies were selected, and insert size was checked by PCR. We spread one clone per individual on an LB plate and let it grow overnight at 37 °C. Plasmids were purified with the QiaPrep Spin Miniprep Kit (Qiagen, Hilden, Germany), and they were sequenced using the M13 forward and reverse primers (supplied with the cloning kit). In addition to the sequences generated, some *E. directus* and *E. terranovensis* COI, ITS1, and ITS2 sequences (Vierna *et al.* 2012) from specimens sampled at Cobscook Bay, Long Pond, and The Wash were included in our data sets (see Table S2 for accession numbers). New DDBJ/EMBL/GenBank accession numbers are HF970346-HF970575 and HF975604-HF975627.

### Bioinformatic analyses

The software BioEdit 7.0.9.0 (Hall 1999) and Geneious Pro 5.4.6 (Drummond *et al.* 2011) were used to examine the electropherograms. To search for stop codons that would be indicative of the presence of pseudogenes, the COI amino acid sequences were obtained from MEGA 5.03 (Tamura *et al.* 2011) using the 'invertebrate mitochondrial genetic code'.

COI and 16S alignments were carried out in ClustalW 2.0 (Larkin *et al.* 2007) under default parameters. ITS1 and ITS2 sequences were aligned using the Q-INS-i strategy as implemented in MAFFT, version 7 (Katoh & Toh 2008; Katoh & Standley 2013). Highly variable regions were deleted from ITS1 and ITS2 alignments using Gblocks (Castresana 2000; Talavera & Castresana 2007) available at [http://molevol.cmima.csic.es/castresana/Gblocks\\_server.html](http://molevol.cmima.csic.es/castresana/Gblocks_server.html) under default (conservative) options. All alignments were trimmed, and identical sequences were collapsed into sequence-types using DnaSP 5.10.01 (Librado & Rozas 2009). Multigene alignments (COI+16S; ITS1+ITS2) were obtained as follows: we

aligned each genomic region independently; then, we concatenated the sequences obtained from each specimen, and finally, we collapsed identical multigene sequences into sequence-types.

Phylogenetic trees were inferred under Bayesian (BA) and maximum-likelihood (ML) methods. BA was carried out using the software MrBayes 3.1.2 (Huelsenbeck & Ronquist 2001) from the CIPRES Science Gateway (Miller *et al.* 2010). Models of evolution for each partition (Table S3) were obtained from MrModelTest 2.3 (Johan Nylander, <http://www.abc.se/~nylander/>). The analysis was performed with 15 000 000 generations initiated with a random starting tree, sampling every 1000 generations, and allowing the program to estimate the likelihood parameters required. Stationarity was assessed using the Web-based software AWTY (Nylander *et al.* 2008). Results collected prior to stationarity were discarded as burn-in. ML phylogenies were obtained using RAxML 7.2.8 (Stamatakis 2006; Stamatakis *et al.* 2008) that was run from the same bioinformatic platform. This software is capable of assigning and estimating separate model parameters for individual genes of multigene alignments (Stamatakis 2006) and implements the general time-reversible (GTR) substitution model for all partitions. Node confidence was assessed using 1000 nonparametric bootstrap replicates (Felsenstein 1985). All phylogenetic trees were edited in Dendroscope 3 (Huson & Scornavacca 2012).

The best resolved phylogenetic trees based on the CO1+16S and ITS1+ITS2 data sets were subjected to a general mixed Yule coalescent analysis (GMYC) (Pons *et al.* 2006; Fontaneto *et al.* 2007). The model was performed with R 2.15.3 (R Development Core Team 2011) package splits (SPecies' LLimits by Threshold Statistics) available at <http://r-forge.r-project.org/projects/splits/>. The required ultrametric phylogenetic trees were generated using penalised likelihood in r8s 1.70 and mid-point rooted (Sanderson 2003; Obertegger *et al.* 2012).

Even though we considered the possibility of including *Pharus legumen* (Linné, 1758) and *Siliqua patula* (Dixon, 1789) sequences as outgroups, they were too much divergent to reliably align them in the case of some genomic regions (ITS1 and ITS2). Therefore and considering that it is not necessary to use outgroups to check whether morphospecies are separately evolving lineages (e.g. Ley & Hardy 2010; James *et al.* 2010), we decided against using outgroup sequences, and thus, all the phylogenetic trees we present here are unrooted.

We also obtained phylogenetic networks for each alignment using the Neighbor-Net algorithm (Bryant & Moulton 2004) and uncorrected p-distances in SplitsTree4 (Huson & Bryant 2006). Finally, the mean intraspecific and interspecific uncorrected p-distances were obtained from MEGA 5.03. Their standard errors were calculated by 1000 bootstraps.

## Results

In general, morphological identifications were confirmed by molecular data. However, the morphologically similar *E. minor* and *E. siliqua* were rather difficult to identify reliably in terms of shell morphology using available keys (Cosel 2009). Nonetheless, we were able to clearly differentiate them from the other European morphospecies *E. magnus* and *E. ensis*. Therefore, they were identified both using phylogenetic clustering and geographic information: because according to Cosel (2009), *E. siliqua* is restricted to the Atlantic, whereas *E. minor* occurs both in the Atlantic and in the Mediterranean, the Atlantic specimens that clustered with the Mediterranean ones were considered as *E. minor*.

Interestingly, two specimens from Jacksonville (FL, USA) that we received from the Florida Museum of Natural History, labelled as *E. minor* Dall, 1899 (*E. coseli coseli*), resulted as belonging to *E. directus*, according to the shape of muscle scars on the inner valves (Cosel 2009) and to phylogenetic clustering.

These specimens are, to our knowledge, the first confirmed record of *E. directus* as far south as north-eastern Florida.

The COI alignment (303 bp) did not display, as expected, any gap. No stop codons were found, and there were only three mutations (from three different sequences) that lead to amino acid substitutions. The proportion of variable sites ( $s$ ) for this region was  $s = 0.337$ . Alignments of 16S (340 bp), 18S (1271 bp) and 5.8S (157 bp) sequences were straightforward. The proportion of variable sites was  $s = 0.126$ ,  $s = 0.009$  and  $s = 0.032$ , respectively. There was only one gap position, in the 18S region.

Regarding the alignments of ITS1 and ITS2 sequences, 285 (44 % of the original 636 positions) were retained after the Gblocks filtering in the case of ITS1. For ITS2, 228 (62 % of the original 364 positions) were kept after filtering. The proportion of polymorphic sites after filtering was  $s = 0.154$  and  $s = 0.224$ , respectively.

The number of haplotypes obtained from the COI alignment was 81 of 109 sequences (74.3 %), whereas the 16S alignment yielded 29 haplotypes of 80 sequences (36.2 %). The 18S region produced six sequence-types of 13 sequences (46.1 %), the 5.8S region yielded only five sequence-types of 41 sequences (12.2 %), and both ITS1 and ITS2 regions yielded 19 of 41 sequences (46.3 %) each. The COI+16S multigene alignment produced 70 haplotypes of 80 sequences (87.5 %), whereas the ITS1+ITS2 alignment yielded 29 sequence-types of 41 sequences (70.7 %).

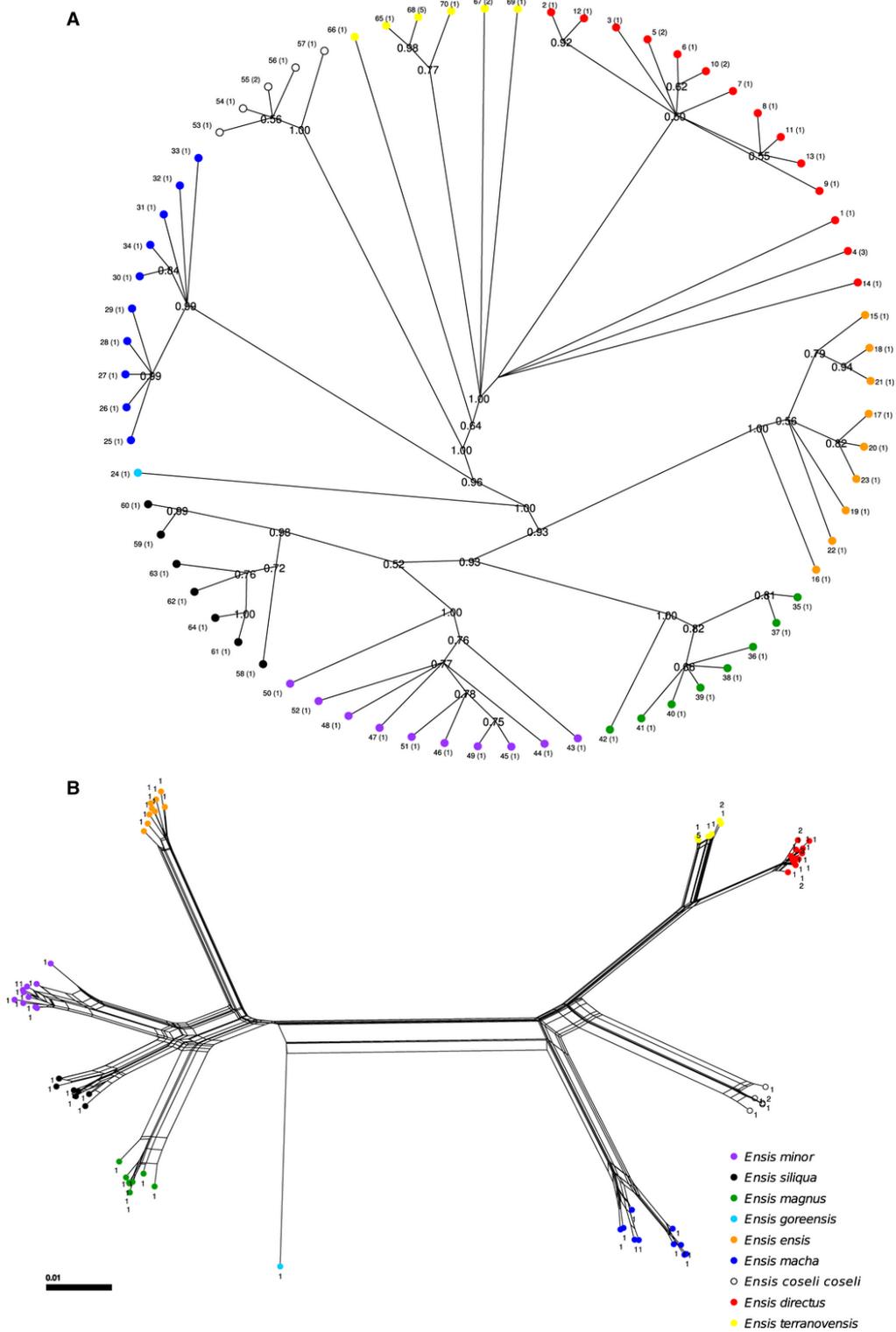
Sequence-type distributions of the 18S and 5.8S regions (that due to their conservation, lacked resolution at the species level and were not considered in the subsequent analyses) were as follows: *E. magnus*, *E. siliqua*, and *E. minor* shared the same 18S sequence-type. However, *E. directus*, *E. terranovensis*, *E. macha*, *E. coseli coseli*, and *E. ensis* had one (non-shared) sequence-type each. The two individuals of *E. macha* from Chile and Argentina whose 18S region was sequenced displayed the same sequence. In the case of the 5.8S region, one of the five sequence-types obtained was shared by all North and South American species. Another one was shared by all European species. The remaining three corresponded to one *E. terranovensis*, one *E. directus*, and one *E. magnus* sequence.

All phylogenetic trees obtained, based either on mitochondrial or on nuclear DNA, recovered morphospecies at each side of the Atlantic as reciprocally monophyletic (Figs 1 and 2; Figs S1-S6).

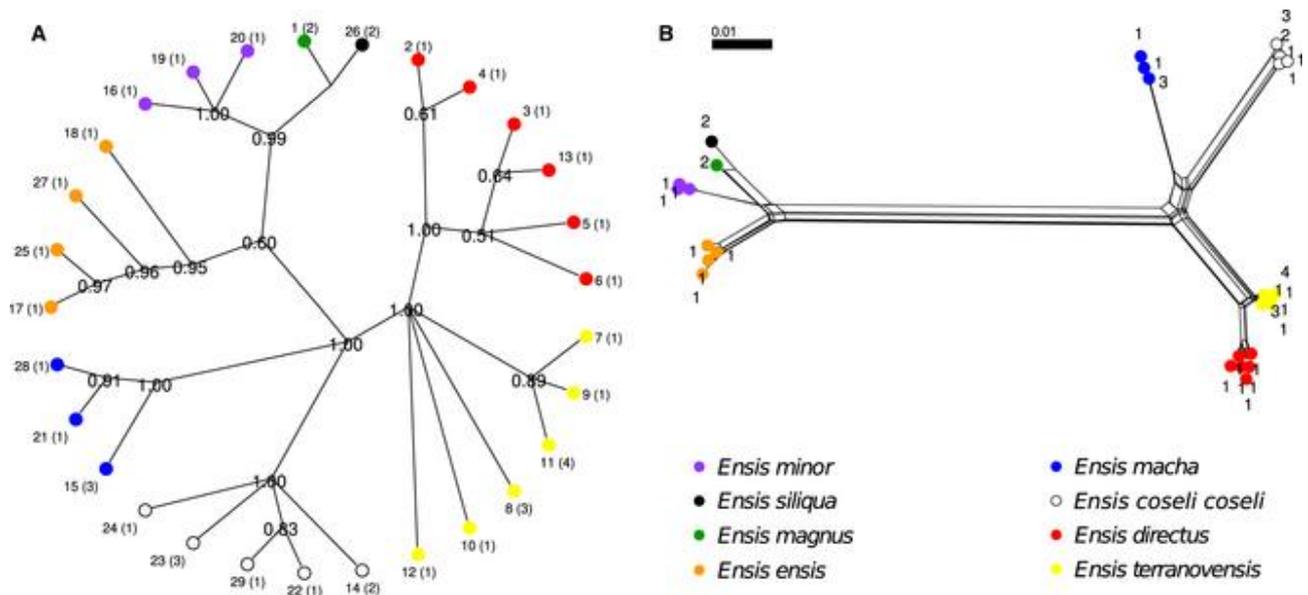
Trees based on the COI alignment recovered all morphospecies as monophyletic, with the exception of *E. terranovensis* in the BA analysis, and both *E. terranovensis* and *E. magnus* in the ML analysis (Fig. S1). The morphospecies *E. goreensis* was represented by only one haplotype.

The 16S alignment produced one haplotype that was shared between *E. terranovensis* and *E. directus*. Both BA and ML phylogenetic trees clustered *E. magnus* and *E. ensis* sequences on monophyletic groups. As for *E. coseli coseli* and *E. goreensis*, only one haplotype was obtained per morphospecies. In the BA analysis, sequences from *E. macha*, *E. siliqua*, and *E. minor* were clustered in monophyletic groups as well (see Fig. S2).

The BA tree reconstructed from the COI+16S multigene alignment recovered all morphospecies as monophyletic, except for one *E. terranovensis* haplotype, which formed a polytomy (Fig. 1). The support values for each of those clades ranged between 0.98 and 1.00 except for *E. terranovensis* and *E. directus*. The ML analysis recovered as monophyletic all the European and African species with the exception of *E. ensis*. In contrast, the sequences from American species did not cluster, in general, into monophyletic groups in the ML tree. Rather, they formed some polytomies, with the exception of the *E. coseli coseli* sequence-types (see Fig. S3).



**Figure 1.** Phylogenetic relationships among mitochondrial (COI+16S) haplotypes. Haplotype frequencies are shown in parentheses in the tree, and at each terminal node, in the network. Terminal nodes are coloured according to which *Ensis* species the sequence was obtained from. —A. Unrooted Bayesian phylogenetic tree. Node confidence values below 0.5 are not shown. —B. Phylogenetic network constructed using the Neighbor-Net algorithm and uncorrected p-distances.



**Figure 2.** Phylogenetic relationships among nuclear (ITS1+ITS2) sequence-types. Sequence-type frequencies are shown in parentheses in the tree, and at each terminal node, in the network. Terminal nodes are coloured according to which *Ensis* species the sequence was obtained from. —A. Unrooted Bayesian phylogenetic tree. Node confidence values below 0.5 are not shown. —B. Phylogenetic network constructed using the Neighbor-Net algorithm and uncorrected p-distances.

No *E. goreensis* ITS1-5.8S-ITS2 sequences could be obtained, probably due to degradation of the extracted DNA, and therefore, we lack nuclear DNA data for this morphospecies. Regarding the ITS1 analysis, sequences from *E. macha*, *E. siliqua*, *E. magnus*, and *E. minor* yielded only one sequence-type per species. Sequences from the other morphospecies were clustered into monophyletic groups, in both BA and ML phylogenetic trees (see Fig. S4).

In the ITS2 analysis, *E. siliqua*, *E. magnus*, and *E. ensis* yielded one sequence-type each. The BA tree recovered all morphospecies as monophyletic with the exception of *E. directus* and *E. terranovensis*, which formed two polytomies. These two species showed a paraphyletic pattern in the ML phylogeny, whereas all others were recovered as monophyletic (see Fig. S5).

Finally, in the ITS1+ITS2 analysis, there were two morphospecies (*E. magnus* and *E. siliqua*) that yielded only one sequence-type each. The BA phylogenetic tree recovered all other morphospecies as monophyletic, with the exception of *E. terranovensis*, which formed a polytomy. Support values for each of these clades ranged between 0.95 and 1.00. In the ML tree, all morphospecies were recovered as monophyletic, and support values were high (94–100) except for *E. terranovensis* and *E. minor* (see Fig. 2; Fig. S6).

The GMYC analysis based on the BA COI+16S phylogenetic tree revealed a model with eight different entities (confidence interval 8–11) as the most likely (likelihood of the null model, 338.693; maximum likelihood of the GMYC model, 352.236; likelihood ratio, 27.085; LR test < 0.0001). The number of morphospecies included in the analysis (nine) falls within the confidence interval. Therefore, the GMYC analysis based on the COI+16S phylogeny is consistent with the existence of the nine different species.

The GMYC analysis based on the ML ITS1+ITS2 phylogenetic tree revealed a model with eight different entities (confidence interval 7–8) as the most likely (likelihood of the null model, 113.846; maximum likelihood of the GMYC model, 120.472; likelihood ratio, 13.253; result of the LR test, 0.004). In the same

way as for the COI+16S phylogeny, the GMYC analysis based on the ITS1+ITS2 data set is consistent with the existence of eight different species (in this case, there were no *E. goreensis* sequences available).

The phylogenetic networks obtained based on the COI, COI+16S, ITS1, and ITS1+ITS2 alignments clearly supported the existence of groups that perfectly matched the currently described morphospecies (Figs S1 and S4; Figs 1 and 2).

The 16S region was not able to completely differentiate among North American species because, as already mentioned, there was one haplotype shared by two of them. Regarding European species, the two *E. siliqua* haplotypes were not clustered together in the network (see Fig. S2).

The ITS2 phylogenetic network clustered the sequence-types obtained from *E. terranovensis* and *E. directus* specimens and also the ones obtained from *E. siliqua*, *E. magnus*, and *E. ensis* (see Fig. S5).

We calculated the intraspecific and interspecific mean p-distance values for the COI, 16S, ITS1, and ITS2 regions using the trimmed alignments containing all sequences (i.e. not the sequence-type alignments; Tables S4–S7).

We also calculated interpopulation mean p-distances for *E. macha* individuals from Puerto Lobos (Argentina) and Playa Dichato (Chile). In this case, no ITS1 or ITS2 values were obtained because the number of sequences available was too small. Divergence between the two *E. macha* populations was in some cases more pronounced than interspecific divergence. For instance, 16S interpopulation divergence was  $0.011 \pm 0.004$ , whereas interspecific divergence in some species pairs was equal or lower: *E. directus* and *E. terranovensis*,  $0.003 \pm 0.003$ ; and *E. siliqua* and *E. magnus*,  $0.011 \pm 0.005$  (Table S5). Similarly, COI interpopulation divergence was  $0.038 \pm 0.010$ , quite comparable to that calculated for *E. directus* and *E. terranovensis* ( $0.058 \pm 0.012$ ) and for *E. siliqua* and *E. magnus* ( $0.064 \pm 0.012$ ; Table S4).

## Discussion

### Species delimitation and molecular systematics

Our results support that the nine morphospecies under study are different evolutionary lineages, according to both nuclear and mitochondrial DNA, and in agreement with Cosel (2009). Therefore, they should be considered as nine different species.

The split between *E. directus* and *E. terranovensis* was already demonstrated by Vierna *et al.* (2012) using morphometrics and nucleotide variation at COI, ITS1, ITS2, and ANT (a fragment of a nuclear single-copy region coding adenine nucleotide translocase). However, in the present work, the split between the two species was not evident for all genomic regions analysed, because in some phylogenetic trees *E. terranovensis*, sequences formed polytomies instead of clustering in a clade. These polytomies could reflect that the two sister species underwent speciation rather recently, a fact that is further supported by the shared 16S haplotype, which is an example of incomplete lineage sorting among closely related taxa.

Despite the absence of tropical west American *Ensis* in our analyses, we have unveiled a strong phylogeographic structure within genus *Ensis*. Species at each side of the Atlantic are reciprocally monophyletic, as shown by molecular data as well as for the shape of the pallial sinus (broad and shaped like an irregular W in American species and narrower and more or less rounded in European species, Cosel 2009).

In Atlantic North America, three lineages were found: whereas *E. directus* and *E. terranovens* were supported as sister species, the position of *E. coseli coseli* was variable. This taxon branched off as sister to *E. directus* and *E. terranovens* in most of the analyses, supporting the monophyly of North American species, and as sister to *E. macha*, in some others. Interestingly, we have confirmed the occurrence of *E. directus* as south as north-eastern Florida.

In South America, *E. macha* specimens sampled at Playa Dichato and Puerto Lobos were rather divergent, again showing phylogeographic structure. The number of individuals analysed was small and prevents any general conclusion about possible population structure of these razor shells along their distributional ranges. However, the degree of divergence displayed by individuals from Playa Dichato and Puerto Lobos according to COI and 16S was in some cases even higher than that obtained for some other *Ensis* interspecific comparisons, and suggests incipient speciation. Significant population structure has also been found in marine gastropod *Nacella magellanica* (Gmelin, 1791) populations from Atlantic and Pacific Patagonia (González-Wevar *et al.* 2012). Taken into account that Camus (2001) identified a major biogeographic break in Chile at 41–43°S and that Playa Dichato is located in the north of these parallels, this biogeographic break could also explain population structure in *E. macha*. Our results highlight the need to study the population structure and diversity of this razor shell along its entire distribution (a survey on shell morphometry along the Patagonian coast is available, Márquez & Van der Molen 2011) to design conservation plans for this intensively fished species (see Hernández *et al.* 2011).

Unfortunately, the phylogenetic relationships between species within the European/African clade were not well resolved as evidenced by the presence of polytomies and low node support values.

Remarkably, we have demonstrated that *E. siliqua* and *E. minor* are different species, despite their morphological similarities. Our results agree with the taxonomic revision by Cosel (2009) based on shell morphology and with González-Tizón *et al.* (2013) who found important differences among the karyotypes of both species.

Some of the specimens studied in this work were also analysed in two previous papers on evolutionary genetics. According to our results, some sequences that were thought to have been obtained from *E. siliqua* specimens (Vierna *et al.* 2009, 2011) were in fact obtained from *E. minor* razor shells. The sequence labels have been updated on DDBJ/EMBL/GenBank. Fortunately, we do not expect that this misidentification of specimens impacts any of the main conclusions of those papers. However, this issue accentuates the importance of DNA barcoding in research fields other than ecology and biodiversity.

Given that both *E. siliqua* and *E. minor* co-occur in Atlantic Europe and that it is not always possible to reliably differentiate among specimens based on a visual inspection of the shell, we suspect that in some previous reports on *E. siliqua* (Darriba *et al.* 2005; Arias *et al.* 2011; Arias-Pérez *et al.* 2012; Varela *et al.* 2012; Rufino *et al.* 2012), specimens from both species could have been confused. The authors presumably determined specimens based on shell morphology, although details on how identifications were carried out were not provided.

Even though Varela *et al.* (2012) found differentiation among northern and southern ‘*E. siliqua*’ populations, Arias *et al.* (2011) and Arias-Pérez *et al.* (2012) found a Portuguese population clustering with an Irish population. In the morphometric study by Rufino *et al.* (2012), they described three different ‘morphs’ based on shell morphometrics, one from an Irish population and two others from Portuguese populations. Taking our results into account, the differentiation that they found among populations might have been produced by a two-species scenario, rather than by population isolation. This can be tested by DNA barcoding some of the specimens analysed in those surveys.

Due to sampling limitations, it was out of the scope of this work to investigate the distribution ranges of species in detail. Nonetheless, because we found *E. siliqua* in Borkum reef, Central North Sea, and Cedeira,

and *E. minor* in Bandol, La Capte, Lira, and Ría de Vigo, one might think that *E. minor* is distributed from the central Galician 'rias' to the Mediterranean, whereas *E. siliqua* has a more northern distribution. However, the distribution range of *E. minor* north of Galicia and northward to Scotland was confirmed by Cosel (2009) based on shell morphology of his own samplings, as were different biotopes and the sympatric occurrence of *E. siliqua* in the same areas. Only in the southern part of Brittany Peninsula, were a few morphological intergrades of *E. minor* x *E. siliqua* found (Cosel 2009).

#### DNA barcodes for Atlantic *Ensis*

The ideal DNA barcode should meet the following conditions: (1) suitable primer pairs should be available, (2) sequence-types should not be shared even by specimens from closely related species, (3) alignments should be straightforward, (4) intragenomic variation should be low to avoid cloning of PCR products before sequencing, and (5) sequences obtained from individuals from the same species should cluster together (and apart from sequences obtained from other species) in a phylogenetic tree or network.

In Atlantic *Ensis*, condition (1) was met by all genomic regions considered, even though in the case of COI, several primer pairs were used. Condition (2) was not met by 5.8S nor by 18S regions so they were not considered in subsequent phylogenetic analyses. Even though 5.8S is not widely used in phylogenetics due to its short length and scarce polymorphism, the 18S region is a popular phylogenetic marker. In *Ensis*, 18S was useful in differentiating the European/African clade from the American clade, but this region failed to differentiate among closely related species, in agreement with Tang *et al.* (2012). Despite the fact that the 16S region did not meet condition (2), it was a useful marker for species delimitation. Conditions (3) and (4) were not met by ITS1 nor by ITS2, as already reported by Vierna *et al.* (2010), but were suitable for species delimitation. Finally, even though condition (5) depends on the particular phylogenetic analyses carried out, it was met by COI and ITS1 in most cases (very clearly in phylogenetic networks, but not as clear in some phylogenetic trees, especially for *E. directus* and *E. terranovensis* sequences). Therefore, we suggest using COI as the main DNA barcode in Atlantic *Ensis* species.

Even though some authors have attempted to establish molecular identification protocols for some *Ensis* using PCR and PCR-RFLPs (Fernández-Tajes & Méndez 2007; Freire *et al.* 2008; Fernández-Tajes *et al.* 2010), because species boundaries were still unclear at that time, the proposed methodologies might give misleading results in some cases. Moreover, DNA barcoding is a much more informative (and therefore reliable) approach compared with PCR/PCR-RFLP based methods, even though the cost is somewhat higher due to the additional sequencing step.

#### Conclusion and future directions

In this work, using mitochondrial and nuclear DNA sequence data, we have clarified the species limits within Atlantic *Ensis*. In addition, we show that DNA barcoding is capable of identifying *Ensis* specimens to the species level and suggest taking advantage of this methodology for reliable and inexpensive identifications of specimens.

Finally, we encourage regional authorities to study the species composition of razor shell communities and establish specific conservation and managing plans, at least in the areas where *Ensis* spp. are an economic resource.

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