

Genetic diversity in fishery-exploited populations of the banded murex (*Hexaplex trunculus*) from the southern Iberian Peninsula^{*†}

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Abstract

This study presents an analysis of the genetic diversity in *Hexaplex trunculus* from three localities of the southern Iberian Peninsula (Ria Formosa and Ria de Alvor in Portugal, and Río Piedras in Spain) using three mitochondrial (12S rRNA, 16S rRNA and cytochrome c oxidase subunit I, COI) and one nuclear (5S rRNA and its non-transcribed spacer, NTS) genes. Restriction digestions of 12S and 16S rRNA genes using four endonucleases were also performed. Low genetic diversity was observed for the four genes studied. Genetic distances ranged from 0 to 0.004 for the mitochondrial genes and these values were slightly higher for the 5S and the NTS. The level of polymorphism within populations, π , was from 0 to 0.0039 (for mitochondrial genes), 0.0111–0.0333 (for 5S rRNA), and from 0.2220 to 0.5079 (for NTS). Furthermore, RFLPs analyses from 12S and 16S rRNA genes showed these localities to be monomorphic. The low genetic variability within populations suggests small population size, and a possible bottleneck due to multiple causes, such as overexploitation, the type of larval development (intracapsular embryos) and/or the peripheral location of the sampled localities considering the geographical distribution of the species. Because these populations show such high genetic similarity, Ria de Alvor and Río Piedras could be potentially used for stock enhancement of the Ria Formosa.

Keywords: Banded murex; Genetic diversity; *Hexaplex trunculus*; Iberian Peninsula; Mitochondrial DNA; 5S rRNA.

Introduction

The family Muricidae constitutes a diverse and important component of marine communities (Vokes, 1996), comprising 1150 (Vokes, 1996) to 1300 species distributed worldwide (Houart, 2001). The banded murex, *Hexaplex trunculus* (Linnaeus, 1758), is found in the Mediterranean Sea, whereas in the adjacent Atlantic Ocean it is mainly present from the Portuguese coast southward to Morocco and to the Madeira and Canary Archipelagos (Poppe and Goto, 1991, Macedo et al., 1999, Houart, 2001). Recently, a first record of this species in NW Spain (O Grove - Galicia), probably due to an accidental introduction from the importation of oysters (Quintas et al., 2005), was followed by the establishment of an apparently stable population of *H. trunculus*, as confirmed by the catch of hundreds of specimens in this area (Rolán and Bañon-Díaz, 2007). This species occurs in the inter-tidal and infra-littoral zones reaching up to 100–120 metres depth (Poppe and Goto, 1991, Muzavor and Morenito, 1999), although they are more frequent at 0.3–30 metres depth (Houart, 2001). It inhabits both hard and soft substrates, from rocky shores (Houart, 2001) to sandy, sandy-muddy and preferentially muddy bottoms (Poppe and Goto, 1991, Macedo et al., 1999, Muzavor and Morenito, 1999) and also occurs in areas enriched with organic matter and in the vicinity of ports (Šimunović, 1995).

The banded murex is regularly or occasionally fished for human consumption in several countries in the Mediterranean Sea (Poppe and Goto, 1991, Houart, 2001), namely in Italy, Cyprus, Turkey (Gaillard, 1987), Croatia (Peharda and Morton, 2006) and Tunisia (Gharsallah et al., 2004). In Iberian waters, *H. trunculus* is commercially exploited both in southern Spain, along the Mediterranean and Atlantic coasts of Andalucía (Anon., 2001, Tirado et al., 2002), and in southern Portugal, mainly inside the coastal lagoons located along the Algarve coast (principally in Ria Formosa, but also in Ria de Alvor) (Muzavor and Morenito, 1999, Vasconcelos et al., 2008).

In the Ria Formosa lagoon, *H. trunculus* is exploited in locally important artisanal fishery. This activity is traditionally undertaken both by manual harvesting during low tide and with an illegal fishing gear locally known as “wallet-line” (Vasconcelos et al., 2008). Additionally, the massive aggregations of females during collective spawning (that can comprise hundreds of individuals) are subjected to hand collecting during low tide, due to the easy capture and high fishing yield (Muzavor and Morenito, 1999, Vasconcelos et al., 2004). More recently, scuba divers operating illegally inside the lagoon have also been collecting considerable quantities of this species (P. Vasconcelos, personal observation). Altogether, these fishing and harvesting activities are regularly or occasionally carried out by tens of professional fishermen and recreational users, but the appraisal of the overall number of persons involved is almost impossible due to the illegal character of some of these activities. Therefore, no reliable official statistics are available on the catches of *H. trunculus* in the Ria Formosa lagoon, because much of the trade is conducted through a parallel economy in which sales are not declared. Probably due to over-fishing, detrimental collecting practices and disregard to the minimum landing size established for this species (50 mm shell length), the abundance of *H. trunculus* in the Ria Formosa lagoon has decreased noticeably over recent years. Furthermore, because *H. trunculus* deposits collective spawns and has direct intracapsular development (i.e., lacking a planktonic larval stage), it is also highly restricted in mobility which limits its ability to colonise new areas (Vasconcelos et al., 2004). Altogether, the reduced catches and the increasing commercial value of the banded murex (reaching values of 10–15€/kg for first sale) have inclusively raised some expectations about the potential of *H. trunculus* as a new species for molluscan aquaculture, both for commercial production and for stock enhancement in the Ria Formosa lagoon.

The availability of information on the species genetics is essential for establishing successful stock enhancement or restocking operations (e.g., Bell et al., 2005, Gaffney, 2006, Wenne et al., 2007), which could ultimately contribute for a long-term sustainability of the fishing resource. This work aimed to characterise genetic variations in *H. trunculus* in the context of its ecology and exploitation. A fishery-exploited population from the Ria Formosa (southern Portugal) was compared with two adjacent populations

(Ria de Alvor in southern Portugal and Río Piedras in Huelva, southern Spain), which could potentially constitute source-populations for stock enhancement of the autochthonous population from the Ria Formosa.

Materials and methods

Sampling

Commercial samples of *H. trunculus* were obtained during 2006 from the autochthonous population in Ria Formosa (Algarve coast in southern Portugal) and from two potential source-populations: Ria de Alvor (Algarve coast in southern Portugal, approximately 30 nautical miles westward) and Río Piedras - El Rompido (Huelva, southern Spain, approximately 20 nautical miles eastward) (Table 1).

In the laboratory, individuals were measured (shell length, SL) using a digital calliper (precision 0.01 mm) and weighed (total weight, TW) on a top-loading digital balance (precision 0.01 g). Routine sexual identification was made in sacrificed specimens by breaking the shells in a bench vice to allow the removal of the soft parts and the exposition of the mantle cavity. Due to the incidence of the imposex phenomenon (development of male secondary sexual characters into females) in local populations of this species (Vasconcelos et al., 2006) males were identified by the presence of a penis and lack of capsule gland, while females were identified by the presence of a vagina and capsule gland.

Subsequently, a small portion (roughly 1 cm²) of foot tissue was cut using dissecting instruments and preserved in 70% ethanol until further processing. To prevent the contamination of the tissue samples, particular attention was taken to avoid cutting the muscle tissue in close proximity to the proboscis (containing digestive enzymes) and to the external surface of the foot (with dark pigmentation).

DNA extraction, amplification and sequencing

Total genomic DNA was extracted as in Winnepennickx et al. (1993) from 20 mg of ethanol-preserved (70% ethanol) muscle tissue. The tissue was homogenised and incubated in CTAB buffer (2% CTAB, 1.4 M NaCl, 20 mM EDTA, 100 mM Tris-HCl pH 8.0). The DNA was purified with phenol/chloroform/isoamylalcohol (25:24:1) and precipitated using two-thirds volume of isopropanol, overnight at room temperature. After DNA centrifugation, the pellet was washed with 90% ethanol and air-dried. Finally, the DNA was dissolved in 1xTE (pH 7.5) to a final concentration of approximately 10 ng/μL.

The 12S rRNA, 16S rRNA and cytochrome c oxidase I (COI) genes were amplified using universal primers. The primers used for 12S rRNA were 12SA-5' and 12SB-3' (Kocher et al., 1989), for 16S rRNA were 16Sar and 16Sbr (Palumbi, 1996), and for COI primers LCO1490 and HCO2198 (Folmer et al., 1994) were used. For nuclear ribosomal 5S gene, the primers were 5S-Univ-F: 5'- accggtgtttcaacgtgat-3' and 5S-Univ-R: 5'- cgtccgatcaccgaagtaa-3'. All PCR amplifications were performed in 25 μL of a solution containing 1 ng/μL DNA, 10 mM Tris-HCl, 50 mM KCl pH 8.3, 1.5 mM MgCl₂, 2.5 mM dNTPs, 1 U of Taq DNA polymerase, and 1 μM of each primer. The PCR profile for 12 S and 16S rRNA consisted of one initial denaturation cycle of 3 min at 94°C, followed by 35 cycles at 94°C for 20 s, 50°C in the case of the 12S rRNA and 51°C for the 16S rRNA for 20 s, and 72°C for 45 s. A final extension was carried out at 72°C for 5 min. For COI, the PCR profile consisted of one initial denaturation cycle of 3 min at 94°C, followed by 35 cycles at 94°C for 1 min, 40°C for 1 min, and 72°C for 1.5 min and a final extension at 72°C for 2 min. The PCR amplification profile for 5 S rRNA consisted of one initial denaturation cycle of 3 min at 94°C, followed by 35 amplification cycles of 45 s at 94°C, 45 s of annealing temperature at 50°C, 1 min at 72°C, and a final extension cycle at 72°C for 10 min. The PCR products were purified with ExoSAP-IT (Amersham

Pharmacia Biotech) following the manufacturer's instructions. Purified PCR products were cloned with TA cloning kit (Invitrogen). Then, the plasmid was purified using the QIAprep®Miniprep kit (Qiagen) based on alkaline lysis of bacterial cells followed by adsorption of DNA onto silica in the presence of high salt. Sequence reactions were carried out in a capillary DNA sequencer (CEQ™8000 Genetic Analysis System) in both directions. The sequences were deposited in GenBank (Accession Nos. listed in Table 1).

Alignment and sequence analyses

Homologue sequences were obtained from GenBank using BLAST program. Alignments for 12S rRNA, 16S rRNA and 5S rRNA were performed with CLUSTAL X (Thompson et al., 1997) using default parameter gap opening 15.0 and gap extension 6.66. To determine the reliability of the data, we tested different orders of sequence input, obtaining identical results. For 16S rRNA we did not detect the hypervariable region corresponding to the loop of domain V, and for this reason, all the sequences obtained were included in the analysis. The alignment of COI nucleotide sequences was constructed on the basis of the translated amino acid sequences using the programs BIOEDIT (Hall, 1999) and CLUSTAL X (Thompson et al., 1997).

Because of the skew in AT content, genetic distances (d) and transition/transversion ratios (R) were calculated according to Tamura's three-parameter model (Tamura, 1992) for the mitochondrial regions, and Kimura two-parameter model (Kimura, 1980) for the 5S rRNA gene. These analyses were performed using the software package MEGA 3.1 (Kumar et al., 2004). Maximum parsimony (MP, heuristic searches) analyses were conducted using equal character weighting, 100 random stepwise addition and tree bisection-reconnection (TBR) branch-swapping. Inferred sequence gaps were considered as missing data. Branch support levels were estimated with bootstrapping (Felsenstein, 1985) (1000 replications, 10 random additions each) using PAUP*. A maximum likelihood (ML) analysis was also performed on the combined matrix under the HKY model (Hasegawa et al., 1985).

Genetic diversity within populations was measured by calculating the nucleotide diversity (π , the average number of nucleotide differences per site between two sequences) (Nei, 1987) using DnaSp software, version 4.0 (Rozas et al., 2003). To test the presence or absence of STRs (Simple Tandem Repeats), we used the Tandem Repeats Finder (TRF) program, version 2.02 (Benson, 1999).

PCR-RFLPs

The amplifications of the 12S and 16S rRNA genes were used to develop PCR-RFLPs. Different endonucleases, *Alu I*, *Hae III*, *Dra I* and *Apa I*, were selected. Restriction reactions were performed in a 20 μ L volume containing 5 μ L of PCR product, 1X reaction buffer and 3 units of endonuclease and incubated at the appropriate temperature overnight. Resulting fragments were resolved after electrophoresis on 2.5% agarose gels in 1X TAE (40 mM Tris-acetate, 1 mM EDTA pH 8.0).

Table 1. Sampling locations and accession numbers of the *Hexaplex trunculus* specimens analysed in this study

Population	Location	Gene							
		12S		16S		COI		5S rRNA	
		clone	ac. no.	clone	ac. no.	clone	ac. no.	clone	ac. no.
Ria Formosa	37°08' N	Formosa-01	AM712298	Formosa-01	AM712596	Formosa-01	AM712604	Formosa-11	AM920312
	7°36' W	Formosa-02	AM712299	Formosa-02	AM712597	Formosa-02	AM712605	Formosa-12	AM920313
		Formosa-03	AM712300		Formosa-03	AM712606	Formosa-21	AM920314	
		Formosa-04	AM712301		Formosa-04	AM712607	Formosa-22	AM920315	
							Formosa-31	AM920316	
							Formosa-32	AM920317	
Ria de Alvor	37°07' N	Alvor-01	AM712302	Alvor-01	AM712598	Alvor-01	AM712608	Alvor-11	AM920318
	8°36' W	Alvor-02	AM712303	Alvor-02	AM712599	Alvor-02	AM712609	Alvor-21	AM920319
		Alvor-03	AM712304	Alvor-03	AM712600	Alvor-03	AM712610		
		Alvor-04	AM712305	Alvor-04	AM712601	Alvor-04	AM712611		
Río Piedras	37°12' N	Piedras-01	AM712306	Piedras-01	AM712602	Piedras-01	AM712612	Piedras-11	AM920320
	7°03' W	Piedras-02	AM712307	Piedras-02	AM712603	Piedras-02	AM712613	Piedras-12	AM920321
								Piedras-21	AM920322
							Piedras-22	AM920323	

Results

A total of 102 specimens of *H. trunculus* (56 males and 46 females) with broad size ranges (SL: 39.29 to 89.10 mm; TW: 5.69 to 87.50 g) were analysed: 40 individuals from Ria Formosa (SL = 62.00 ± 5.01 mm; TW = 22.62 ± 5.58 g), 40 individuals from Ria de Alvor (SL = 49.14 ± 5.29 mm; TW = 11.55 ± 4.17 g) and 22 individuals from Río Piedras (SL = 71.29 ± 11.99 mm; TW = 43.12 ± 20.73 g).

The partial amplification of 12S rRNA showed a length of 370 bp, and an AT content higher than 66%, as expected for the 12S rRNA region III. The alignment of nucleotide clones (data not shown) revealed a low number of nucleotide changes among them. Genetic distance and nucleotide diversity values among clones within populations (Table 2) were very low, being zero for Ria Formosa. Likewise, the values of genetic distance among localities (Table 3) were also low (0.001–0.007).

Table 2. Mean pairwise distances (d) and nucleotide diversity (π) within populations

Population	12S		16S		COI		5S		NTS	
	d	π	d	π	d	π	d	π	d	π
Ria Formosa	0	0	0.004	0.0039	0	0.0010	0.0258	0.0250	0.647	0.4204
Ria de Alvor	0.001	0.0014	0	0	0	0	0.0341	0.0333	0.974	0.5079
Río Piedras	0.003	0.0027	0	0	0	0	0.0112	0.0111	0.508	0.2220

Within group means are arithmetic means of all individual pairwise distances between specimens within a group.

The internal region of 16S rRNA was 513 bp in length and the AT content was higher than 63%. Again, alignment among the different clones displayed low variability (data not shown). Genetic distance values among clones within populations (Table 2) and among populations (Table 3) were lower than for the 12S rRNA gene. This was also supported by the low value of nucleotide diversity (Table 2), which was zero for Ria de Alvor and Río Piedras. Ria Formosa displayed higher values than those for the 12S rRNA gene.

Table 3. Distance between populations average, below diagonal for 12S gene, above diagonal for 16S

	Ria Formosa	Ria de Alvor	Río Piedras
Ria Formosa	-	0	0.002
Ria de Alvor	0.001	-	0.002
Río Piedras	0.007	0.007	-

The average distance is the arithmetic mean of all pairwise distances between two groups in the inter-population comparisons.

The E1 region corresponding to the COI gene was 657 bp in length and showed AT content >62%. The translation of the nucleotide sequences into amino acid sequences, following the *Drosophila yakuba* mitochondrial genetic code (Clary and Wolostelholme, 1985), allowed us to obtain open reading frames in all the clones, not finding stop codons in the middle of the sequence. The length of the amino acid chains was of 219 amino acids. The percentages of each amino acid were calculated, this region being rich in hydrophobic amino acids of which leucine was the most abundant. Alignment of COI nucleotide sequences revealed that all clones were identical (Table 2), only showing one variable site out of 657 nucleotides. Genetic distance and nucleotide diversity values (Table 2) were even lower than for the 12S rRNA and 16S rRNA genes.

The PCR-RFLPs from 12S and 16S rRNA were developed to analyse genetic polymorphism in 30 individuals for each of the Portuguese localities and 20 for the Río Piedras locality. After digestion with the four restriction endonucleases, *Alu I*, *Hae III*, *Dra I* and *Apa I*, all the samples were monomorphic, displaying only one haplotype.

Genetic variability was also analysed by cloning and sequencing the 5S rRNA gene. The cloned fragments showed high variability in the size of the insert, ranging from ~250 bp to 1000 bp (Table 4). Different clones were randomly selected to study the transcribed (5S) and non-transcribed (NTS) regions. For the 5S coding region, all clones showed 120 bp length and a GC content ranging from 53.3 to 58.3 (Table 4), displaying higher genetic variability than the mitochondrial genes. Genetic distances among clones within each locality (Table 2) revealed Río Piedras as the lowest variable (lowest *d* and π values), while Ria de Alvor displayed the highest genetic variability. Furthermore, genetic distances among populations (Table 5) showed Río Piedras as being more distant from Ria de Alvor and Ria Formosa. The maximum parsimony phylogenetic analysis of the 5S region (Fig. 1) showed one cluster containing all the clones of *H. trunculus* and the rest of the gastropod species included in this analysis and available in the Genbank (*Arion rufus*, *Aplysia kurodai* and *Helix pomatia*). The topology showed the monophyly of *H. trunculus* with regard to the rest of gastropods and grouped the clones from Río Piedras separately from those from Ria de Alvor and Ria Formosa.

Table 4. Lengths and guanine-cytosine content of coding and spacer regions of 5S rRNA, and microsatellite sequences with positions

Clone	5S rRNA + NTS		5S rRNA		NTS		Microsatellite	Position
	Length	%CG	Length	%CG	Length	%CG		
Formosa-11	458	38.0	120	54.2	338	32.3	(TATT)8	164–195
Formosa-12	454	37.5	120	53.3	334	31.5		
Formosa-21	426	46.0	120	56.7	306	41.8		
Formosa-22	422	45.3	120	54.1	302	42.0		
Formosa-31	507	38.3	120	55.0	387	32.8	(ATC)15	355–398
Formosa-32	956	40.6	120	55.0	836	38.1	(CAA)35	348–449
							(CAG)10	454–482
							(GCCT)11	523–567
							(CT)17	757–789
							(TGATC)9	817–862
Alvor-11	258	36.8	120	55.9	138	22.5		
Alvor-21	428	46.0	120	53.3	308	42.2		
Piedras-11	819	36.3	120	57.5	699	33.2	(TCA)8	446–468
Piedras-12	809	36.6	120	57.5	689	32.9	(TCA)8	405–427
Piedras-21	959	40.6	120	58.3	839	38.5	(CAA)30	362–446
							(CAG)10	451–479
							(GCCT)11	520–564
							(CT)15	756–784
							(TGATC)9	812–857
Piedras-22	962	41.1	120	58.3	842	38.6	(CAA)35	355–447
							(CAG)10	452–480
							(GCCT)12	521–569
							(CT)15	759–787
							(TGATC)9	815–860

Table 5. Distance between populations average, below diagonal for 5S gene, above diagonal for NTS

	Ría Formosa	Ría de Alvor	Río Piedras
Ría Formosa	-	0.623	0.583
Ría de Alvor	0.027	-	0.760
Río Piedras	0.050	0.052	-

The average distance is the arithmetic mean of all pairwise distances between two groups in the inter-population comparisons.

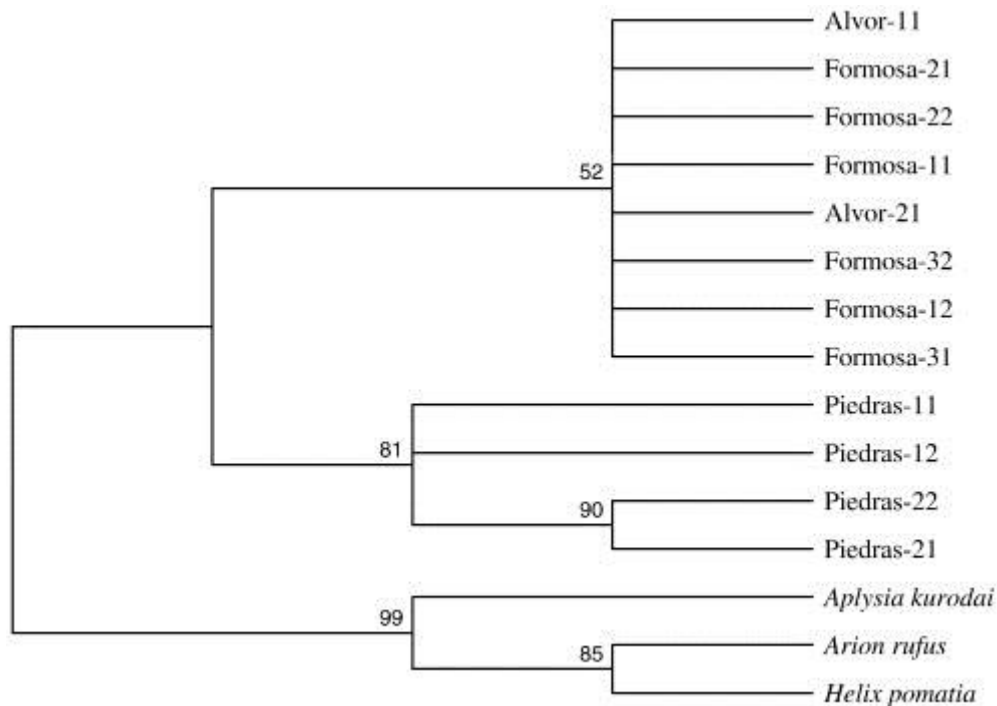


Figure 1. The single most-parsimonious tree (1000 bootstrap replicates; L = 34, CI = 0.8056, RI = 0.8444, RC = 0.6802) obtained from the maximum parsimony analysis of 5S transcribed sequences in *Hexaplex trunculus*. Maximum likelihood analysis produced a largely congruent topology (HKY model, $-\ln L = 353.85784$). Sequences from the three gastropod species deposited in Genbank (*Aplysia kurodai* X04305, *Arion rufus* J01888, and *Helix pomatia* J01889) were incorporated into the tree. Bootstrap values over 50% are shown above branches. (For more details of the populations, clones and accession numbers, see [Table 1](#)).

Regarding the NTS, the lengths ranged from 138 to 842 nucleotides and the GC content from 31.5 to 42.2% (Table 4). This region showed higher variability than the 5S coding region, as observed from the analysis of the genetic distances within populations (Table 2) and among the different populations (Table 5). Sequence alignments of clones revealed the existence of four groups: i) Formosa-21, Formosa-22 and Alvor-21, ii) Piedras-11, Piedras-12 and Formosa-31, iii) Formosa-11, Formosa-12, and Alvor-11, and iv) Formosa-32, Piedras-21, and Piedras-22. Ria de Alvor was the locality with the highest variability and genetic distance values. This grouping is observed in the maximum parsimony tree (Fig. 2), which shows that types II and III are more closely linked to each other than to the other two NTS types.

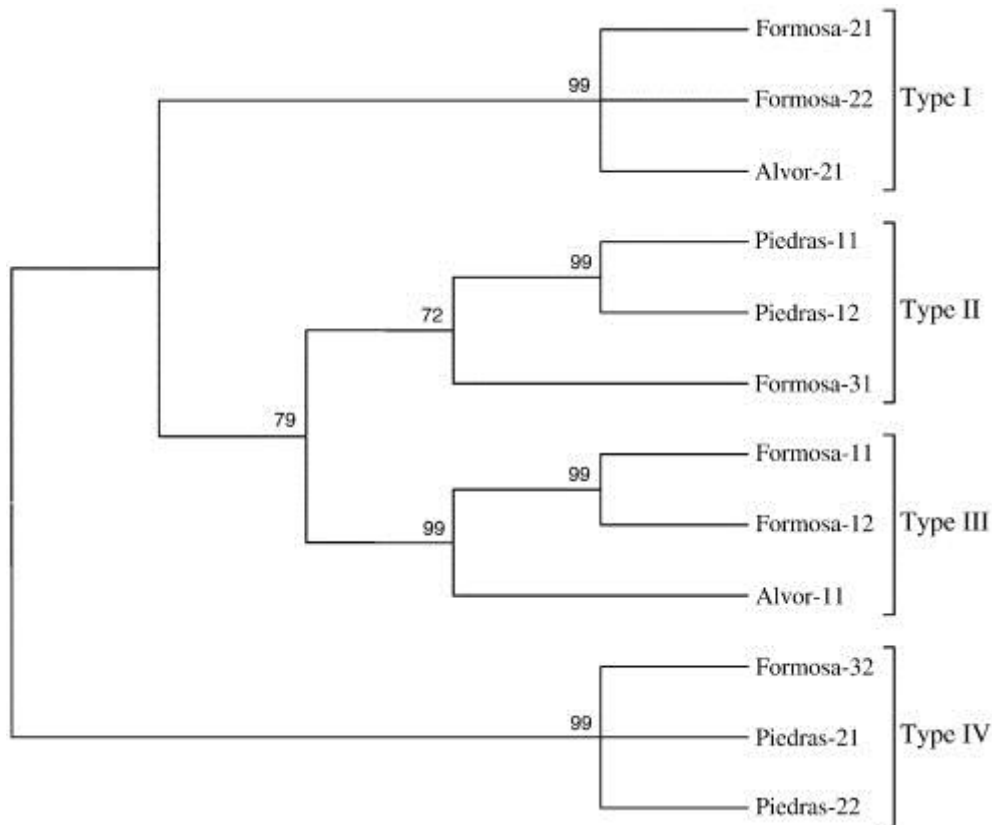


Figure 2. The single most-parsimonious tree (1000 bootstrap replicates; $L = 275$, $CI = 0.8667$, $RI = 0.8906$, $RC = 0.7719$) obtained from the maximum parsimony analysis of 5S non-transcribed sequences in *Hexaplex trunculus*. Maximum likelihood analysis produced a largely congruent topology (HKY model, $-\ln L = 3637.83163$). Bootstrap values over 50% are shown above branches. (For more details of the populations, clones and accession numbers, see [Table 1](#)).

The analysis of NTS regions using the TRF program allowed us to detect the existence of microsatellite sequences in some of the clones studied (Table 4). Using a penalty of 2 for the match, 7 for the mismatch, and 7 for the presence of indels, and using alignment score values of 50 and maximum period size of 500, we obtained five different types of microsatellites in the clones of NTS type IV (Formosa-32, Piedras-21, Piedras-22). Repetitive units are basically trinucleotides, with the triplet CAA repeated at least 30 times.

Discussion

Mitochondrial DNA (mtDNA) is a useful marker to investigate genetic diversity and it has been used to analyse polymorphism and define stocks in many invertebrate species (Boudry et al., 1998, O'Foighil et al., 1998, Arnaud-Haond et al., 2003). However, mtDNA must be used in conjunction with nuclear markers to accurately identify populations for conservation (Moritz, 1994), because when considering a low effective number of genes (Birky et al., 1989), mtDNA can diverge while nuclear genes do not. The value of combining mtDNA with nuclear markers has been demonstrated for a great variety of some intensively managed species (see Moritz, 1994 for review). In some cases, low mtDNA diversity is correlated with low nuclear gene diversity. This suggests that it is inappropriate to set conservation or management priorities on the basis of within-population mtDNA diversity (Moritz, 1994). Thus, in this study, we combined the analysis of three mitochondrial genes (12S rRNA, 16S rRNA and cytochrome c oxidase subunit I) and the

nuclear 5S rRNA gene to investigate the genetic diversity in three populations of the banded murex (*H. trunculus*) subjected to different levels of fishery exploitation (intensively exploited in Ria Formosa and moderately exploited in Ria de Alvor and Río Piedras).

A considerable number of works have revealed high levels of both inter- and intra-specific variability in 12S rRNA, 16S rRNA and COI genes in marine molluscs and also from RFLPs analyses. However, our results showed that the *H. trunculus* populations examined have low genetic variability for the four genes analysed. For the mitochondrial genes, in the analyses of the 12S and 16S rRNAs the sequence length and the AT content were similar to those found in other gastropods (Marko and Vermeij, 1999, Holznagel and Lydeard, 2000, Kirkendale and Meyer, 2004), while nucleotide diversity values, π , were lower than those obtained from 16S rRNA for *Biomphalaria pfeifferi* (Angers et al., 2003) (Table 6). Furthermore, the analysis from 12S and 16S RFLPs showed that they were monomorphic, confirming the low genetic differentiation among and within populations. For COI, there was also low variability for nucleotide diversity, in this case being lower than that detected in the genera *Adriohydrobia* (Wilke and Falniowski, 2001), *Cerithidea* (Kojima et al., 2006), and in the species *Nucella lamellosa* and *N. ostrina* (Marko, 2004), *Batillaria cumingi* (Kojima et al., 2004), *Buccinum tsubai* (Iguchi et al., 2007) and *Nassarius reticulatus* (Couceiro et al., 2007) (Table 6).

Table 6. Nucleotide diversity (π values) for 16S and COI mitochondrial genes in different gastropod species

Species	Nucleotide diversity		Reference
	16S	COI	
<i>Biomphalaria pfeifferi</i>	0.0045	-	Angers et al. (2003)
<i>Adriohydrobia gagatinella</i>	-	0.0051	Wilke and Falniowski (2001)
<i>Batillaria cumingi</i>	-	0.012	Kojima et al. (2004)
<i>Buccinum tsubai</i>	-	0.00055 to 0.00277	Iguchi et al. (2007)
<i>Cerithidea cingulata</i>	-	0.0004 to 0.0119	Kojima et al. (2006)
<i>Cerithidea djadjariensis</i>	-	0.0000 to 0.0088	Kojima et al. (2006)
<i>Cerithidea largillierii</i>	-	0.0059 to 0.0068	Kojima et al. (2006)
<i>Cerithidea rhizophorarum</i>	-	0.0007 to 0.0054	Kojima et al. (2006)
<i>Hexaplex trunculus</i>	0 to 0.0039	0.0000 to 0.0010	Present work
<i>Nassarius reticulatus</i>	-	0.0023 to 0.0041	Couceiro et al. (2007)
<i>Nucella lamellosa</i>	-	0.005	Marko (2004)
<i>Nucella ostrina</i>	-	0.002	Marko (2004)

However, the analysis of the nuclear 5S rRNA revealed higher values of π than for those obtained from the mitochondrial ones. This gene displayed guanine-cytosine richness, with a percentage surpassing 52% for the coding region (Table 4), similar to that obtained for *A. kurodai* and *A. rufus* (Komiya et al., 1986), and *H. pomatia* (Fang et al., 1982). Values of nucleotide diversity obtained from the 5S region were higher than those obtained from the mitochondrial genes, possibly due to the high conservation degree of this gene through species (Drouin and Moniz de Sá, 1995). Regarding NTS, this is the first time that this region is used to perform an analysis of genetic diversity in a gastropod. The results obtained revealed that the guanine-cytosine content was 22.5% – 42.2% and the length ranged from 138 bp to 839 bp, which is due to the absence/ presence of microsatellite sequences.

For most invertebrate fisheries, the key to understanding the population structure lies, in part, in knowing the mobility of the species and their larvae (Thorpe et al., 2000). Most of them have reduced mobility or are sessile, thus the only mode of dispersion is through the larval stages (or ultimately through accidental introductions of larvae, juveniles or adults). In the case of the gastropods, the scale of genetic differentiation between populations or stocks (or localities) appears to be related to larval type (Thorpe et al., 2000). The banded murex, *H. trunculus*, is a direct-developer species (lacking planktonic larva), and both juveniles and

adults are highly restricted in mobility (Vasconcelos et al., 2004). Furthermore, unintended transport by the currents is most unlikely, because they normally live adhered to the bottom, and occasionally even burrow shallowly into soft sediments to avoid both the unfavourably low and high water temperatures (Spanier, 1981, Spanier, 1986, Spanier and Karmon, 1987), being often observed partially buried in the sand (Spanier, 1986, Rilov et al., 2004). Altogether, this could explain the low genetic diversity and differentiation found in the banded murex. In this sense, the analysis of different mitochondrial genes in gastropod species with pelagic and nonpelagic larvae has revealed a significant reduction in nucleotide diversity in species with nonpelagic larvae (Grant and da Silva-Tatley, 1997, Foltz, 2003). This is consistent with the suggestion that the nonpelagic lineages have reduced effective population sizes, which is correlated with the lower levels of gene flow among populations (Foltz, 2003). Furthermore, if the population size is small, the low values of genetic diversity could be also caused by genetic drift. The reduced mobility of *H. trunculus* crawling hatchlings (benthic early post-metamorphic individuals) implies low dispersal (and hence, small population size), therefore stocks are much more vulnerable to the multiple effects of natural and anthropogenic stressors than those species which have planktonic larvae with high dispersal.

The relationships between larval type and gene flow are complex, and the level of gene flow in a species probably depends on other life history factors in addition to the mode of reproduction, such as the adult body size, reduced fecundity, or ecological conditions (Wilke and Davis, 2000). Taking this into account, another cause for the low genetic variability in *H. trunculus* could be the eventual existence of a recent bottleneck and an event of decreasing population size, possibly due to overfishing, although this would be more likely to have occurred in the case of the population from Ria Formosa. This, together with detrimental collecting practices (hand collecting in communal spawns during low tide and illegal catches through scuba-diving) might have dramatically worsened the status of genetic diversity in this locality.

However, overfishing solely does not explain the low genetic variability detected in the populations from Ria de Alvor and Río Piedras, both subjected to lower levels of fishery exploitation (only moderately exploited, when compared to the intensively exploited population from Ria Formosa), and thus other factors must be considered, namely the species distributional range and the geographical location of the localities sampled. Indeed, *H. trunculus* is essentially a Mediterranean species, and therefore Río Piedras, Ria Formosa and Ria de Alvor are peripheral localities in its geographic distribution, constituting the westernmost populations along the southern Iberian Peninsula (moreover, the distances between them are relatively short). It is known that the distribution of most species is characterised by central dense populations and peripheral marginal populations, and that this distribution is regulated by biotic and/or abiotic factors that make peripheral habitats less suitable for the maintenance of populations (Brown et al., 1995, Brown et al., 1996). Demographic characteristics of these margins together with spatial and temporal variations of the environment determine the exact range of the boundary (Brown et al., 1996). Usually, marginal or peripheral populations are fragmented in patches or areas that favour metapopulation processes (Holt and Keitt, 2000). Each population within a metapopulation is relatively independent of the rest, and may eventually be endangered or become extinct as a consequence of demographic stochasticity. Furthermore, individuals from marginal habitats are expected to have lower fitness and population growth than those from central areas of distribution (Holt and Keitt, 2000, Maurer and Taper, 2002), and subsequently, to have low or reduced genetic variability. This situation, in conjunction with human activities that lead to a reduction of the suitable habitats, may ultimately result in a reduction of the species' range (Faugeron et al., 2004).

In addition, all these assumptions should be integrated in a scenario of global climate change that will most probably aggravate the overall situation. Indeed, recent rapid climate change has resulted in modifications in the biogeographic ranges, abundance, and population structure of several species (including Lusitanian species) (Mieszkowska et al., 2005). Moreover, it is believed that species with small and/or isolated populations and low genetic variability (often indicated by recent bottlenecks in population numbers) are least likely to withstand impacts of climate change (Inkley et al., 2004), increasing their risk of extinction. In

this context, species vulnerable to genetic bottlenecks may benefit from conservation efforts that enhance the genetic diversity (Hellmann and Pineda-Krch, 2007), consequently increasing the population's ability to adapt to environmental changes (Ray, 2001, Bell and Okamura, 2005, Frankham, 2005).

In terms of the stock enhancement of *H. trunculus* from Ria Formosa, several genetic considerations must be taken into account, being one of the most important to ascertain the geographic distribution of genetic diversity. This must be investigated before performing a stock selection for restoration. To compare the distribution of genetic diversity between populations in central areas of distribution and peripheral populations (also including more distant locations such as northern Africa, Madeira and Canary Archipelagos), could provide qualitative indications about the genetic status of the banded murex. The analysis of genetic diversity and genetic differentiation of populations will supply fundamental data for conservation and stock enhancement, allowing the sustainable exploitation of the fishing resources and preserving biodiversity.

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