Population differentiation of the shore crab *Carcinus maenas* (Brachyura: Portunidae) on the southwest English coast based on genetic and morphometric analyses

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SUMMARY: *Carcinus maenas* has a planktonic larval phase which can potentially disperse over large distances. Consequently, larval transport is expected to play an important role in promoting gene flow and determining population structure. In the present study, population structuring on the southwest coast of England was analysed using molecular and morphometric approaches. Variation at eight microsatellite loci suggested that the individuals sampled within this region comprise a single genetic population and that gene flow among them is not restricted. Nevertheless, the *F*<sub>st</sub> values estimated across loci for all populations suggested that the Tamar population was significantly different from the Exe, Camel and Torridge populations. This differentiation is not explained by isolation by distance, and coastal hydrological events that are apparently influencing larval flux might be the cause of this pattern. Morphometric analysis was also performed. Analysis of carapace and chela shape variation using landmark-based geometric morphometrics revealed extensive morphological variability, as the multivariate analysis of variance showed significant morphometric differences among geographic groups for both sexes. Thus, the morphological differentiation found may be a plastic response to habitat-specific selection pressures.

Keywords: *Carcinus maenas*, microsatellites, population structure, geometric morphometrics, phenotypic plasticity, gene flow.

RESUMEN: Diferenciación poblacional del cangrejo *Carcinus maenas* (Brachyura: Portunidae) en la costa sudoeste de Inglaterra basada en análisis genéticos y morfométricos. – El cangrejo *Carcinus maenas* tiene una fase larvaria planctónica que potencialmente puede dispersarse a grandes distancias. Como consecuencia, es esperable que el transporte larvario tenga un papel importante en el flujo genético y en la determinación de la estructura de las poblaciones. En el presente estudio se ha analizado la estructura de las poblaciones de la costa sudoeste de Inglaterra, utilizando marcadores moleculares y morfométricos. La variación en ocho loci de microsatélites sugiere que los individuos muestreados en esta región constituyen una única población genética y que el flujo genético entre ellos no es limitado. Sin embargo, los valores de *F*<sub>st</sub> estimados para los ocho loci de todas las poblaciones muestreadas sugieren que la población de Tamar es significativamente diferente de las poblaciones de Exe, Camel y Torridge. Esta diferencia no se explica por el aislamiento por distancia. En contrapartida, el flujo genético parece estar influenciado por mecanismos hidrológicos costeros, que contribuyen así al patrón de diferenciación encontrado. También se realizó un análisis morfométrico, utilizando la técnica de la morfometría geométrica. Estos análisis, ejecutados según la forma del caparazón y de la pinza, revelaron una gran variabilidad morfológica, del mismo modo en que el análisis de la varianza multivariado reveló diferencias morfológicas significativas entre poblaciones, para ambos sexos. Así, la variación morfológica encontrada podría ser una respuesta plástica a presiones selectivas relacionadas con la especificidad del hábitat ocupado.

Palabras clave: *Carcinus maenas*, microsatélites, estructura de las poblaciones, morfometría geométrica, plasticidad fenotípica, flujo genético.
INTRODUCTION

Population genetic analyses can reveal important aspects of evolution and ecology of a species, such as natural selection, gene flow, genetic variation and differentiation. As an example, the amount of genetic variability can determine the adaptation to environmental changes, while gene flow allows connectivity between populations, preventing genetic erosion and inbreeding (Rousset, 2001). Furthermore, from the distribution of molecular genetic markers, important ecological traits, such as mating system and spatial population boundaries, can be inferred (Rousset, 2001). However, understanding these processes in marine species is particularly difficult because barriers to gene flow are far less obvious compared to continental species (Patarnello et al., 2007). Population genetic studies of marine invertebrate species have shown that high-dispersal potential due to planktonic larvae is often associated with mild genetic differentiation over large scales, which implies high levels of gene flow within populations (e.g. Reuschel and Schubart, 2006).

A number of recent studies have detected extensive phenotypic variability in shore crabs within relatively restricted geographical areas (e.g. Brian, 2005; Todd et al., 2006). Given that population divergence is prevented by gene flow, patterns of phenotypic variability are likely to reflect differences between the local environmental conditions, as was suggested by Brian et al. (2006). Phenotypic plasticity (i.e. induced changes resulting in different phenotypes in different environments) is today recognized as an important evolutionary mechanism that modulates inherited differences of individuals (Hollander et al., 2006) and can also be an important adaptive strategy in variable or changing environments (Schlichting and Pigliucci, 1998). In fact, if organisms encounter predictable environments, fixed development is expected, whereas in organisms that cannot predict their future environment, phenotypic plasticity would be optimal to increase local adaptation (Hollander et al., 2006). As an example, inducible defences are a ubiquitous form of plasticity that involves the production of chemicals, morphologies, or behaviours by prey species in response to predator cues (Trussell and Smith, 2000). These changes reduce prey vulnerability and examples include the diel vertical migration in marine zooplankton (Bollens et al., 1992) and shell shape and thickness in gastropods and bivalves (Trussell, 1996). Despite improved understanding of the cues inducing these changes and their immediate adaptive value (Schlichting and Pigliucci, 1998), our understanding of how this phenomenon contributes to broader temporal and spatial patterns of phenotypic variation remains poor.

The shore crab *Carcinus maenas* is common in the intertidal throughout Europe and has been well studied due to the ease with which it can be found, identified, sexed and measured. This highly adaptable crab has recently gained notoriety due to its globally invasive nature associated with drastic ecological and economic effects. Once established, it becomes the dominant intertidal crab in some areas (Yamada, 2001), affecting the abundance, size structure and defence response of native species (Roman and Palumbi, 2004). Thus, investigations of the population structure of this crab within its native range can provide insights into processes of invasions in other regions.

*Carcinus maenas* has a long planktonic larval phase (up to 50 days in the plankton; Tresher et al., 2003), and its offshore dispersal may account for considerable exchange of individuals between local populations (Peliz et al., 2007). According to Queiroga (1996), the zoal stages I and II are concentrated in the surface layer, but a gradual ontogenic displacement to deeper waters is observed from then on. Horizontally, there is a clear association of the first zoal with the sites where hatching occurs, while the older zoal stages are dispersed progressively offshore (Queiroga, 1996). Therefore, in accordance with the population genetics theory that suggests that marine animal species with long planktonic larvae have less genetic structure and higher connectivity than those with direct development (Palumbi, 2003), *C. maenas* would be expected to have low levels of population differentiation. However, previous studies of the shore crab *C. maenas* have suggested several different conclusions. Bulnheim and Bahns (1996) reported a slight geographic cline from north to south Europe in one allozyme locus. Bagley and Geller (1999) found no population structure in a microsatellite DNA study of Atlantic European crabs, and a recent study by Pascoal et al. (2009), also with microsatellite data, revealed very weak but significant structuring between populations on the Portuguese coast. A more extensive study by Roman and Palumbi (2004), with the mitochondrial cytochrome oxidase I gene, detected a slight population structure between the central North Sea and populations to the south, and a break between populations to the Faeroe Islands and Iceland and continental populations.

The shore crab *Carcinus maenas* (Linnaeus, 1758) is a typical inhabitant of the European coastline, which lives in the tidal and in the subtidal zone (Bulnheim and Bahns, 1996). In its native range, *C. maenas* has a distribution from northern Norway and Iceland to Mauritania, and it has successfully colonized Australia, Tasmania, South Africa, Japan and both coasts of North America (Roman and Palumbi, 2004 and references therein). On the English coast, *C. maenas* occurs throughout the coastline, the estuarine populations being linked by open coast populations. It has a complex life cycle comprising an exported planktonic larval phase that develops in shelf waters and takes four to six weeks to reach the megalopa stage (Tresher et al., 2003). Megalopae migrate back to the coast during spring tides, and transport is accomplished by selective tidal stream transport (Moksnes et al., 1998). Once they reach a suitable environment, the megalopae of *C. maenas* settle in a variety of intertidal and subtidal...
habitats, though they show a preference for those that are structurally complex (Paula et al., 2006).

Our study explored local patterns of population structure of onshore populations of *C. maenas* on the coastline of southwest England, using eight microsatellite loci as genetic markers. Morphological differences among these populations were assessed to show whether genetic and morphometric tools provide coinciding or contrasting results in population differentiation.

**MATERIALS AND METHODS**

**Sampling sites and procedures**

Adult shore crabs were collected using baited traps from the intertidal zone at nine estuarine sites on the southwest English coast during the Autumn of 2004 and 2005 (Fig. 1). The sites were the Axe, Exe, Teign, Tamar, Looe, Fowey, Hayle, Camel and Torridge estuaries. After collection, specimens were transported to the laboratory, where they were frozen at -20°C for preservation, and the fourth and fifth pereiopods were preserved in absolute ethanol. Specimens were separated by size and sex and labelled, and the carapaces and chelae were processed to eliminate those that were not suitable for the study (broken carapaces and regenerated and/or broken claws). Females of the species *C. maenas* with a carapace width of less than 1.5 cm and males with one of less than 2.5 cm were also removed from the morphometric analyses. These sizes correspond to the minimum sizes of sexual mature specimens (Neal and Pizzolla, 2008). Thus, this procedure eliminated most allometric growth variation. Morphometric analyses of the carapaces were made following Silva and Paula (2008), using the major chelae in males and right chelae in females.

**Genetic data analysis**

A total of 270 individuals were collected for genetic analyses. Forty-five specimens from each of the Exe, Teign, Tamar, Looe, Camel and Torridge populations were analysed. The Axe, Fowey and Hayle populations were not considered for the genetic analyses due to logistical constraints.

Total genomic DNA was extracted from muscle tissue of pereiopods using sequential phenol-chloroform extraction steps as described by Hillis et al. (1996). The DNA obtained was resuspended in low TE buffer (Tris-HCl 10 mM pH 8.0, EDTA 0.1 mM pH 8.0) and used as a template in polymerase chain reactions (PCR). Before PCR amplification, DNA concentration was estimated using the Thermo Scientific NanoDrop™ 1000 Spectrophotometer. Genetic diversity was screened at 8 microsatellite loci (Cma10EPA, Cma11EPA, Cma12EPA, Cma14EPA, developed by Tepolt et al. (2006); and SP107, SP229, SP280 and SP495 developed by Pascoal et al., 2009). Microsatellite polymorphism was detected using a fluorescent detection method, the forward primer for each locus being 5'-labelled with one of three fluorophores. The labelled PCR products from the 8 loci were then divided into three sets, taking into account the size of the fragments (Cma10EPA + Cma11EPA + SP107; Cma12EPA + Cma14EPA + SP229; SP280 + SP495). PCR amplifications were carried out in 15-μl reactions with 10 pmol of each primer set, 10-100 ng of genomic DNA, 1x PCR buffer, 1.3-1.8 mM MgCl2, 0.2 mM of each dNTP (Promega) and 0.5 U Taq DNA polymerase (Promega). Thermal cycling parameters were: initial denaturation at 94°C for 5 min followed by 26-35 cycles of denaturation at 94°C for 1 min, primer-specific annealing temperature (Table 1) for 1 min, extension at 72°C for 1 min and final extension at 72°C for 10 min. Following PCR amplification, the extension products were resolved on 1.5% agarose gels. PCR products were run on a Beckman Coulter™ sequencer. Quality of PCR products and allelic length were determined using the CEQ™ Genetic Analysis System software. Alleles were designated according to their sizes and the MICROVANES toolkit, a specific tool for Microsoft Excel, was used as a conversion program in order to obtain the GENEPOP, FSTAT and ARLEQUIN input data files.

Raw data were analysed with MICRO-CHECKER software (van Oosterhout et al., 2004) to check microsatellite data for null alleles and scoring errors. The observed and expected heterozygosity for each locus and population were calculated using GENEPOP 1.2 (Raymond and Rousset, 1995). The number of alleles forward primer for each locus being 5'-labelled with one of three fluorophores. The labelled PCR products from the 8 loci were then divided into three sets, taking into account the size of the fragments (Cma10EPA + Cma11EPA + SP107; Cma12EPA + Cma14EPA + SP229; SP280 + SP495). PCR amplifications were carried out in 15-μl reactions with 10 pmol of each primer set, 10-100 ng of genomic DNA, 1x PCR buffer, 1.3-1.8 mM MgCl2, 0.2 mM of each dNTP (Promega) and 0.5 U Taq DNA polymerase (Promega). Thermal cycling parameters were: initial denaturation at 94°C for 5 min followed by 26-35 cycles of denaturation at 94°C for 1 min, primer-specific annealing temperature (Table 1) for 1 min, extension at 72°C for 1 min and final extension at 72°C for 10 min. Following PCR amplification, the extension products were resolved on 1.5% agarose gels. PCR products were run on a Beckman Coulter™ sequencer. Quality of PCR products and allelic length were determined using the CEQ™ Genetic Analysis System software. Alleles were designated according to their sizes and the MICROVANES toolkit, a specific tool for Microsoft Excel, was used as a conversion program in order to obtain the GENEPOP, FSTAT and ARLEQUIN input data files.

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and the allelic richness for each locus and population were estimated using fstat 2.9.3.2 (Goudet, 2002). To test for deviations from Hardy-Weinberg expectations and linkage disequilibrium a probability test was performed, with unbiased \( p \)-values and associated standard errors (SE) estimated by the Markov Chain algorithm. Bonferroni correction of the \( p \)-values was used to correct multiple tests. As suggested by Raymond and Rousset (1995), number of batches (B) and number of iterations per batch (C) were increased to SE \( \geq 0.01 \) (B =400, C =4000). These estimations were achieved with genepop 1.2 (Raymond and Rousset, 1995). The Weir and Cockerman (1984) analogues of Wright’s \( F \)-statistics, \( F_{ST} \), and their significance per locus were estimated to assess levels of population differentiation over all loci, using fstat 2.9.3.2 (Goudet, 2002).

Isolation-by-distance was examined employing Mantel’s t-test (Mantel, 1967), computed with mantel 1.18 (Cavalcanti, 2005). Pairwise comparisons were made of geographic distances and \( F_{ST} \). A Bayesian approach was used to assess population structure by model-based clustering methods with the program BAPS 3.2 (Corander et al., 2005), which identifies the optimum number of partitions among groups of samples. Furthermore, genotypes were plotted in a multidimensional space, using a principal-component analysis (PCA) with the program pca-gen 1.2 (Goudet, 1999) and labelled as 1 (Exe), 2 (Teign), 3 (Tamar), 4 (Looe), 5 (Camel) and 6 (Torridge).

Finally, a hierarchical analysis of molecular variance (AMOVA, Excoffier et al., 1992) was conducted using arlequin 3.11 (Excoffier et al., 2005) using an a priori design which divided all populations into three groups corresponding to a geographic distance greater than 100 kilometres (Group 1: Exe, Teign; Group 2: Tamar, Looe; Group 3: Camel, Torridge).

Morphometric data analysis

Fifty males and fifty females from each population were used in the morphometric analysis. To quantify possible shape variation in the carapaces and in the chelae, landmark-based geometric morphometrics was used. Images of each specimen were taken using a Canon Power Shot A85 digital camera with a resolution of 4.0 megapixels and consistent zoom and distance, in order to avoid false perspectives and allow accurate comparisons. Fifteen carapace landmarks and nine claw landmarks were digitized using the program tpsDig 2.10 (Rohlf, 2006) (Fig. 2). Only one side of the carapace (the left) was used to avoid duplication of equivalent landmarks and computation problems (Rufino et al., 2004).

The digitized landmark configurations were then subjected to a generalized Procrustes analysis (GPA; Rohlf and Slice, 1990) to remove the effects of size, position and orientation in the digital images. The aligned landmark configurations produced by this analysis were used to generate the shape variables, the relative warp scores and the uniform component, from a thin-plate spline analysis (TPS). These variables were obtained with tpsRelw 1.45 (Rohlf, 2007a). The software tpsRegr 1.34 (Rohlf, 2007b) was used to detect the presence of allometry, and when detected, it was removed by the regression of each relative warp against a measure of body size (centroid size), thus estimating residual shape variation.

To understand patterns of morphometric differentiation among populations, a multivariate analysis of variance (MANOVA) was performed on the relative warps scores. After assessment of the degree of variation among populations, pairwise comparisons among these populations were done with the post-hoc Tukey HSD test. In addition, discriminant function analysis was used to determine Mahalanobis distances between each pairs of sites, for a more accurate differentiation between populations. Finally, a cluster analysis was generated from the Squared Mahalanobis distance matrix to assess phenotypic relationships among the populations. These procedures were performed in statistica 6.0.

RESULTS

Genetic data

A total of 270 Carcinus maenas collected from southwest England were typed for eight microsatel-
literate loci. All loci were highly polymorphic (number of alleles: 30 for Cma10EPA, 47 for Cma11EPA, 36 for Cma12EPA, 12 for Cma14EPA, 11 for SP107, 24 for SP280 and 10 for SP495), with the exception of locus SP229, for which only one allele was detected. Therefore, SP229 was not used in further analyses. All the populations were characterized by a considerable mean number of alleles per locus, ranging from 13.71 to 16.29, and by a number of private alleles ranging from 3 to 8. The expected heterozygosity (H_{exp}) per locus ranged from 0.356 to 0.957 and the observed heterozygosity (H_{obs}) from 0.347 to 0.795 (Table 1). Three loci showed a significant heterozygosity defect (Table 1, p-value <0.05). Considering the Hardy-Weinberg Equilibrium (HWE) results by locus and by population, loci Cma10EPA, Cma11EPA and SP280 deviated from HWE in all populations. In the other four loci, no evidence of heterozygote deficiencies was found, so it was possible to assume that all populations were in HWE. The observation of many cases of excess homozygosity at a single locus may suggest the presence of null alleles. This was confirmed by the results of MICRO-CHECKER 2.2.3. After correction for null alleles with the Brookfield null allele estimator 1 (Brookfield, 1996), subsequent analyses were performed on the corrected data set. Genotyping errors due to stuttering and large allele dropout were not found (confidence interval = 95% and 10000 iterations).

Fisher exact test for linkage disequilibrium among loci across populations revealed no significant p-values among 21 possible combinations.

Table 1. – Main genetic variability measures by locus of Carcinus maenas. T (ºC), annealing temperature; N, number of alleles found per locus; H_{exp}, expected heterozygosity; H_{obs}, observed heterozygosity; Ar, allelic richness; p, Hardy-Weinberg test p-value.

<table>
<thead>
<tr>
<th>Locus</th>
<th>Primer sequence 5’-3’</th>
<th>Repeat</th>
<th>T (ºC)</th>
<th>N</th>
<th>H_{exp}</th>
<th>H_{obs}</th>
<th>Ar</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cma10EPA</td>
<td>F:GAGACCCTGCAATGCGATTTCCTCT</td>
<td>(CA)_{2}</td>
<td>59</td>
<td>30</td>
<td>0.947</td>
<td>0.784</td>
<td>22.22</td>
<td>0.012</td>
</tr>
<tr>
<td>Cma11EPA</td>
<td>F:GGGAGAAGCTGCTGAGTCACT</td>
<td>(CA)_{3}</td>
<td>55</td>
<td>47</td>
<td>0.96</td>
<td>0.732</td>
<td>29.71</td>
<td>0.001</td>
</tr>
<tr>
<td>Cma12EPA</td>
<td>F:GGCGGAGCTGCTGAGTCACT</td>
<td>(GT)_{3}</td>
<td>60</td>
<td>36</td>
<td>0.952</td>
<td>0.795</td>
<td>26.1</td>
<td>0.054</td>
</tr>
<tr>
<td>Cma14EPA</td>
<td>F:GGGAGAAGCTGCTGAGTCACT</td>
<td>(CC)_{4}</td>
<td>60</td>
<td>12</td>
<td>0.356</td>
<td>0.347</td>
<td>7.38</td>
<td>0.352</td>
</tr>
<tr>
<td>SP107</td>
<td>F:GTACCCGGGAAAGCGAGTACG</td>
<td>(GAG)_{1}</td>
<td>51</td>
<td>11</td>
<td>0.592</td>
<td>0.719</td>
<td>7.2</td>
<td>0.434</td>
</tr>
<tr>
<td>SP280</td>
<td>F:GTACCCGGGAAAGCGAGTACG</td>
<td>(GTA)_{2}</td>
<td>49</td>
<td>24</td>
<td>0.717</td>
<td>0.37</td>
<td>9.72</td>
<td>0.001</td>
</tr>
<tr>
<td>SP495</td>
<td>F:AAAGTCAGGCGGCTAGTGA</td>
<td>(GAG)_{10}</td>
<td>52</td>
<td>10</td>
<td>0.649</td>
<td>0.731</td>
<td>6.7</td>
<td>0.358</td>
</tr>
</tbody>
</table>

Table 2 shows F_{ST} estimates across all loci for all population pairs. F_{ST} values ranged from 0.0015 to 0.0458, illustrating low levels of genetic structuring. Following Bonferroni correction, only three of the 15 pairwise F_{ST} values were found to be significant (between Exe-Tamar; Tamar-Camel; Tamar-Torridge). No significant isolation-by-distance was shown when pairwise F_{ST} values and geographic distance were compared (Mantel test, 10000 randomizations, r =-0.224, p =0.205). The hierarchical analysis of molecular variance (AMOVA) showed that genetic variation within and among populations amounted to 97.98% and 1.4% of total variation, respectively. The genetic variation among pre-defined groups was close to zero (0.62%). Overall F_{ST} and the F_{SC} values were low (0.0202 and 0.01404 respectively), and not significant (p >0.05). This indicates no genetic differentiation among all populations sampled and among populations within groups, respectively. The F_{ST} value found was also very small (0.00622) and non-significant (p >0.05), indicating low genetic differentiation among the proposed groups. The BAPS analyses, conducting assumed different numbers of groups of population, indicated that the best partition included all six sampled populations (ln likelihood = -8753.2). The result of PCA on genotypes is

Table 2. – Genetic differentiation among Carcinus maenas populations. F_{ST} values are below diagonal, and geographic distances (km) among populations are above diagonal. * p<0.05: significance of F_{ST} after sequential Bonferroni correction.

<table>
<thead>
<tr>
<th></th>
<th>Exe</th>
<th>Teign</th>
<th>Tamar</th>
<th>Looe</th>
<th>Camel</th>
<th>Torridge</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exe</td>
<td>0.0099</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Teign</td>
<td></td>
<td>0.0458</td>
<td>0.0014</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tamar</td>
<td></td>
<td></td>
<td>0.352</td>
<td>0.279</td>
<td>0.356</td>
<td></td>
</tr>
<tr>
<td>Looe</td>
<td></td>
<td></td>
<td></td>
<td>0.0021</td>
<td>0.02</td>
<td>0.952</td>
</tr>
<tr>
<td>Camel</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.0106</td>
<td>0.124</td>
</tr>
<tr>
<td>Torridge</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.0015</td>
</tr>
</tbody>
</table>

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Fig. 3. – Scores of microsatellite genotypes plotted on the first two axes (PC-I and PC-II) of a principal component analysis performed using PCA-GEN. 1; Exe; 2; Teign; 3; Tamar; 4; Looe; 5; Camel; 6; Torridge.
shown in Figure 3. Scores were plotted onto the two principal axes (PC-I and PC-II), which cumulatively explain 74.7% of the total genetic diversity. The $p$-value for the overall $F_{ST}$ was 0.01 and the global $F_{ST}$ was 0.02616. Per axis, the $F_{ST}$ values achieved were, respectively, 0.01287, 0.00667, 0.00270, 0.00208 and 0.00184. This analysis showed a separation of all populations, so no clear clustering was formed. However, the Exe (1) and Torridge (6) populations seemed to be closer together than to the remaining populations, and the Tamar (3) population was further apart from all the other populations.

**Morphometric data**

Analyses performed in the male carapaces revealed that the first relative warp (RW1) explained 47.6%, the second (RW2) 12.5% and the third (RW3) 8.1%, summing 68.2% of variance explained. The first axis revealed shape variation in the rostrum and on the overall carapace shape, with the negative end of this axis corresponding to more round shapes. RW2 explained shape variation in the lateral-posterior spines and the distances among them, and RW3 accounted for differences due to the variation in the carapace width (Fig. 4). A similar variation pattern was observed in females, but the rounder carapaces corresponded to the positive values of relative warp 1. In this case, the first three axes explaining 43.3%, 11.5% and 9%, respectively (Fig. 4). Shape variation in the chelae was more marked. For the males, RW1 explained 53.7% of the variation explained by all the relative warps, RW2 explained 12.8% and RW3 explained 9.4%. The first warp accounted for the variation due to the size of the pollex, the height of the manus and the flattening of the posterior part of

![Fig. 4. – Scatterplot of individual scores from the relative warp analysis. Grids show deformation from consensus configuration of the carapace of *Carcinus maenas* males (A) and females (B).](image-url)
the claw (Fig. 5). The variation explained by RW2 corresponded to the orientation of the pollex and the total length of the chela. Relative warp 3 accounted for the shape variation due to the size of the gape. In accordance with the shape variation of the carapaces, a similar shape variation pattern was found in the females, with the first three axes explaining 53.3%, 11.3% and 8.9%, respectively (Fig. 5).

The multivariate analysis of variance revealed significant morphometric differences among the nine populations of *Carcinus maenas* (males: Wilks’ $\lambda = 0.189$, $p < 0.0001$; females: Wilks’ $\lambda = 0.226$, $p < 0.0001$). The comparisons made among populations, obtained with the post-hoc Tukey HSD test, showed that the Torridge population was significantly different from almost all the other populations. Additionally, the post-hoc test revealed the absence of significant differences between the sites that were closer together. In contrast, the sites further apart tended to show significant differences between each other (Table 3). The discriminant function analysis and cluster analysis revealed that there was extensive inter-population variability in the morphology of both male and female *C. maenas*, but differences were more pronounced in males, as was detected by the squared Mahalanobis distance. Nevertheless, the patterns exhibited by each sex were similar. Distances between the morphology of male crabs showed two well-defined clusters (Exe, Axe and Teign; Hayle, Looe and Tamar). Males from Fowey, Camel and Torridge exhibited the greatest differences from these groups (Fig. 6). The cluster analysis performed in the females revealed three clusters on the north coast (Hayle, Camel and Torridge) and two overlapping clusters on the south coast (Exe, Teign, Looe; and Axe, Fowey and Tamar). However, Hayle, Camel and Torridge females were most distinct in terms of their morphology.
**DISCUSSION**

*Carcinus maenas* exhibited no clear evidence of genetic differentiation among the sites sampled on the southwest English coast. Different analyses, including AMOVA and the Mantel test, revealed no pattern of geographic separation among populations. Therefore, genetic variation was not associated with geographic subdivision. Furthermore, the Bayesian approach used to assess genetic population structure suggested that all sampled individuals comprised a single genetic population. Pairwise *F_{ST}*, low values suggested no restriction of movement among sites and, consequently, no impediment to gene flow between populations.

Migrations during larval development to the marine environment promote transport over large distances, allowing gene flow between populations (Bilton et al., 2002). *Carcinus maenas* has a long planktonic larval phase (up to 50 days) that can be dispersed extensively in the water column before settling to the substratum and undergoing metamorphosis. Peliz et al. (2007) conducted a study with *C. maenas* along the west coast of the Iberian Peninsula, and suggested a larval dispersal distance of about 60 km (s = 20 km), which is mostly representative of alongshore larval dispersion. Because sampled populations along the studied coast are separated by distances of 15 to 432 km, such a dispersal radius may account for considerable exchange of individuals between local populations. Thus, the long planktonic development mode of *C. maenas* can allow larval flow between distant populations and, consequently, might promote population homogenization. These results further suggest that there are high degrees of connectivity within the study area with little evidence of reduction in gene flow, either resulting from barriers that block the dispersion potential or failures in the reproduction of migrant individuals within the new population to which they are dispersed.

A recent study by Pascoal et al. (2009) with microsatellite data detected very weak but significant structuring between *C. maenas* populations on the Portuguese coast, which, according to the authors, might result from hydrodynamic or topographic barriers. On the Portuguese coast, *C. maenas* forms large populations in estuaries and on their adjacent rocky shores. The geographic distance that separates estuaries may be characterized as a barrier, thus limiting larval flux between populations. On the other hand, *C. maenas* has a continuous distribution on the English coast, occurring on all types of shores (Neal and Pizzola, 2008), and no evidence of genetic differentiation among the sites sampled was found in our study. These differences detected in the population structure of *C. maenas* could therefore be due to the contrasting distribution patterns exhibited on both coasts.

The population from the River Tamar Estuary revealed statistically significant levels of population differentiation when compared with the Exe, Camel and Torridge populations. The River Tamar is a large river, and its drainage to the Atlantic Ocean may affect nearshore currents and salinity, and influence larval transport. Newly hatched larvae migrate vertically in synchrony with the tidal cycle, attaining their highest position in the water column during ebb (Queiroga, 1996). This migration, under behavioural control (Zeng and Naylor, 1996), ensures that larvae are exported to the sea shortly after spawning. The influence of estuarine currents and brackish water can also limit the pattern of offshore larval dispersal and onshore megalopal transport, favouring genetic differentiation from the rest of the populations. A similar pattern of population differentiation was found in the Portuguese Tagus Estuary and on its adjacent rocky shores for the shore

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**Table 3.** Pairwise comparisons among populations achieved with the post-hoc Tukey HSD test. Female values are above diagonal and male values are below diagonal. **p<0.001; n.s., non significant value.

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**Fig. 6.** UPGMA dendrogram derived from cluster analysis of squared Mahalanobis distances for relative warp scores, for the major chela of *Carcinus maenas* males.
The results achieved in the current study, in accord-
ance with Brian et al. (2006), suggest that genetic and
phenotypic characters are in some way independent,
and that environmental factors, such as habitat, wave
action, predation, food supply and salinity can play
an important role in the patterns of shape variability.
However, the extent of plasticity in shore crabs is un-
clear. Two aspects of the life history of the shore crab
may have led to the evolution of plastic responses
(Todd et al., 2006): due to its pelagic larval stage, high
levels of genetic homogeneity are expected, so there
is limited opportunity for local adaptation (Roman and
Palumbi, 2004); and post-settlement movements can
occur, particularly in response to predator pressures,
even though settling megalopaes have swimming and
habitat-selection ability (Moksnes et al., 1998). Fur-
thermore, the morphology of C. maenas may respond
plastically to site-specific selection pressures, causing
changes in the expression of a particular morphological
trait (Brian et al., 2006).

Carcinus maenas is a heterochelous species, and
Silva and Paula (2008) have shown that the crusher
claw is more suitable for population differentiation
studies. In the current study, clearer patterns of shape
variation were seen in the crusher claw than in the cara-
pace. Previous studies have shown that claw size is ca-

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