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Geometric morphometrics, Fourier analysis of otolith shape, and nuclear-DNA markers distinguish two anchovy species (*Engraulis* spp.) in the Eastern Mediterranean Sea

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ABSTRACT

Two anchovy forms identified from their external morphology and designated as ‘Silver’ and ‘Blue’ anchovies were captured in Mersin Bay (Eastern Mediterranean Sea). Silver anchovy seemed to inhabit shallower waters (9-77 m) than Blue anchovy (majority at depths of 69-111 m). Significant differences in body shape, which was analyzed through geometric morphometrics, and in otolith shape, examined using Fourier analysis, separated the two forms. Length polymorphisms at two intron and 9 microsatellite loci showed strong genetic differences between the two forms. Mersin-Bay Blue anchovy were genetically related to *Engraulis encrasicolus* from the Western Mediterranean Sea. Mersin-Bay Silver anchovy possessed an original genetic composition that distinguished them from both *E. encrasicolus* and *E. albidus* at the two intron loci, while presenting affinities to *E. albidus* based on microsatellite markers. Mersin-Bay Silver anchovy were assigned to the latter species, of which they are considered as a geographically isolated population. This is the first record of *E. albidus* in the Levantine basin and confirmation of its presence in the Eastern Mediterranean Sea. These results have implications for the management of anchovy populations and their habitats.

Keywords: *Engraulis albidus*; *E. encrasicolus*; introns; microsatellites; Mersin Bay

1. Introduction

The degree of geographic isolation among European anchovy populations has been addressed through a number of population genetic and phylogeographic studies (Spanakis et al., 1989; Bembo et al., 1996a, b; Magoulas et al., 1996; Magoulas et al., 2006; Bouchenak-Khelladi et al., 2008; Kristofferson and Magoulas, 2008; Erdogan et al. 2009). Complex patterns of geographical structure in the European anchovy have been revealed, where the level of genetic differentiation between inshore and offshore forms within a basin (e.g. Western Mediterranean Sea) exceeds that within the offshore form at the scale of the whole Mediterranean Sea (Borsa, 2002; Grant, 2005; Bouchenak-Khelladi et al., 2008).

It has been hypothesized that the remarkable degree of anchovy population genetic structure in the Mediterranean Sea derives from its higher geographic and oceanographic complexity relative to other regions (Grant, 2005). While such extrinsic barriers to gene flow likely determine part of the genetic differences between anchovy populations, intrinsic (reproductive) barriers have also been documented (Borsa, 2002). Restriction in gene flow between two habitat-specific forms sampled locally in a context where the anchovy form inhabiting the pelagic habitat shows relative genetic homogeneity over a broad geographical range (the Mediterranean Sea), indicates reproductive isolation between the two forms. This has led Borsa et al. (2004) to distinguish the two forms as different species, where *Engraulis albidus* is the species inhabiting the inshore waters of the north Western Mediterranean Sea and presumably the northern Adriatic Sea, and *E. encrasicolus* is the species inhabiting the pelagic waters of the north Western Mediterranean Sea. Two genetically distinct forms also co-occur in the Bay of Biscay, where they segregate according to habitat (Zarraonaindia et al., 2012) and where distinct coastal and oceanic spawning areas have been reported (Motos et al., 1996). Some of the populations sampled in the Bay of Biscay are genetically closer to some populations from the Mediterranean Sea than to adjacent populations in the northeastern Atlantic (Zarraonaindia et al., 2012). Kristofferson and Magoulas (2008) have compared anchovy populations from the central Aegean Sea, the Ionian Sea, the Adriatic Sea and the Gulf of Lions with regard to mtDNA clade proportions, nuclear allele frequencies, body shape and otolith shape. They concluded that several reproductively isolated anchovy populations also exist in the western and central regions of the Eastern Mediterranean Sea.

Using morphometric characters, Turan et al. (2004) reported dissimilarity among the anchovy populations inhabiting the eastern Levantine basin and those found in the Black Sea and the Aegean Sea. However, the authors did not consider possible dissimilarities within regions.

In the present study, two anchovy forms distinct by their general appearance and colour were sampled in Mersin Bay (northeastern area of the Eastern Mediterranean Sea). The morphology of individuals was examined using geometric morphometric analysis, and otolith shape was described using Fourier analysis. The aim of this study was to determine whether the morphological differences between the two forms reflected genetically distinct populations, so the two forms were further characterized by their allelic composition at nuclear genetic markers. The two forms were further compared to reference samples of *E. albidus* and *E. encrasicolus* from the Western

Mediterranean Sea to determine their genetic relationships to either species.

2. Materials and methods

2.1. Study site and samples

Mersin Bay, in the Levantine basin, Eastern Mediterranean Sea (36°31'N-36°48'N; 34°13'E-34°44'E) (Fig. 1) is a potentially important area for commercial fisheries (Abdulla et al., 2008). Two hundred and twenty nine anchovy were collected by trawling in Mersin Bay (Fig. 1) in June 2010, from the research vessel *Lamas* of the Middle East Technical University, Institute of Marine Sciences, Mersin, Turkey. Anchovy were visually sorted into two groups according to their external characteristics by the fisheries scientists present on board: a proportion of individuals possessed a slender and rounded body shape, with a firmer body and a darker (blue) back than the other individuals, which were generally paler with a well-delineated longitudinal silver stripe along the flank (Fig. 2), and with a broader belly and a softer texture. Hereafter, the two forms will be referred to as “Blue” and “Silver” anchovies, respectively. Blue anchovies (Fig. 2a) were captured above the bottom at depths between 9 - 111m (the majority occurring between 69 -111m) whereas Silver anchovies (Fig. 2b) were caught in the inshore waters of northeastern Mersin Bay, at depths of 9 m -77 m. Immediately after collection, the anchovy were frozen at -20°C, shipped to the Middle East Technical University Institute of Marine Sciences, Mersin and kept frozen until processed. Ninety three Blue anchovy and 136 Silver anchovy were used for geometric morphometric analysis; 22 Blue anchovy and 20 Silver anchovy were used for Fourier analysis of otolith shape; and 48 individuals randomly selected from each form were used for analysis of genetic variation. Genetic analyses were carried out in the population genetics laboratory of the Middle East Technical University, Ankara.

Anchovy samples from Annaba, Algeria (sample Ann; $N=15$), from the Benguela current system, southern Africa (Ben; $N=15$), and from Cul-de-Beauduc (Cul; $N=7$) and Sète (Set; $N=15$) both located in the northern region of the Western Mediterranean Sea [i.e. subsamples of, ANN, BEN, CUL and SET samples respectively of Bouchenak-Khelladi et al. (2008)] were used as reference samples for genetic comparison. The CUL sample consists of reference material for *E. albidus* while the SET sample consists of *E. encrasicolus* specimens. Cul-de-Beauduc and Sète are the respective type localities of these two species (Borsa et al., 2004).

2.2. Analysis of body shape by geometric morphometrics

Geometric morphometrics is based on body landmarks. Differences in the relative positions of landmarks indicate differences in shape between specimens (Bookstein, 1991). The geometric morphometrics method involves five steps: (1) capturing the images of individuals; (2) quantifying shape by digitization of landmarks on the images using the TPS software (Rohlf, 2001); (3) superimposing the landmarks; (4) visualization of shape variation; and (5) analysing the shape

variables using principal component analysis (PCA) (Pearson, 1901) and discriminant analysis (DA) (Lebart et al., 1997). Procrustes superimposition consists of translating the configurations defined by the raw coordinates to a common centroid, scaling to unit centroid size, and rotating until the sum of the squared distances between corresponding landmarks is minimized (Rohlf and Slice, 1990). General Procrustes analysis and digitization of deformation grids (Bookstein, 1989) for 20 landmarks on Blue and Silver anchovy samples were run using the Morpho-J, version 1.02d software (Klingenberg, 2011). Deformation grids, which employ the Procrustes distance as a metric, provide a visually interpretable description of deformation, with the same number of variables as there are statistical degrees of freedom.

Principal component analysis was run using PAST version 2.01 (Hammer et al., 2001). Discriminant analysis was run on the Procrustes coordinates for body shape, using cross-validation to assess the effectiveness of the algorithm (Polly and Head, 2004). To mitigate possible effects of allometric growth, only adult specimens were measured.

The left side of each individual was photographed with a digital Olympus C-7070 camera (image resolution: 7.1 megapixels) (Fig. 2a,b). Twenty landmarks (Fig. 2c) were digitized from each photograph using the TpsDig version 2.12 software. The X-Y coordinates of the landmarks were recorded. Sex was determined from the visual aspect of the gonads, which were examined after the individuals had been photographed.

2.3. Fourier analysis of the outline of otoliths

The outlines of the images of otoliths (sagittae) of 1+ year-old Blue ($N=15$) and Silver ($N=12$) anchovies, and those of 2+ year-old Blue ($N=7$) and Silver ($N=8$) anchovies were subjected to Fourier analysis. Both left and right sagittae were extracted, cleaned, and photographed under a dissecting microscope (Olympus SZx12) using an Olympus C-7070 camera. The age (in years) of each individual was estimated from the number of translucent zones in the sagittae, under the assumption that a translucent zone represents the winter season (Waldron, 1994). Variation in otolith shape was assessed using elliptical Fourier analysis (Kuhl and Giardina, 1982; Ferson et al., 1985). The outline of the otolith was traced using the softwares tpsDig2 and EFAWin (Rohlf, 1993; Isaev, 1995). The x - and y - coordinates of 120 landmarks were obtained from points equally spaced along the otolith's outline using tpsDig2. Next, a set of harmonics that describe the shape of the otolith was produced using EFAWin. Twenty one harmonics were used to obtain a good balance between the ratio of sample size to the number of variables, and the accuracy of the description of otolith shape (Fig. 3). The harmonics were made invariant to otolith size, rotation, and starting position of the tracing of the outline (Ferson et al., 1985; Kristofferson and Magoulas, 2008). Principal component analysis was run using the EFAWin scores with the PAST statistical program. The generalized Mahalanobis distance (Mahalanobis, 1936) was used to estimate differences in otolith shape among anchovy populations. Mahalanobis distances were computed using Morpho-J. Canonical discriminant analysis (CDA) was run under SYSTAT v. 12.02.00 to estimate differences in otolith shape between age groups of Blue and Silver anchovies.

2.4. Genetic analyses

Genomic DNA was extracted from 100 mg muscle tissue using the CTAB protocol (Stewart and Via, 1993). The amount of DNA from each sample was subsequently quantified by spectrophotometry. Individuals were genotyped at 12 size-polymorphic nuclear loci, including two introns and 10 microsatellites (Appendix A). The forward primers were fluorescently labeled with HEX, TAMRA or FAM. The quality of the PCR amplifications was checked by subjecting the PCR products to electrophoresis in 0.8-4 % agarose gels. Allele size was determined in an ABI-PRISM 3100 sequencer (Applied Biosystems, Foster City CA, U.S.A.). Raw data from the sequencer were processed with the PEAK SCANNER1.0 software for peak identification and fragment sizing (Applied Biosystems).

Expected heterozygosity (H_e ; Nei, 1978) and observed heterozygosity (H_o) were calculated using GENETIX v.4.05.2 (Belkhir et al., 2000). Allelic richness (AR) and the inbreeding coefficient (f ; Weir and Cockerham, 1984) were estimated using FSTAT v. 2.9.3.2 (Goudet, 2002). Genetic differentiation between samples was estimated as Weir and Cockerham's (1984) θ using GENETIX. The Bonferroni correction (Rice, 1989) was applied for multiple tests. The presence and frequency of null alleles was tested using the expectation maximization algorithm of Dempster et al. (1977) implemented in FreeNA (Chapuis and Estoup, 2007). The null hypothesis of Hardy-Weinberg equilibrium at a locus was tested by using the GENEPOP version 4.1.1- (Raymond and Rousset, 1995; Rousset, 2008) which implements Fisher's exact test for multiple alleles (Guo and Thompson, 1992). Outlier f values were detected using Minitab v.16.0 (Minitab Inc., 2010). A value was considered to be an outlier when it lay outside the interval delimited by one interquartile subtracted from and added to the values of the first and third quartiles respectively.

Bayesian structure analysis was run using the clustering algorithm implemented in the software STRUCTURE 2.1 (Pritchard et al., 2000; Falush et al., 2007). The number of populations assumed by STRUCTURE is K , each of which is in the Hardy-Weinberg and linkage equilibrium and characterized by a set of allele frequencies. To estimate the assignment of each individual to a cluster, ten replicate simulations were run for each prior value of K in the total sample. K was allowed to range from 2 to 7, using 10,000 burn-in repetitions and a final run of 100,000 Markov-chain-Monte Carlo steps under an admixture model with independent allele frequencies. The CLUMPP software (Jakobsson and Rosenberg, 2007) was used to calculate similarity coefficients. To determine the actual number of populations, we used Evanno et al.'s (2005) method, which checks the second-order rate of the likelihood function ($\Delta K = mL''(K) / sL(K)$) (where $mL''(K)$ is the mean of the absolute values of $L''(K)$ and $sL(K)$ is the standard deviation of $L(K)$).

The correlation between genotypes at pairs of loci was determined using Weir's (1979) measure of genotypic disequilibrium, R_{ij} . Random permutations of genotypes at a locus (GENETIX v. 4.05.02) allowed us to generate the distribution of R_{ij} under the null hypothesis of genotypic equilibrium. Correspondence analysis (CA; Benzécri, 1982) was run to visualize samples characterized by their allele frequencies, by reducing the multidimensional allelic frequency space

to a bidimensional space. Two CAs were run on allele-frequency data, the first one on the matrix of samples x allele-frequencies derived from Appendix B (26 samples x compound-alleles at 2 intron loci), and the second one on the matrix derived from Appendix C (12 samples x compound-alleles at 3 microsatellite loci). Hierarchical clustering analysis (Lê et al., 2008) was used to delineate clusters of samples; for this, pairwise distances between samples were Euclidean distances in the space defined by the first five axes of the CA.

3. Results

3.1. Geometric morphometrics

Clear-cut differences were observed in the body shapes of Blue and Silver anchovies from Mersin Bay, whereas no effect of sex on body shape was visible (Fig. 2e). The Mahalanobis distance between male and female specimens was 1.72 for Blue anchovy and 1.68 for Silver anchovy. These differences were not significant (test on T-square statistic; 1000 permutations; $P > 0.05$), hence, males and females were pooled for subsequent analysis. The Mahalanobis distance between the two morphotypes was 4.59 ($P < 0.001$). According to DA, the assignment of Blue anchovy to their own group was 95.6% and that of Silver anchovy was 96.8% (Fig. 2f), thus formalizing the initial distinction of two forms from their general, external features.

The deformation grid for the body shape of Blue, relative to Silver anchovy is given in Fig. 2d. This figure points out remarkable deformation at the head and posterior parts of the ventral region.

3.2. Otolith shape

The results of the Fourier analysis of otolith shape are summarized in Fig. 3b,c. Otoliths of individuals belonging to different age groups were analyzed separately. The otoliths of Silver anchovy differed from those of Blue anchovy in both the 1+ and 2+ age groups. Although some marginal overlap was present in 1+ individuals (Fig. 3b), no overlap was observed between Silver and Blue anchovies aged 2+ (Fig. 3c). The CDA discriminated Blue from Silver anchovies with reclassification scores of, respectively, 80% and 85% for 1+ individuals (Wilk's $\Lambda = 0.017$; $P < 0.001$), and, respectively, 86% and 100% for 2+ individuals (Wilk's $\Lambda = 0.003$; $P < 0.001$).

3.3. Genetic analyses

Summary statistics (allele number, allelic richness, expected heterozygosity, observed heterozygosity, inbreeding coefficient estimates, estimated null allele frequencies) for Blue and Silver anchovies from Mersin Bay and for four reference anchovy populations, all genotyped at 12 nuclear loci, are reported in Table 1. To assess the suitability of the genetic markers for investigating population structure, we first checked the general conformity of the observed genotype distributions relative to those expected under the Hardy-Weinberg equilibrium; this was

done with the two samples from Mersin Bay which unlike the other samples were large enough to allow meaningful estimations of f . Locus *EeJ27.1* exhibited much higher f values (0.502 and 0.526 in Silver and Blue anchovies, respectively) than the other loci (Table 1), hence was designated as an outlier in both Silver and Blue anchovies (Appendix B). Moreover, the null allele frequencies inferred at locus *EeJ27.1* (Table 1) in the two anchovy populations from Mersin Bay (0.24, 0.25) were higher than the generally accepted threshold value ($r = 0.20$) above which the impact of null alleles on θ estimates can be considered a problem (Chapuis and Estoup, 2007). Thus, *EeJ27.1* was excluded from further analysis. The most polymorphic marker was *EeOV04* with 34 different alleles while *CK6-1* was the least polymorphic with 10 alleles (Table 1). The distributions of allele frequencies by sample are summarized in Supplementary material, Fig. S1.

The outcomes of Bayesian structure analysis for $K = 2-7$ are summarized in Supplementary Material, Table S2 and Fig. S2. Using Evanno et al.'s (2005) approach, the most likely number of clusters (K) was determined as 2 (Supplementary material, Fig. S3). No significant genetic disequilibrium (R_{ij}) value was detected (Supplementary material, Table S3).

3.4. Comparison with other anchovy populations

Pairwise θ estimates between populations ranged from 0.017 (between Ann and Set) to 0.094 (between Blue anchovy and Cul) and were all significant (Table 2). Mersin-Bay Silver anchovy appeared to be genetically closer to the reference *E. albidus* than to the reference *E. encrasicolus*. Likewise, Mersin-Bay Blue anchovy were genetically closer to *E. encrasicolus* (Table 2). To analyze the placement of Mersin-Bay Silver and Blue anchovies in the phylogeographic context of the entire Mediterranean Sea, we compared them on the basis of allele frequencies to the widest possible range of samples of individuals. We thus compared the allele-frequency data at loci *CK6-1* and *CK6-2* obtained in this study with previously published, homologous data from across the Mediterranean Sea (Bouchenak-Khelladi et al., 2008; Kristofferson and Magoulas, 2008). This required pooling alleles as explained in Appendix B. The resulting matrix of allele frequencies per sample (Appendix B) was subjected to correspondence analysis (CA). Comparisons were also made using microsatellites, between this study and that of Borrell et al. (2012). We obtained the genotype data of Borrell et al. (2012) concerning loci *EeOV04*, *Ee-10*, *Ee2-508*, *Ej-27.1* and *Ej-41.1* (Y.J. Borrell, pers. comm.). Locus *Ej-27.1* was discarded because of the inferred high proportion of null alleles in our samples (see above) and so was *Ej-41.1* because we were unable to establish cross-study relationships between alleles at this locus. Alleles at the three remaining loci were pooled as explained in Appendix C. The resulting matrix of allele frequencies per sample (Appendix C) was subjected to CA. The outcomes of the CAs on allele frequency data are presented in Fig. 5a (introns) and Fig. 5b (microsatellites). The Mersin-Bay “Blue” anchovy sample was included within the *E. encrasicolus* cluster (Fig. 5a, b), thus confirming its identity as *E. encrasicolus*. Correspondence analysis on microsatellite data further suggested “Silver” anchovy from Mersin Bay to be a genetically differentiated, local population of *E. albidus* (Fig. 5b).

4. Discussion

A trivial but crucial and sometimes overlooked problem in fish ecology and fisheries is to accurately identify the fish to species. This is a requirement for correctly estimating biomass, productivity and population trends, which in turn helps define management strategies (Ward, 2000). Inadvertently or intentionally confounding multiple species in an assessment would amount to ignoring the different population dynamics of each individual species due to different life histories. In the case where fish are commercially exploited, it is also important that they be correctly labelled to species through the market chain.

4.1. Two anchovy species in Mersin Bay

Anchovy specimens from Mersin Bay presented two distinct phenotypes or forms differing by their general morphology, here coined “Blue” and “Silver” because of correlated colour differences. Here, we confirmed that the two forms also differ in their body shape as defined by landmarks, and that they differ significantly by their otolith shape. The two forms were segregated according to habitat: Blue anchovy tended to occur in the deeper areas of the bay, whereas Silver anchovy occurred in shallower, inshore waters.

Morphometric analysis has long been used as a tool for stock identification (Cadrin and Friedland, 1999). Silva (2003) examined morphometric variability using both the distance and landmark morphometric data of *Sardina pilchardus* from the northeastern Atlantic. Both morphometric methods indicated that southern Iberian (southern Portugal, Gulf of Cadiz) and northern Moroccan *S. pilchardus* specimens had a morphotype distinct from those in the other areas. Discrimination of the two morphotypes was also confirmed statistically by the significant difference between group centroids and by the high percentage of correct classification (85%). The assignment of individuals to their own group was even higher (>95 %) for both anchovy populations from Mersin Bay. This high score demonstrates a high level of morphological differentiation between the two sympatric forms.

Otolith shape has been widely used to discriminate closely related fish species (L’Abée-Lund and Jensen, 1993; Tuset et al., 2003; Ponton, 2006; Wakefield et al., 2014). It has also been used successfully to distinguish populations within a single species, *Melanogrammus aeglefinus* (Begg and Brown, 2000). Tracey et al. (2006) compared otolith forms of two widely separated *Latris lineata* populations using elliptical Fourier analysis. They achieved reasonable success in demonstrating population differentiation based on otolith shape, which was further supported by mitochondrial DNA differences. DeVries et al. (2002) used otolith Fourier coefficient analysis to distinguish Atlantic king mackerel (*Scomberomorus cavalla*) populations from the eastern Gulf, reaching high classification success (71-86%). In the present study, the reclassification success of Mersin Bay Blue and Silver anchovies was between 80% and 100%, which again indicates strong differentiation.

The two anchovy forms from Mersin Bay also differed by their allelic frequencies at eleven

nuclear loci, suggesting that the morphological differences do not result from pure phenotypic plasticity.

Altogether, these observations are strongly reminiscent of the differences observed between the two forms (also sometimes coined “Blue” and “Silver” anchovies) reported from the Adriatic Sea (Bembo et al. 1996a) and from the north Western Mediterranean Sea (Pasteur and Berrebi, 1985; Borsa, 2002). The latter observations have eventually led to recognizing two anchovy species, *E. encrasicolus* and *E. albidus*, in the Western Mediterranean Sea (Borsa et al., 2004; Bouchenak-Khelladi et al., 2008). The present results indicate a level of genetic differentiation between Silver and Blue anchovies from Mersin Bay comparable to that observed between *E. albidus* and *E. encrasicolus* in the Western Mediterranean Sea.

4.2. Comparison of Mersin Bay anchovies with other populations

The recognition of at least two anchovy species in the Western Mediterranean Sea (Borsa, 2002; Borsa et al., 2004; Bouchenak-Khelladi et al., 2008) was originally inferred from higher similarities at allozyme loci between inshore anchovy of the north Western Mediterranean Sea and the northern Adriatic Silver anchovy, than between either of these populations and their geographically closest counterparts, namely Blue anchovy from a few tens of kilometers apart in the north Western Mediterranean Sea and Blue anchovy from the Central Adriatic Sea. Likewise, the genetic similarities between Blue anchovy populations throughout the northeastern Atlantic and Mediterranean Sea were greater than between Blue anchovy and Silver anchovy within, e.g., the Adriatic Sea. This indicated restricted gene flow at a local scale in spite of a high level of genetic exchange on a regional level. Even though no diagnostic allozyme locus has been detected between the two forms, the patterns of genetic clustering were correlated with habitat (offshore vs. inshore) and not with geography.

Blue anchovy from Mersin Bay were genetically more similar to *E. encrasicolus* from the Western Mediterranean Sea than from typical *E. albidus*. Silver anchovy from Mersin Bay were genetically closer to *E. albidus* from the Western Mediterranean Sea, albeit slightly so, than they were to typical *E. encrasicolus*. In the absence of linkage data, one may argue that the two CK intron markers scored by Bouchenak-Khelladi et al. (2008) could well be linked, hence additional markers would be welcome to enforce the genetic distinction between the two species. Because allozyme genes encode enzymes involved in metabolism, one may also argue that genetic variation at allozyme loci in anchovies from the Western Mediterranean (Pasteur and Berrebi, 1985) may also have been potentially affected by selection. Here, the result that Blue anchovy from Mersin Bay are *E. Encrasicolus* and Silver anchovy from Mersin Bay are *E. albidus* was based on two sets of nuclear markers including microsatellite markers (see Fig. 5b). The latter are a priori unlikely to be linked to the two intron markers scored by Bouchenak-Khelladi et al. (2008), or to the six polymorphic allozyme loci scored by Pasteur and Berrebi (1985).

The difference between Mersin Bay Blue anchovy and *E. encrasicolus* was much higher than the allozyme-based estimate of genetic differentiation reported among *E. encrasicolus* populations

[i.e., Group II of Borsa (2002)] Mediterranean-wide. It is possible that Mersin-Bay Blue anchovy actually belong to another population, e.g. Group III of Borsa (2002), which includes samples from Sicily and from the Aegean Sea. The genetic difference, at nuclear DNA loci, between Mersin-Bay Silver anchovy and *E. albidus* was much higher than the (allozyme-based) comparison between *E. albidus* from the Western Mediterranean Sea and Silver anchovy from the northern Adriatic Sea (Borsa, 2002). This last result can be interpreted in terms of genetic differentiation between geographically distant populations.

5. Conclusion

Species are independently evolving metapopulation lineages (de Queiroz, 2007). Expected properties of separate species are distinct morphologies and distinct genetic composition. The occurrence of genetically distinct forms in sympatry provides evidence for reproductive isolation, which is another expected property of separate species. Here, we demonstrated both morphological distinction and significant, substantial genetic differences between anchovy forms at a single site, Mersin Bay. In our view, this is evidence for two separate species. These two anchovy forms also seem to segregate according to habitat (i.e., inshore vs. offshore), which is a further expected property of separate species (de Queiroz, 2007).

The present findings provide new insights into the biogeography of a common, but, from a taxonomic viewpoint, still incompletely described fish of the northeastern Atlantic and Mediterranean waters. Future studies on the biology and ecology of anchovy populations will have to take into account the occurrence of two genetically and morphologically differentiated forms that occur sympatrically in several regions in both the Western and the Eastern Mediterranean basins and possibly elsewhere. This has further implications regarding the fishery management of anchovy, and the conservation of important biodiversity areas, such as Mersin Bay.

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Table 1. Summary of genetic diversity at 2 intron- and 10 microsatellite loci in European anchovy (*Engraulis* spp.) from Mersin Bay and from reference samples. *Ann*, Annaba, south Western Mediterranean Sea; *Ben*, Benguela current system off South Africa; *Cul*, Cul-de-Beauduc, north Western Mediterranean Sea; *Set*, off Sète, north Western Mediterranean Sea (sampling details for reference samples in Bouchenak-Khelladi et al., 2008). *N*, sample size; *k*, number of alleles; *AR*, allelic richness; *He*, expected heterozygosity; *Ho*, observed heterozygosity; *f*, inbreeding coefficient; *Null* frequency of null alleles under the hypothesis of the Hardy-Weinberg equilibrium.

Sample, parameter	Locus											
	<i>CK1</i>	<i>CK2</i>	<i>EeOV04</i>	<i>Ee-10</i>	<i>Ee2-91a</i>	<i>Ee2-91b</i>	<i>Ee2-135</i>	<i>Ee2-407</i>	<i>Ee2-452a</i>	<i>Ee2-508</i>	<i>Ej-27.1</i>	<i>Ej-41.1</i>
Silver (<i>N</i> =48)												
<i>k</i>	5	8	26	19	12	7	15	18	15	13	23	12
<i>AR</i>	2.482	4.468	8.908	7.516	6.506	5.025	5.881	6.484	7.564	7.757	8.678	6.281
<i>Ho</i>	0.375	0.604	0.681	0.854	0.745	0.750	0.674	0.833	0.830	0.688	0.468	0.896
<i>He</i>	0.514	0.675	0.936	0.891	0.868	0.795	0.808	0.827	0.900	0.912	0.934	0.851
<i>f</i>	0.273	0.105	0.275*	0.042	0.143	0.057	0.167	-0.008	0.079	0.248*	0.502*	-0.054
<i>Null</i>	0.08	0.034	0.128	0.000	0.048	0.012	0.074	0.007	0.040	0.116	0.237	0.000
Blue (<i>N</i> =48)												
<i>k</i>	8	12	24	20	13	7	13	12	17	9	27	10
<i>AR</i>	3.622	5.495	8.955	6.211	6.850	5.007	5.506	6.216	7.520	4.980	9.606	6.532
<i>Ho</i>	0.5	0.521	0.804	0.813	0.833	0.792	0.617	0.771	0.833	0.792	0.457	0.958
<i>He</i>	0.617	0.746	0.941	0.755	0.885	0.794	0.823	0.807	0.901	0.745	0.957	0.873
<i>f</i>	0.192	0.304*	0.147	-0.078	0.059	0.003	0.252*	0.045	0.076	-0.064	0.526*	-0.099
<i>Null</i>	0.083	0.133	0.073	0.007	0.024	0.012	0.113	0.012	0.029	0.000	0.251	0.000
Ann (<i>N</i> =15)												
<i>k</i>	2	7	19	13	10	10	9	14	10	11	21	10
<i>AR</i>	1.683	4.972	9.832	7.848	7.251	5.928	6.090	7.473	7.412	7.007	10.778	7.057
<i>Ho</i>	0.143	0.533	0.929	1.000	0.929	0.867	0.533	0.667	0.933	0.867	0.857	0.867
<i>He</i>	0.137	0.786	0.958	0.897	0.905	0.816	0.846	0.878	0.910	0.878	0.982	0.883
<i>f</i>	-0.04	0.329	0.032	-0.12	-0.027	-0.064	0.378	0.247	-0.026	0.014	0.131	0.019
<i>Null</i>	0	0.128	0.000	0.000	0.000	0.000	0.139	0.067	0.000	0.016	0.048	0.000
Ben (<i>N</i> =15)												
<i>k</i>	3	6	7	10	11	7	5	10	10	9	16	8
<i>AR</i>	2.705	4.386	4.860	6.671	7.381	5.840	4.017	7.132	7.057	6.502	9.582	6.435
<i>Ho</i>	0.4	0.467	0.786	0.733	0.933	0.714	0.600	0.733	0.857	0.867	0.643	0.733
<i>He</i>	0.358	0.72	0.688	0.855	0.903	0.855	0.722	0.899	0.889	0.878	0.960	0.881
<i>f</i>	-0.125	0.360	-0.149	0.147	-0.034	0.169	0.174	0.189	0.037	0.014	0.339*	0.172
<i>Null</i>	0	0.122	0.000	0.079	0.000	0.045	0.055	0.064	0.000	0.000	0.151	0.073
Cul(<i>N</i> =7)												
<i>k</i>	2	2	10	10	6	5	6	6	4	6	9	6
<i>AR</i>	2.000	2.000	9.099	8.978	5.692	4.835	5.429	5.549	4.000	5.549	8.253	5.692
<i>Ho</i>	1.000	0.167	0.857	1.000	0.571	0.429	0.857	0.571	0.667	0.714	0.429	0.714
<i>He</i>	0.538	0.410	0.956	0.945	0.835	0.780	0.780	0.747	0.636	0.747	0.934	0.846
<i>f</i>	-1.000	0.615	0.111	-0.063	0.333	0.471	-0.108	0.25	-0.053	0.048	0.561*	0.167
<i>Null</i>	0.000	0.168	0.000	0.000	0.112	0.177	0.000	0.082	0.000	0.000	0.236	0.000
Set (<i>N</i> =15)												
<i>k</i>	1	8	14	14	7	6	8	10	10	7	18	8
<i>AR</i>	1.000	5.602	7.799	7.441	5.545	4.395	5.121	6.470	7.328	5.264	9.760	6.08
<i>Ho</i>	0	0.786	0.733	0.867	0.923	0.933	0.467	0.643	0.800	0.750	0.714	0.733
<i>He</i>	-	0.762	0.901	0.837	0.822	0.724	0.756	0.852	0.906	0.754	0.960	0.862
<i>f</i>	0.001	-0.033	0.192	-0.037	-0.129	-0.302	0.391	0.252	0.12	0.005	0.263	0.154
<i>Null</i>	NA	0	0.078	0.000	0.000	0.000	0.157	0.058	0.048	0.000	0.104	0.034

* Value significant after Bonferroni correction for multiple tests (Rice, 1989).

Table 2. Mediterranean anchovy, *Engraulis* spp. Pairwise genetic differentiation (θ) estimates between populations, based on 2 intron- and 9 microsatellite loci.

	Sample				
Sample	Blue	Ann	Ben	Cul	Set
Silver	0.041*	0.038*	0.066*	0.063*	0.073*
Blue	0	0.043*	0.087*	0.094*	0.057*
Ann		0	0.051*	0.057*	0.017*
Ben			0	0.092*	0.094*
Cul				0	0.095*

*Value significant after Bonferroni correction for multiple tests (Rice, 1989)

LEGENDS TO FIGURES

Fig. 1. Trawling locations for anchovy (*Engraulis* spp.) in Mersin Bay, Eastern Mediterranean Sea (36°31'N-36°48'N; 34°13'E-34°44'E) in June 2010. Pie charts indicate catch ratios of Silver and Blue anchovies, as determined from external appearance; *grey*, Silver anchovy; *blue*, Blue anchovy.

Fig. 2. Body shape and colour differences between the two anchovy forms (*Engraulis* spp.) from Mersin Bay, Eastern Mediterranean Sea, and landmarks for geometric morphometrics. **(a)** Anchovy of the “Blue” phenotype typical of the deeper waters of Mersin Bay; *scale*: 1 square = 0.2 cm. **(b)** Anchovy of the “Silver” phenotype typical of the shallow inshore habitat of Mersin Bay; *scale*: 1 square = 0.2 cm. **(c)** Position of 20 morphometric landmarks (black dots): 1, tip of premaxillary; 2, pit of upper jaw; 3, border of head; 4, anterior base of the dorsal fin; 5, posterior base of the dorsal fin; 6, 7, 8 points of maximum curvature of the caudal peduncle; 9, posterior base of anal fin; 10, anterior base of anal fin; 11, base of pelvic fin; 12, base of pectoral fin; 13, ventral tip of cleithrum; 14, tip of lower jaw; 15, anterior border of eye; 16, posterior border of eye; 17, most anterior point of cleithrum; 18, dorsal tip of cleithrum; 19, 20, insertions of pectoral fin. **(d)** Pattern of shape change along PC1 of principal component analysis (see Fig. 2e). Circles indicate the locations of the landmarks in the mean shape of the sample; arrows indicate the changes in the relative positions of the landmarks as the score on PC1 increases (scaling factor 0.1). **(e)** Principal component analysis of body shape for 229 Silver and Blue anchovy specimens from Mersin Bay, based on 20 landmarks (Fig. 2c). *Closed triangles* (▲): male Silver anchovy; *inverted open triangles* (▼): female Silver anchovy; *open squares* (□): male Blue anchovy; *closed squares* (■): female Blue anchovy. **(f)** Cross-validation discriminant function analysis of Blue and Silver anchovy body shape. Rate of correct classification was 95.6% for Silver anchovy and 96.8% for Blue anchovy. *Blue*: Blue anchovy; *grey*: Silver anchovy; *dark blue*: misclassified individuals.

Fig. 3. Otolith morphometrics of Mersin Bay *Engraulis* spp. **(a)** Photographed are the sagittae of 2+ year-old Silver anchovy (left) and Blue anchovy (right). Sagittae traced by 120 equally spaced points using the TPSDig2 computer program (Rohlf, 1993). *Load*: points of 120 landmarks around otolith (blue contour); *H:1, H:2 and H:21* harmonic numbers (red contour) from EFAWin program. Twenty one harmonics were used to obtain a balance between ratio of sample size to the number of variables, and accuracy of the description of otolith's shape. **(b)** Principal component analysis on 21 harmonics of sagitta contour for 1+ Blue ($N=15$) and Silver ($N=13$) individuals. **(c)** Principal component analysis on 21 harmonics of sagitta contour for 2+ Blue ($N=7$) and Silver ($N=8$) individuals. *Black circles* (●): Silver anchovy; *diamonds* (◊): Blue anchovy.

Fig. 4. Results of Bayesian structure analysis of individual genotypes in anchovy samples (*Engraulis* spp.) from Mersin Bay (Silver, Blue) together with reference samples from the Benguela current system (Ben) and from the Western Mediterranean Sea (Ann, Set, Cul), based on 11 nuclear loci (*EJ27.1* excluded) assuming $K=2$ clusters. Twenty replicate simulations were run for each K value,

ranging from 2 to 7, using 10,000 burn-in repetitions and a final run of 100,000 Markov chain-Monte Carlo steps under an admixture model with independent allele frequencies. Evanno et al.'s (2005) test designated $K=2$ as the most likely (see Supplementary material, Figs. S2 and S3). Each individual, represented by a vertical bar, is segmented with assorted colours representing the different clusters to which it was assigned, with segment length proportional to assignment probability. Figure drawn using Distruct v1.1 (Rosenberg, 2004).

Fig. 5. Genetic relationships of Mersin-Bay *Engraulis* spp. (*Silver* and *Blue*) to samples from the Atlantic and the Mediterranean as inferred from correspondence analysis (CA) (FACTOMINER: Lê et al., 2008). Ellipses delineate clusters of samples determined by hierarchical clustering (FACTOMINER). *Grey lettering*: samples set as inactive in the analyses (see Appendix B for justification). *In brackets*: inertias of axes. **(a)** Samples of the present study, together with those of Bouchenak-Khelladi et al. (2008), including reference samples for *E. encrasicolus* (*SET*) and *E. albidus* (*CUL*), and those of Kristofferson and Magoulas (2008). Samples were characterized by their compound-allele frequencies at intron loci *CK6-1* and *CK6-2* (Appendix B). **(b)** Samples of the present study, together with those of Borrell et al. (2012), characterized by their compound-allele frequencies at microsatellite loci *EeOV04*, *Ee-10* and *Ee2-508* (Appendix C).

Figure1

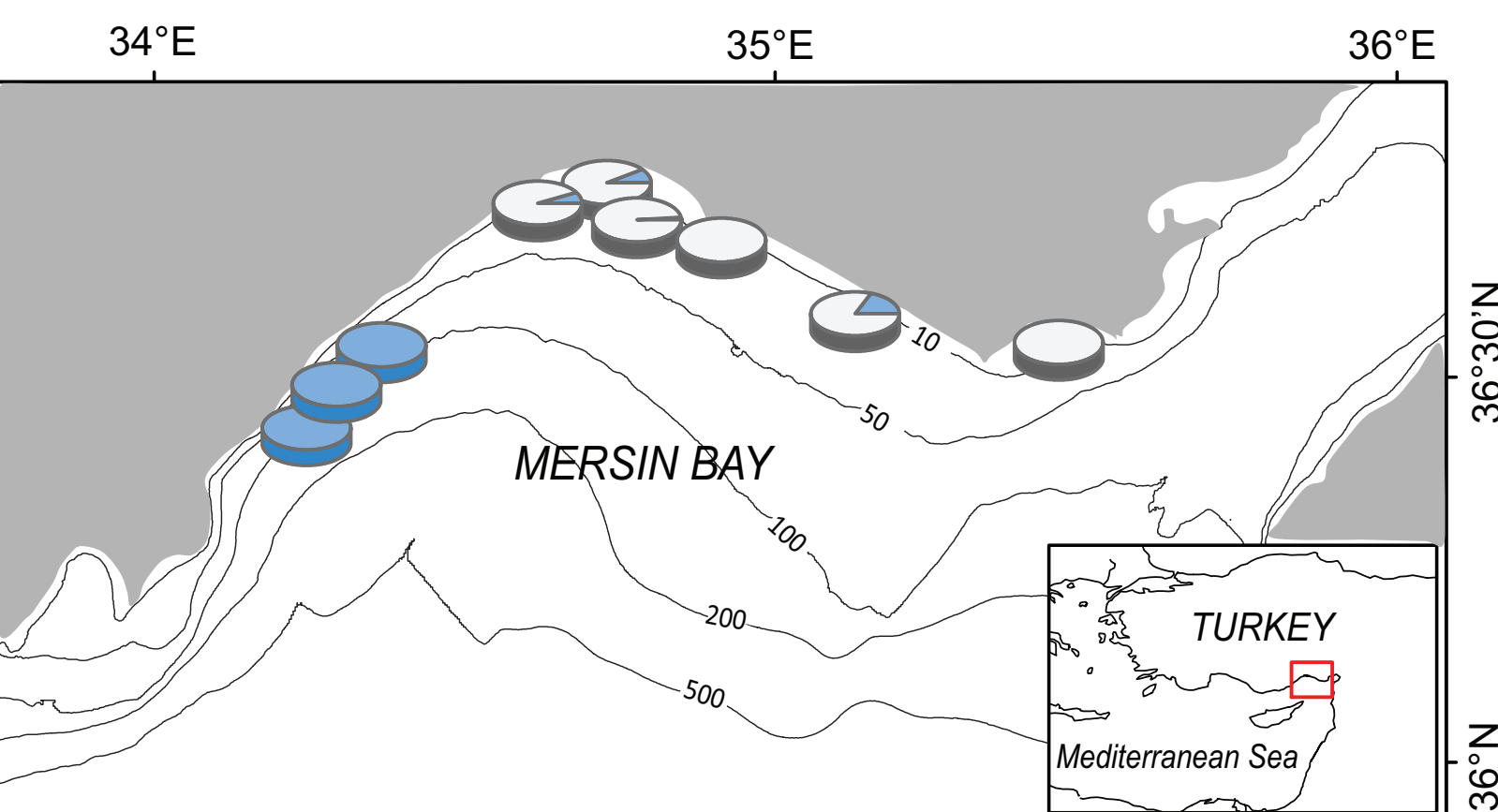


Figure2

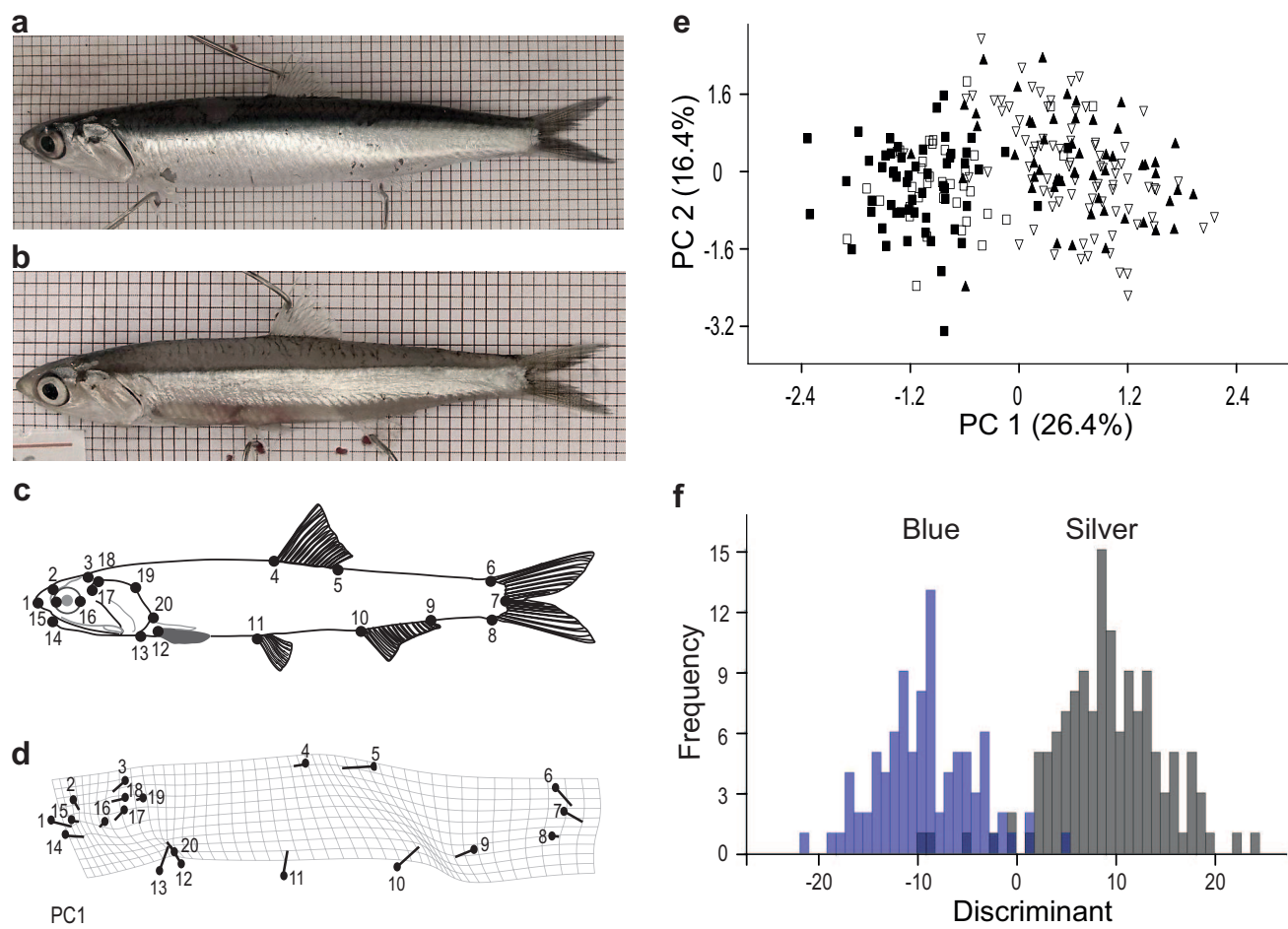


Figure3

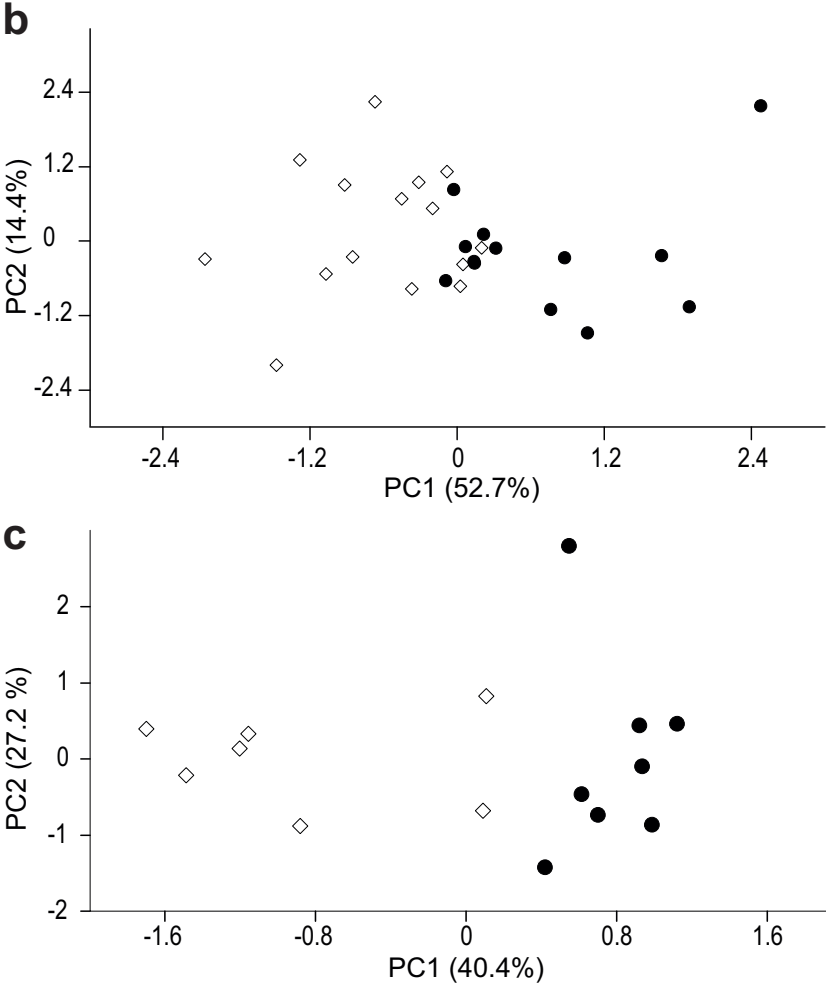
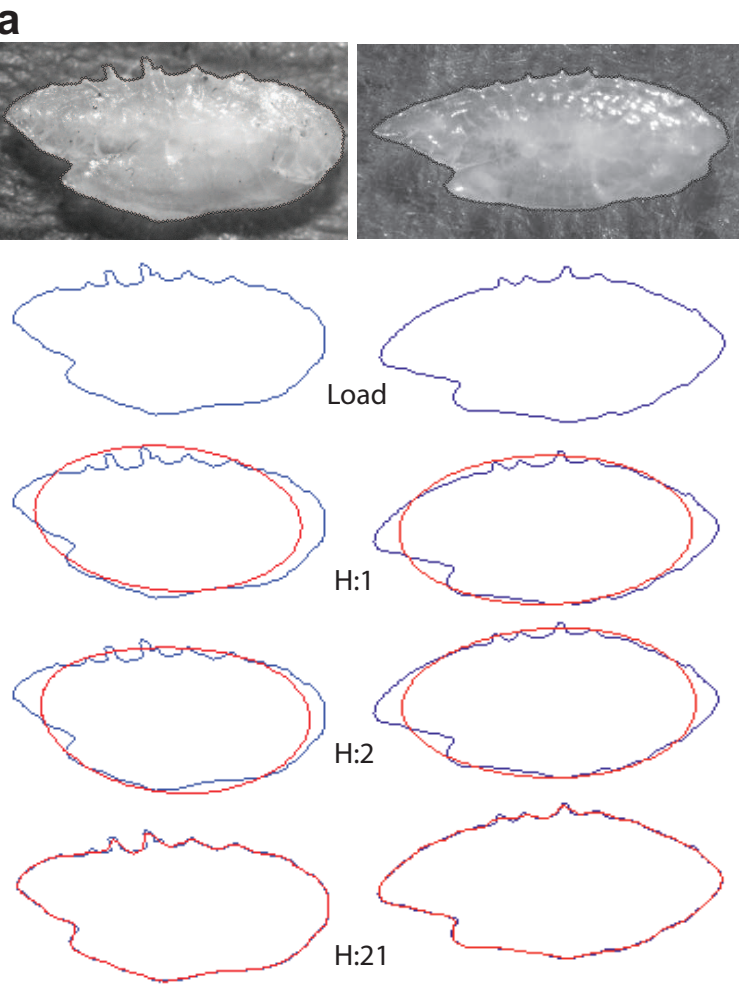


Figure4

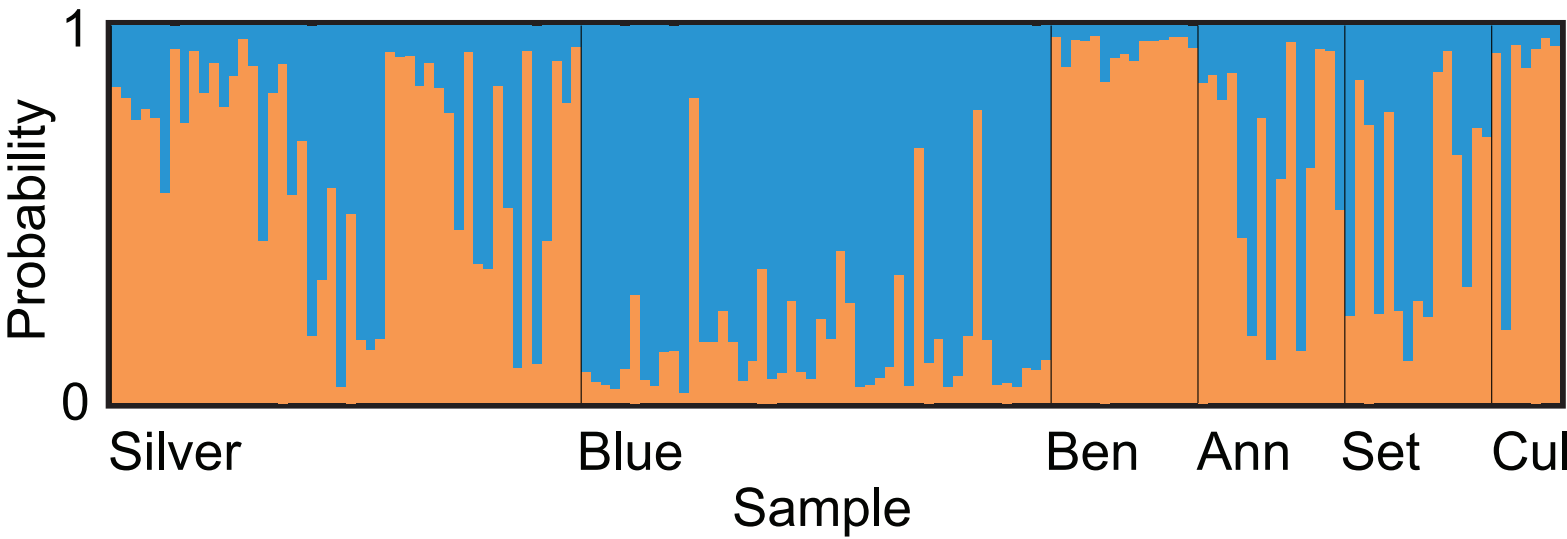
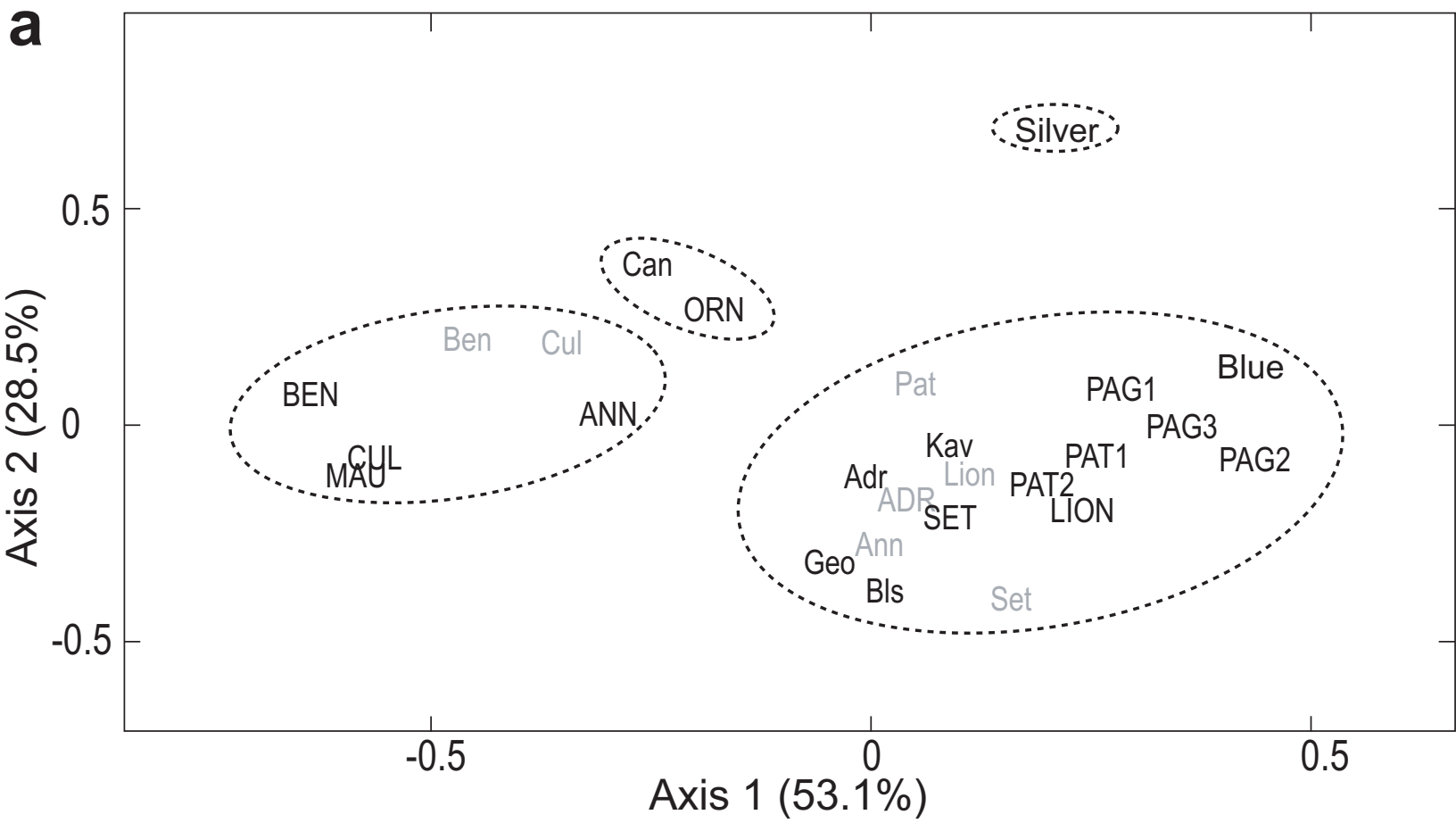
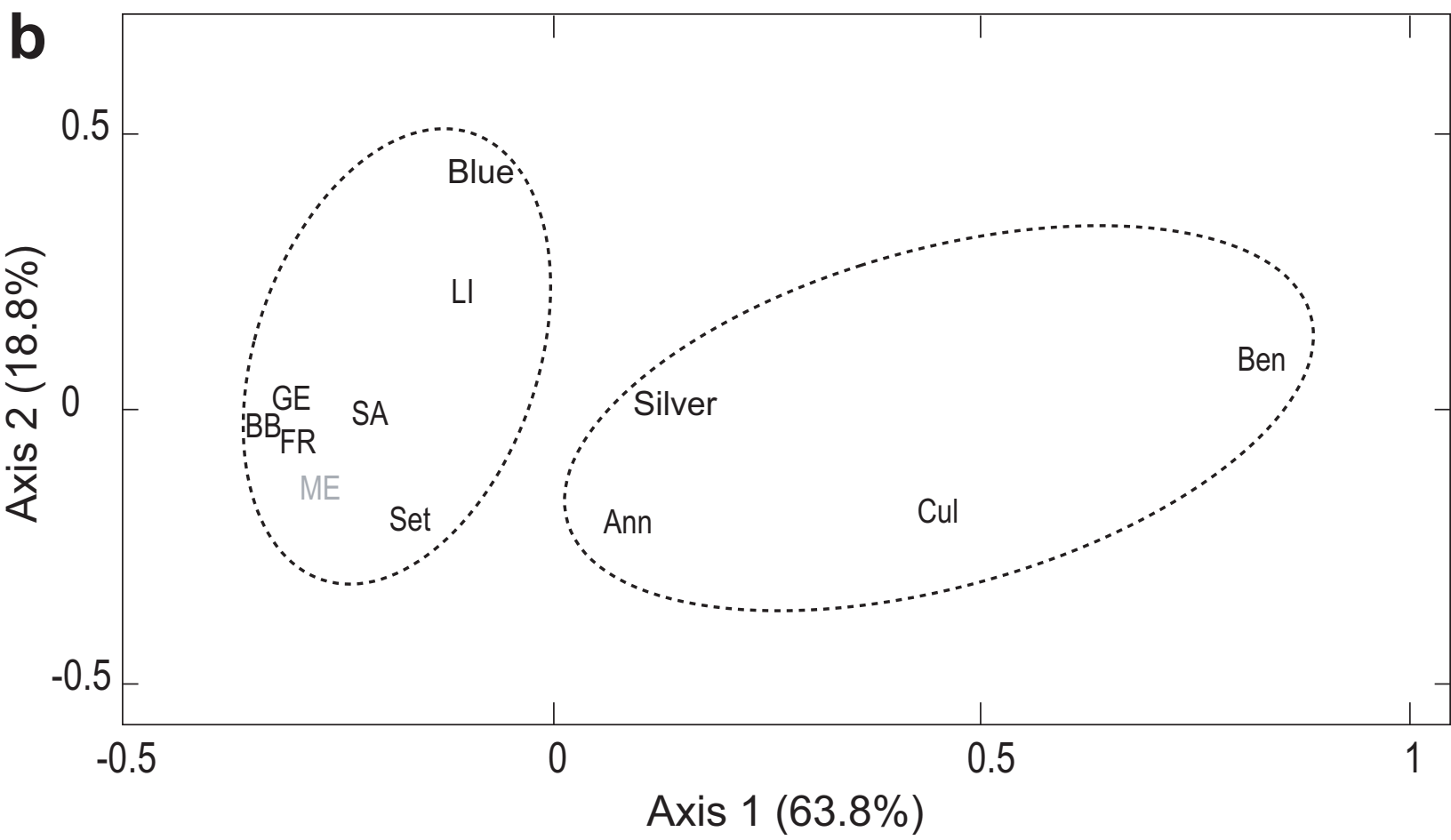


Figure5

a



b



Appendix A

Primers and PCR conditions for two intron and 10 microsatellite loci in European anchovy

Locus	Repeat motif (5'-3')	Forward (F) and reverse (R) primer sequences	T _a (°C)	[Mg ²⁺] (mM)	Allele size (bp)	Reference
<i>CK6-1</i>	-	F 5'-CGACATTGTAATGATGTTACAATGA-3' R 5'-ATTTCTTTTGGGTTGGCTCTTCTCT-3'	54	1.7	320-1040	Borsa et al., 2004
<i>CK6-2</i>	-	F 5'-CTCAGAACTACATACCAAACCAATG-3' R 5'-ACTCACTGTAATTCTGAATAGAGCT-3'	52	1.7	472-527	Borsa et al., 2004
<i>EeOV04</i>	(CA) ₂	F 5'-CCCAGGAAGTGTATTGGGTGGCTT-3' R 5'-TGTGTGTCCCAAACACAGTGCCTGA-3'	56	1.5	275-400	Borrell et al., 2011
<i>Ee-10</i>	[(GT) ₉ CT] ₂ [(GT) ₂ CT] ₃	F 5'-GGTGGATGAAGTGGCAATCT-3' R 5'-CTGGGGTGGCATAAAGTGAAG-3'	56	1.5	200-268	Landi et al., 2005
<i>Ee2-91a</i>	(AGG) ₁₂	F 5'-AGAGCAGGTTCTTGTGTGG-3' R 5'-TGTGGTGGCTACTATCAGG-3'	55	1.5	250-292	Pakaki et al., 2009
<i>Ee2-91b</i>	(CCGCA) ₈	F 5'-GGTCTTGAGCTTGGCATAGG-3' R 5'-CCGGAAGACACTCTGCACAC-3'	60	1.5	115-170	Pakaki et al., 2009
<i>Ee2-135</i>	(ATTAG) ₁₀	F 5'-AGGGCAGTGACAGGAGAGTC-3' R 5'-TCGTTACCTGCGTTTATACTG-3'	55	1.5	110-165	Pakaki et al., 2009
<i>Ee2-407</i>	(CA) ₁₃	F 5'-AGGAATCTCCTTCCCGTCTC-3' R 5'-GTGGGTCTGTGGGTGTTTG-3'	60	1.5	136-196	Pakaki et al., 2009
<i>Ee2-452</i>	(AC) ₁₃	F 5'-CCCAACCCTAGGGAGACATC-3' R 5'-TCGTTTCAGCAAGCATACACC-3'	59	1.5	250-298	Pakaki et al., 2009
<i>Ee2-508</i>	(AGG) ₈	F 5'-CACATGCTCGCTAAACATTG-3' R 5'-CCTGATGCTGCTTGGTAGC-3'	55	1.5	154-199	Pakaki et al., 2009
<i>Ej-27.1</i>	(GA) ₃₆	F 5'-GACTGTGAAGGAACGCTGGT-3' R 5'-AATAGGATTAGTCATCACAGGG-3'	56	1.5	150-268	Chiu et al., 2002
<i>Ej-41.1</i>	(CACAA) ₈	F 5'-TCTACCCCTGGAGGACACAC-3' R 5'-ACAGGGGGTTGAGAAAGAGG-3'	55	1.5	110-175	Chiu et al., 2002

Appendix B

Comparison of allele frequencies at intron loci *CK6-1* and *CK6-2* for the samples analyzed in this study with those in Bouchenak-Khelladi et al. (2008) (*Bouchenak-Khelladi*) and, indirectly, Kristofferson and Magoulas (2008) (*Kristofferson*)^a. *S*, *O*, *F*, compound alleles as defined in footnotes^{b,c}. *N* sample size.

Locus,	Sample																										
Allele	Bouchenak-Khelladi ^a														Kristofferson ^a						This study ^a						
	Adr	ANN	BEN	Bls	Can	CUL	Geo	Kav	MAU	Lion	ORN	Pat	SET	ADR	LION	PAG1	PAG2	PAG3	PAT1	PAT2	Ann	Ben	Blue	Cul	Set	Silver	
<i>CK6-1</i> ^b																											
<i>S</i>	0	0.01	0	0.02	0	0	0	0.02	0	0	0	0.02	0.01	0.01	0	0.01	0.03	0.02	0	0	0	0	0.08	0	0	0.02	
<i>O</i>	0.74	0.86	0.75	0.94	0.61	0.93	0.95	0.78	0.93	0.93	0.58	0.73	0.85	0.78	0.86	0.71	0.65	0.73	0.85	0.83	0.93	0.80	0.54	0.50	1	0.60	
<i>F</i>	0.26	0.13	0.24	0.04	0.39	0.07	0.05	0.21	0.06	0.07	0.41	0.25	0.14	0.20	0.14	0.28	0.33	0.25	0.15	0.17	0.07	0.20	0.38	0.50	0	0.38	
(N)	(52)	(49)	(49)	(56)	(49)	(112)	(55)	(58)	(46)	(15)	(24)	(53)	(95)	(37)	(47)	(117)	(40)	(83)	(83)	(82)	(15)	(15)	(48)	(7)	(15)	(48)	
<i>CK6-2</i> ^c																											
<i>S</i>	0.47	0.17	0.05	0.51	0.14	0.10	0.48	0.46	0.10	0.43	0.25	0.36	0.50	0.49	0.55	0.50	0.67	0.55	0.51	0.54	0.47	0.03	0.57	0.25	0.61	0.14	
<i>O</i>	0.09	0.24	0.07	0.10	0.25	0.13	0.12	0.18	0.12	0.30	0.17	0.26	0.16	0.10	0.23	0.28	0.15	0.27	0.30	0.21	0.17	0.27	0.20	0	0.14	0.64	
<i>F</i>	0.44	0.58	0.87	0.39	0.6	0.77	0.41	0.36	0.78	0.27	0.58	0.38	0.34	0.41	0.21	0.22	0.17	0.18	0.19	0.25	0.37	0.70	0.23	0.75	0.25	0.23	
(N)	(46)	(49)	(48)	(50)	(48)	(115)	(53)	(36)	(46)	(15)	(24)	(29)	(93)	(43)	(47)	(114)	(46)	(103)	(92)	(78)	(15)	(15)	(48)	(7)	(15)	(48)	

^a Samples Lion, Adr and Pat of Bouchenak-Khelladi et al. (2008) are partly the same as samples LION, ADR and either PAT1 or PAT2 respectively of Kristofferson and Magoulas (2008); they were used to determine allele homologies across the two studies. Similarly, samples Cul, Set, Ann and Ben of this study are subsamples of, CUL, SET, ANN and BEN respectively of Bouchenak-Khelladi et al. (2008) and were used for the same purpose. To eliminate duplications, samples Lion, Pat, ADR, Cul, Set, Ann and Ben were defined as inactive elements in the CA (Fig. 5a)

^b At locus *CK6-1*, compound allele *S* corresponds to alleles 120, 110 and 105 of Bouchenak-Khelladi et al. (2008), 1-3 of Kristofferson and Magoulas (2008) and 990, 940, 615 and 563 of this study; allele *O* corresponds to allele 100 of Bouchenak-Khelladi et al. (2008), 4 of Kristofferson and Magoulas (2008), and 524 of this study; compound allele *F* corresponds to alleles 98, 95, 90, 52, c050, 47, 30, 10 and 1 of Bouchenak-Khelladi et al. (2008), 5-21 of Kristofferson and Magoulas (2008), and 444, 414, 412, 404 and 320 of this study

^c At locus *CK6-2*, compound allele *S* corresponds to alleles 150, 110, 107 and c105 of Bouchenak-Khelladi et al. (2008), 1-2 of Kristofferson and Magoulas (2008), and 527, 525 and 523 of this study; compound allele *O* corresponds to alleles c102 of Bouchenak-Khelladi et al. (2008), 3-5 of Kristofferson and Magoulas (2008), and 521 and 518 of this study; compound allele *F* corresponds to alleles 100, 98, 97, 95, 90 and 10 of Bouchenak-Khelladi et al. (2008), 6-14 of Kristofferson and Magoulas (2008), and 516, 514, 512, 510, 508, 504, 502, 474 and 472 of this study.

Appendix C

Comparison of allele frequencies at microsatellite loci *EeOV04*, *Ee-10* and *Ee2-508* for the samples analyzed in this study, with those in Borrell et al. (2012) (*Borrell*). *N* sample size.

Locus, Allele	Sample						This study ^a					
	Borrell ^a											
	BB	FR	GE	LI	ME	SA	Ann	Ben	Blue	Cul	Set	Silver
<i>EeOV04</i> ^b												
>306	0.45	0.51	0.64	0.54	0.55	0.55	0.54	0.36	0.59	0.36	0.37	0.60
<i>c304</i>	0.33	0.31	0.17	0.10	0.22	0.30	0.25	0	0.21	0.21	0.20	0.22
<300	0.22	0.18	0.19	0.36	0.23	0.15	0.25	0.64	0.26	0.50	0.43	0.21
(N)	(181)	(39)	(21)	(25)	(43)	(48)	(14)	(14)	(46)	(7)	(15)	(47)
<i>Ee-10</i> ^c												
>214	0.75	0.74	0.76	0.58	0.84	0.69	0.64	0.17	0.34	0.50	0.83	0.57
214	0.20	0.18	0.12	0.30	0.10	0.22	0.04	0.30	0.48	0.14	0.03	0.23
<214	0.05	0.09	0.12	0.12	0.06	0.09	0.32	0.53	0.18	0.36	0.13	0.20
(N)	(182)	(40)	(21)	(25)	(44)	(48)	(14)	(15)	(48)	(7)	(15)	(48)
<i>Ee2-508</i> ^d												
>163	0.41	0.45	0.36	0.44	0.50	0.56	0.67	0.97	0.43	0.93	0.46	0.79
163	0.51	0.49	0.50	0.44	0.46	0.39	0.27	0	0.44	0.07	0.46	0.07
<163	0.07	0.06	0.14	0.12	0.04	0.04	0.07	0.03	0.14	0	0.08	0.14
(N)	(182)	(40)	(21)	(24)	(42)	(47)	(15)	(15)	(48)	(7)	(12)	(48)

^a To establish identities of alleles across studies, the Blue and Set samples of the present study were compared with sample ME of Borrell et al. (2011), which was located in the southern Central Adriatic and, hence was assumed to consist of Blue anchovy (Bembo et al., 1996a).

^b The most common allele at locus *EeOV04* in Blue anchovy (this study) was *304* (frequency = 0.14). The most common allele in sample ME of Borrell et al. (2012) was *305* (frequency = 0.16), indicating that the sizing of alleles in Borrell et al. (2012) was shifted by one bp to the right, relative to this study. All alleles larger than *306* (*307* of Borrell et al., 2012) were lumped into compound allele >*306* while all alleles smaller than *300* (*301* of Borrell et al., 2012) were lumped into compound allele <*300*. All alleles from *300* to *306* (from *301* to *307* in Borrell et al., 2012) were lumped into compound allele *c304*

^c The most common allele at locus *Ee-10* in Blue anchovy (this study) was *214* (frequency = 0.48) and was assumed to be homologous to allele *215* of Borrell et al. (2012). All alleles larger (resp. smaller) than *214* (*215* in Borrell et al., 2012) were lumped into compound allele >*214* (resp. <*214*)

^d The most common allele at locus *Ee2-508* in Blue anchovy (this study) was *163* (frequency = 0.44) and was assumed to be homologous to allele *165* of Borrell et al. (2012). All alleles larger (resp. smaller) than *163* (*165* in Borrell et al., 2012) were lumped into compound allele >*163* (resp. <*163*)

Appendix D. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/xxx>

Supplementary material to:

Geometric morphometrics, Fourier analysis of otolith shape, and nuclear-DNA markers distinguish two anchovy species (Engraulis spp.) in the Eastern Mediterranean

by Arzu Karahan, Philippe Borsa, Ali Cemal Gucu, İrfan Kandemir, Emel Ozkan, Yesim Ak Orek, Sinan Can Acan, Evren Koban and Inci Togan

Table S1-S3 and Figs. S1-S3 here appended

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Table S1. Detection of outlier loci with respect to the distribution of *Fis* values estimated for two anchovy (*Engraulis* spp.) samples from Mersin Bay. To determine the outliers one interquartile range (IQR) (which is defined as the interval between the upper limits of the first and third quartiles) was added to the upper value of the third quartile, and subtracted from the upper value of the first quartile.

Locus	Silver anchovy			Blue anchovy		
	<i>Fis</i>	Rank	Quartile	<i>Fis</i>	Rank	Quartile
<i>CK6-1</i>	0.273	10	Q4	0.192	9	Q3
<i>CK6-2</i>	0.105	6	Q2	0.304	11	Q4
<i>EeOV04</i>	0.275	11	Q4	0.147	8	Q3
<i>Ee-10</i>	0.042	3	Q1	-0.078	2	Q1
<i>Ee2-91a</i>	0.143	7	Q3	0.059	6	Q2
<i>Ee2-91b</i>	0.057	4	Q2	0.003	4	Q2
<i>Ee2-135</i>	0.167	8	Q3	0.252	10	Q4
<i>Ee2-407</i>	-0.008	2	Q1	0.045	5	Q2
<i>Ee2-452a</i>	0.079	5	Q2	0.076	7	Q3
<i>Ee2-508</i>	0.248	9	Q3	-0.064	3	Q1
<i>Ej27.1</i>	0.502	12	outlier	0.526	12	outlier
<i>Ej41.1</i>	-0.054	1	minimum	-0.099	1	minimum
Lower limit	-0.175	-	-	-0.332	-	-
Upper limit	0.488	-	-	0.521	-	-

Table S2. Membership coefficients determined from Bayesian structure analysis (STRUCTURE: Falush et al., 2007) based on genotype data at two intron and 9 microsatellite loci, for two anchovy samples from Mersin Bay (i.e., samples Blue and Silver, which consisted, respectively, of all blue and silver individuals sampled from the bay), with prior number of populations (*K*) allowed to range from 2 to 4. Each Cluster is represented by a particular colour in the STRUCTURE diagrams (Fig. S2).

Sample	<i>K</i>	Cluster to which assigned			
		Cluster 1	Cluster 2	Cluster 3	Cluster 4
Blue	2	0.50	0.50	-	-
	3	0.43	0.12	0.45	-
	4	0.33	0.21	0.38	0.08
Silver	2	0.57	0.43	-	-
	3	0.27	0.49	0.24	-
	4	0.18	0.24	0.18	0.41

Table S3. Locus-pairwise linkage disequilibrium values [R_{ij} ; Weir (1979) in Belkhir et al. (2000)] in Silver anchovy (below diagonal) and Blue anchovy (above diagonal) from Mersin Bay, north Eastern Mediterranean. No significant value was detected (permutation tests and Bonferroni correction).

Locus	Locus										
	<i>CK6-1</i>	<i>CK6-2</i>	<i>EeOV04</i>	<i>Ee-10</i>	<i>Ee2-91a</i>	<i>Ee2-91b</i>	<i>Ee2-135</i>	<i>Ee2-407</i>	<i>Ee2-452a</i>	<i>Ee2-508</i>	<i>EJ41.1</i>
<i>CK6-1</i>	-	0.105	0.103	0.099	0.101	0.118	0.093	0.112	0.136	0.106	0.112
<i>CK6-2</i>	0.115	-	0.122	0.089	0.104	0.119	0.101	0.114	0.106	0.105	0.120
<i>EeOV04</i>	0.139	0.108	-	0.126	0.106	0.117	0.148	0.114	0.106	0.111	0.112
<i>Ee-10</i>	0.098	0.104	0.108	-	0.109	0.111	0.134	0.112	0.101	0.099	0.110
<i>Ee2-91a</i>	0.098	0.118	0.107	0.114	-	0.124	0.107	0.106	0.109	0.111	0.126
<i>Ee2-91b</i>	0.106	0.096	0.117	0.119	0.146	-	0.130	0.134	0.118	0.123	0.138
<i>Ee2-135</i>	0.083	0.143	0.119	0.121	0.147	0.119	-	0.103	0.104	0.108	0.156
<i>Ee2-407</i>	0.156	0.104	0.103	0.122	0.125	0.124	0.117	-	0.115	0.115	0.128
<i>Ee2-452a</i>	0.108	0.113	0.111	0.112	0.113	0.126	0.125	0.125	-	0.114	0.111
<i>Ee2-508</i>	0.096	0.126	0.110	0.114	0.123	0.126	0.115	0.108	0.127	-	0.123
<i>EJ41.1</i>	0.093	0.126	0.109	0.103	0.114	0.113	0.113	0.105	0.117	0.117	-

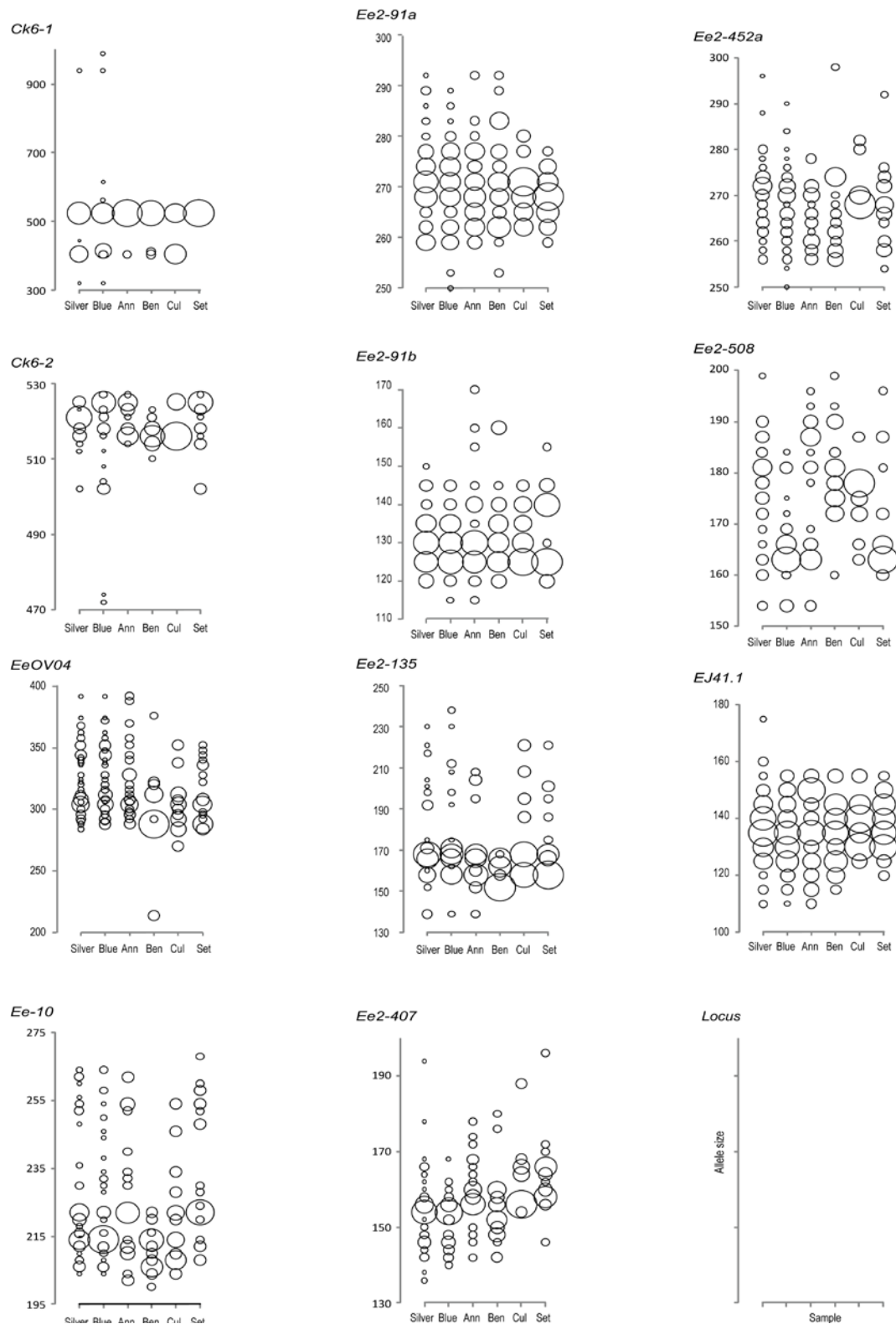


Figure S1. Anchovies, *Engraulis* spp. Size-allele frequencies at 11 nuclear loci in Silver and Blue anchovies from Mersin Bay, Eastern Mediterranean, and comparison with reference samples from the Western Mediterranean Sea (*Ann*, *Cul*, *Set*) and the southern Atlantic (*Ben*). Ordinate: allele size (in base pairs).

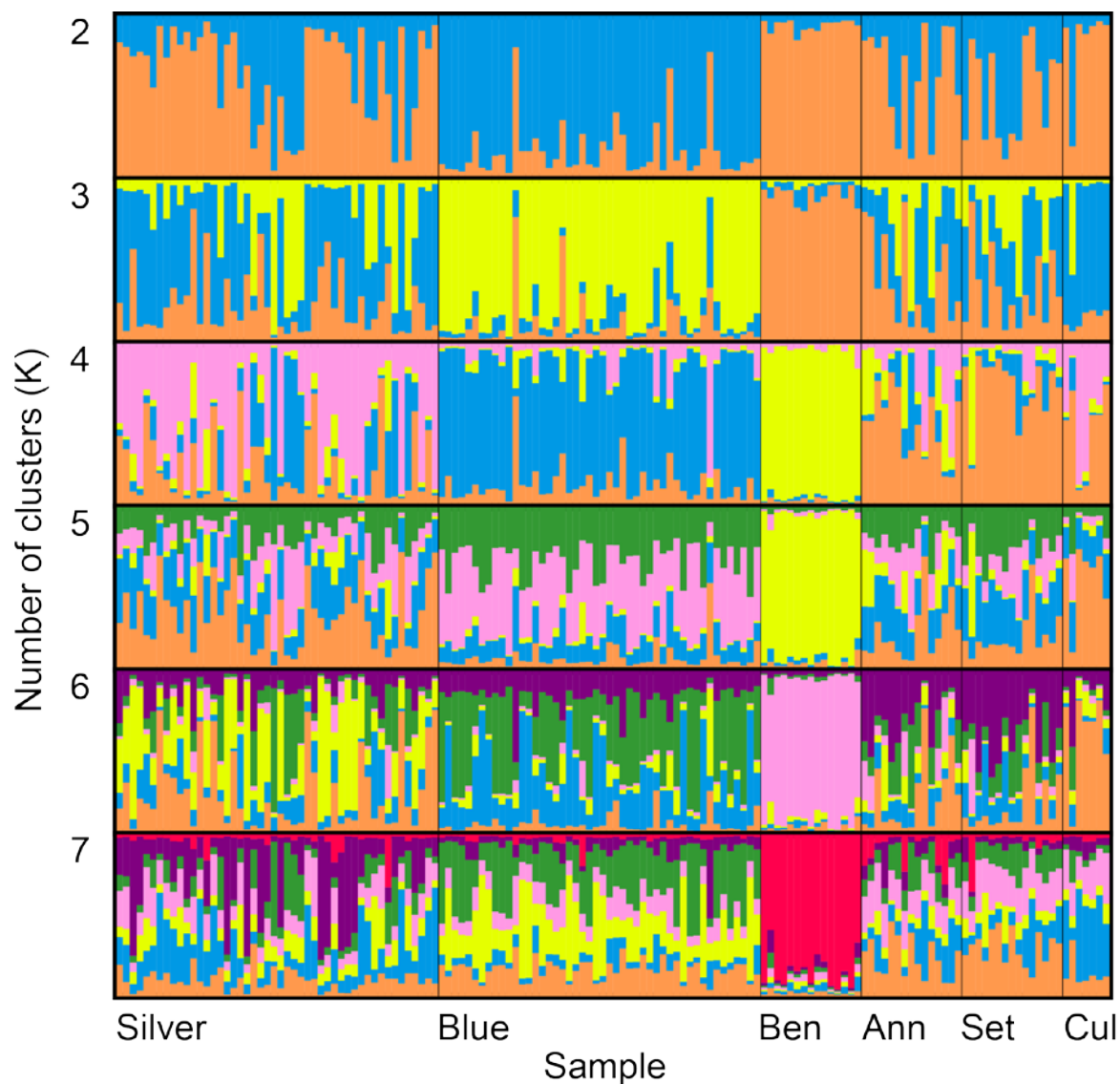


Fig. S2. Results of Bayesian structure analysis of individual genotypes in anchovy samples (*Engraulis* spp.) from Mersin Bay and four other samples from the Atlantic and the Western Mediterranean Sea, based on 11 nuclear loci (*EJ27.1* excluded) assuming $K=2$ to 7 Clusters. Ten replicate simulations were run for each K value, ranging from 2 to 7, using 10,000 burn-in repetitions and a final run of 100,000 Markov chain-Monte Carlo steps under an admixture model with independent allele frequencies. Each individual, represented by a vertical bar, is segmented with assorted colours representing the different clusters to which it was assigned, with segment length proportional to assignment probability. Figures drawn using Distruct v1.1 (Rosenberg, 2004). *Abbreviations for samples* as in Fig. S1.

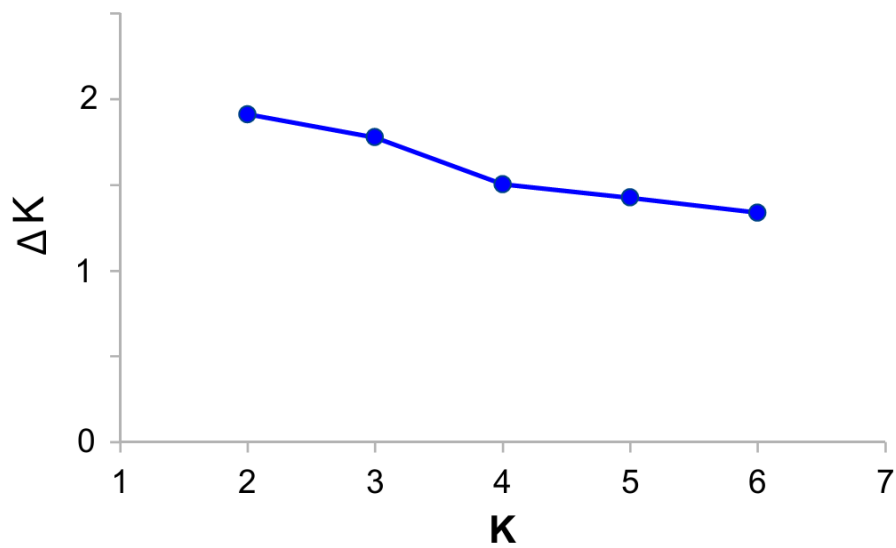


Fig. S3. Delta K values with respect to K, according to the calculation method by Evanno et al. (2005): $\Delta K = \text{mean}[|L''(K)|] / \text{sd}[L(K)]$.