

Genetic variability of Chilean and Peruvian surf clams (Donax marincovichi and Donax obesulus)

D. Carstensen, M. Herrmann, J. Laudien, S. Schiel, W. Arntz, F. Leese, C. Held

Introduction

Exposed intertidal sandy beaches are commonly dominated by surf clams of the genus Donax (Ansell 1983). In Peru and Chile these bivalves play an important role for artisanal fisheries. Within this genus, the taxonomic status of Donax marincovichi and D. obesulus, distributed along the Peruvian coastline, is controversially discussed (Coan 1983, Guzmán et al. 1998). As morphometric comparisons reveal no significant differences we possibly deal with a single, rather than with two species.

The aim of this study is twofold: First, we want to establish a molecular marker suitable for barcoding and providing evidence, based on sequence data, concerning this taxonomic controversy. Second, we want to estimate the genetic relatedness among geographically distant populations along the Peruvian coastline



D. marincovichi (23 mm

(17 mm



Fig. 1: Geographical distribution (------) and sampling localities (1-11) of *D. marincovichi* and *D* Chilean and Peruvian coast. Locality 12 showes the beach where *D. hanleyanus* was sampled.

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Material and Methods

Specimens of the two putative species D. marincovichi and D. obesulus were sampled at nine beaches along their distribution (S18°27'53.8"/W70°18'24.3" to S3°33'57"/W80°27'5"). Additionally, specimens of *D. asper* and *D. hanleyanus* were sampled as outgroups from one beach each. Opened and closed shells were conserved in >80% ethanol. DNA was extracted with the Qiagen Dneasy kit according to the manufacturer's recommendations. 10-100 ng DNA were used for PCR-amplification of a fragment of the cytochrome oxidase I (COI) using the primers HCO (5'-TAAACTTCAGGTGACCAAAAAATCA-3') and LCO (5'-GGTCAACAAATCATAAAGATATTGG-3') (Folmer et al. 1994) in 25 µl reactions. PCR products were purified using the Qiagen DNA purification kit according to the manufacturer's recommendations. Sequencing was conducted on an ABI 3730xl automated sequencer. Sequence data were processed and aligned using the ClustalW algorithm. Phylogenetic analysis was performed using PAUP4beta10.



Results and Discussion

Sequence data from the COI proved to be useful for species discrimination within the genus Donax: The taxonomic status of both species, D. asper and D. hanleyanus is well supported. However, there is no indication of reproductive isolation between D. marincovichi and D. obesulus in the COI data (Fig. 2a). With 0 - 1.2% sequence divergence, the divergence between D. marincovichi and D. obesulus is on the order of known intraspecific variability in the COI gene (Held 2000; and Fig. 2b). Therefore, the taxonomic status of the two species must be auestioned.

No genetic differentiation between the geographically separated D. marincovichi populations could be observed from the sequence data. In the future, molecular markers with higher resolving power (e.g. AFLP, Microsatellites) must be analysed for these puposes

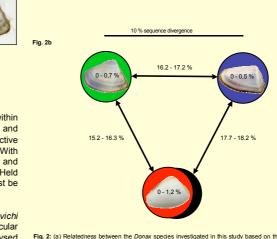


Fig. 2: (a) Relatedness between the Donax species investigated in this study based on the investigation of a 606 bp fragment of the militochondrial glochrome oxidase I. (b) interspecific and intraspecific (in cricles) genetic distances between D aspec, D harleyenas, D obsulus, and D marinovichih. D obsulus and D. marinovichih cannot be distinguished based on the COI data

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Alfred Wegener Institute for Polar and Marine Research Am Alten Hafen 26 27568 Bremerhaven : +49-471-48311315 tel: +49-471-48311315 fax: +49-471-48311918

Daniel Carstensen:

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mail: dears nsen@awi-bremerhaven.de