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Variola louti (Perciformes Epinephelidae) in the Mediterranean Sea: Incidental introduction or aquarium release?

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On the presence of the yellow-edged lyretail grouper, Variola louti (Perciformes: Epinephelidae), in the Mediterranean: Lessepsian immigrant or aquarium release?

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Abstract:	Lessepsian immigrants are altering the composition and functioning of Eastern Mediterranean ecosystems. Here, we report the first confirmed and second published record of the yellow-edged lyretail grouper, Variola louti (Forsskål, 1775) in the Mediterranean Sea and Cyprus, supported by morphological and and genetic analysis. Phylogenetic analyses revealed that none of the samples from the Red Sea or the Indian Ocean (Mozambique, South Africa and India), clustered with our samples; indicating that aquarium release is the most possible pathway.

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- 1 On the presence of the yellow-edged lyretail grouper, *Variola louti* (Perciformes: Epinephelidae), in
- 2 the Mediterranean: Lessepsian immigrant or aquarium release?
- 3 Short title: Variola louti in Cyprus and Mediterranean
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11 Introduction

Lessepsian immigrants (i.e. Red Sea species entering the Mediterranean through the Suez Canal) have greatly altered the composition and functioning of many Eastern Mediterranean shelf ecosystems and are expected to continue doing so, as new species arrive and establish self-sustaining populations in the region (Michailidis et al., 2019). Some of these species can potentially become invasive and through a series of mechanisms substantially change the community structure, cause the loss of native genotypes, modify habitats, affect food web properties and ecosystem processes, impede the provision of ecosystem services, impact human health, and cause substantial economic losses (Katsanevakis et al., 2014).

Here, we report the first confirmed record of the yellow-edged lyretail grouper, *Variola louti* (Forsskål, 1775) in the Mediterranean Sea and Cyprus. *Variola louti* is an Indo-Pacific reef-associated grouper of commercial importance in its native range, and a popular species in the aquarium trade. It was first reported in Cyprus approximately a year ago based on an underwater observation, and its introduction was attributed to an aquarium release (Kousteni et al., 2019). In this report, the species was identified using both morphological characteristics and genetic analysis. The phylogenetic relationship with available sequences from specimens analysed in other regions was further examined and discussed.

26 Materials and Methods

On 29 August 2019, a recreational spearfisher caught a *V. louti* individual at the north-western tip of Cyprus,
near Pomos village (32.55 E 35.17 N). The fish was caught around noon on a rocky bottom by 15 m depth,
approximately 90 km ocean distance from the area of its first Mediterranean sighting a year ago (Kousteni
et al., 2019) (Figure 1).

The specimen was morphologically examined for confirmation of its identity. DNA was extracted and the mitochondrial barcode gene COI (Cytochrome oxidase 1) was sequenced following published protocols

33 (Bariche et al., 2015). Briefly, the amplification of COI used fish specific primers VF2T1 and VR1dT1

34 (Ivanova et al., 2007). PCR amplified fragments were sequenced in both directions using the primers used 35 for the amplification, and then compared with available sequences in GenBank. Phylogenetic reconstructions were based on the Neighbour-Joining method generated in R (RCoreTeam, 2016) with the 36 use of the ape package (Paradis, Claude, & Strimmer, 2004). Genetic distances were based on the Kimura 37 2 parameter method. The maximum likelihood (ML) method was also used as a second phylogenetic 38 reconstruction approach, as implemented in GARLI (Zwickl, 2006). To estimate support for the nodes, 39 1000 bootstrap replicates were performed and we retained only the values supporting the nodes accounting 40 41 for more than 50% of the bootstrap replicates.

42 **Results**

43 Morphology

The study specimen was a sexually immature individual of 2145 g wet weight. It had an oblong body, with 44 the maximum body depth and head length 3 and 2.7 times in standard length respectively. The dorsal head 45 profile and the interorbital area were convex. Both jaws had a pair of large canines at the front, while the 46 47 lower jaw also had two large canines at the midsides. Palatines and vomer were also toothed. The caudal fin was lunate with the upper and lower lobes produced, about twice the length of middle rays, and the 48 pelvic fins extended past the anus. The body and median fins were brown to orange-red with numerous 49 small round or elongate spots of lavender and pink, while caudal, dorsal, anal, and pectoral fins had a broad 50 51 yellow rear margin. All morphological characteristics and morphometric and meristic measurements (Table 52 1) are in agreement with V. louti, as described in the literature (Heemstra, Randall, Carpenter, & Niem, 53 2001).

54 *Genetic analysis*

The PCR amplification and sequencing of the cytochrome oxidase 1 resulted in a 658 bp fragment 55 (GenBank accession number MN475883). A BLAST comparison of this sequence with available sequences 56 in GenBank placed it in a cluster of 23 sequences, all identified as V. louti. Seven of those sequences were 57 identical to the one obtained for our sample. These seven sequences belonged to samples collected in 58 59 Australia, Indonesia, Philippines, and China. Two V. louti sequences available in GenBank were obtained from samples collected in the Red Sea. One sequence from Egypt has a 99.69% similarity with our sequence 60 61 (MH707293, two mismatches), and one sequence from Eilat, Israel, has a 97.7% similarity with our 62 sequence (MF124078, 15 mismatches).

Phylogenetic analyses were performed by comparing our sequence to *V. louti* sequences extracted from
GenBank, using four *V. albimarginata* (the only other *Variola* species) sequences as outgroups. Maximum
likelihood and Neighbor-Joining methods resulted in identical tree topologies, therefore only the NJ tree is
shown here (Figure 2). As indicated above from the BLAST results, our sequence clustered with *Variola louti* samples, and was very well separated from the *V. marginata* sequences. When all available sequences

68 of V. marginata were used, and additional grouper species were used as outgroups, results remained 69 unchanged (not shown).

70 No samples from the Red Sea or the Indian Ocean (Mozambique, South Africa and India), clustered with 71 our samples, however, resolution at the level of this genetic marker is relatively low due to low mutation 72 rate in CO1, therefore such a geographic association is difficult to rule out.

Discussion 73

Morphology and genetic results with both BLAST and phylogenetic analyses unambiguously identify this 74 specimen as the yellow-edged lyretail grouper, Variola louti. While only four samples from GenBank are 75 76 from the Indian Ocean (two from the Red Sea and two from Mozambique), none of them cluster with our sample (a result that would be consistent with a Lessepsian immigrant). In contrast, results tentatively 77 indicate an aquarium release, since most aquarium trade fish are imported from other regions (e.g. Indonesia 78 and Philippines). The presence of a single large individual in Cyprus, which could in fact be the same 79 specimen previously observed in Cyprus (although that individual was recorded as being of a larger size 80 >70 cm TL) (Kousteni et al., 2019), is also consistent with this hypothesis. Yet, further work, sampling and 81 82 observations, as well as the engagement of citizen scientists, are necessary to conclusively settle this 83 auestion. Je.

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87 **Conflict of Interest**

88 None.

89 **Data Availability Statement**

90 The data that support the findings of this study are openly available in "GenBank" at https://www.ncbi.nlm.nih.gov/genbank/, accession number MN475883. 91

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Table 1. Morphometric and meristic measurements of the examined Variola louti specimen.

- Figure 1. (A) The location of the capture, indicated with a red circle, and of the previous Mediterranean
 record indicated with a black circle. (B) Specimen analysed in this study.
- **Figure 2.** Phylogenetic reconstruction of *Variola* groupers based on the cytochrome oxidase marker. Tree
- topology is based on the Neighbour-Joining, NJ, method (identical to Maximum Likelihood, ML,
- topology), numbers on nodes are bootstrap values derived from 1000 replicates (only numbers above 50%
- are shown). Firs number is for NJ, second number for ML. Mediterranean sample is from Cyprus and is
- 160 in red. All other sequences are from GenBank and are in black. Their sample origin is indicated after their
- 161 accession number.

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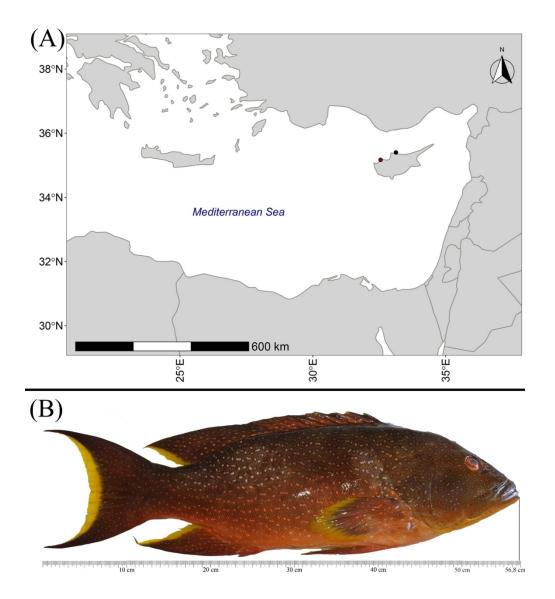


Figure 1. (A) The location of the capture, indicated with a red circle, and of the previous Mediterranean record indicated with a black circle. (B) Specimen analysed in this study.

169x197mm (220 x 220 DPI)

