

Resolution of the *Portunus gladiator* species complex: taxonomic status and identity of *Monomia gladiator* (Fabricius, 1798) and *Monomia haanii* (Stimpson, 1858) (Brachyura, Decapoda, Portunidae)

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Abstract

The United States Food and Drug Administration (FDA) has recently adopted DNA barcoding for the purpose of determining the species identity of commercial seafood products. This effort has revealed instances of incongruence between current scientifically accepted taxon names and those utilized by the seafood industry in product labelling. One such case is that of “*Portunus haanii*”, a name utilized by the seafood industry to label commercial products under the market name “red swimming crab.” However, carcinologists currently regard *P. haanii* as synonym of *Portunus gladiator* Fabricius, 1798, which itself is the subject of debate over whether it is a secondary homonym of *Cancer gladiator* Fabricius, 1793. Further complicating matters, DNA barcode sequences from commercial products match GenBank sequences identified as *Portunus pseudoargentatus* Stephenson, 1961. Here the complicated taxonomic history of the *Portunus gladiator* complex is reviewed and a resolution proposed based on combined morphological descriptions and molecular phylogenetic analyses. It is demonstrated that, given the provisions of the International Code of Zoological Nomenclature and the current elevation of *Monomia* Gistel, 1848, to full genus rank, its type species, *Portunus gladiator* Fabricius, 1798, should be treated as a valid and available taxon name. It is also shown, upon examination and comparison of types and topotypic material that *Monomia haanii* (Stimpson, 1858) is a distinct taxon from *M. gladiator*, and *Portunus pseudoargentatus*

Stephenson, 1961, is a junior subjective synonym of *M. haanii* (Stimpson, 1858). Furthermore, it is shown that crab meat sold in the US currently labeled as “*Portunus haanii*” and/or “red swimming crab” is in fact *M. haanii* using comparative analysis of DNA barcode sequences between museum-vouchered reference specimens, whole crabs provided directly by a seafood importer, and processed commercial products purchased at retail.

Keywords

Commercial species, DNA barcoding, molecular phylogenetics, morphology, seafood, swimming crab, taxonomy

Introduction

The United States Food and Drug Administration (FDA) has adopted DNA barcoding for the purpose of species identification to assure the accurate labelling of seafood products as well as to address issues with species substitution and fraud (Handy et al. 2011; Eischeid et al. 2016). This effort has identified instances of incongruence between currently accepted taxon names and the names utilized by industry in product labelling. At the same time that FDA was beginning to build its reference library of decapod crustacean DNA standards (Food and Drug Administration 2017), species substitution of brachyuran crabs was highlighted by a survey of crab cakes from restaurants in the Maryland / Washington, DC metropolitan area (Warner et al. 2015). DNA testing from that survey of products advertised by restaurants to contain crabmeat only from “local” *Callinectes sapidus* Rathbun, 1896, revealed at least one of six species of substituted portunoid crab in 38% of the crab cakes tested (Warner et al. 2015). According to the report, the most commonly detected substitute species was *Portunus pelagicus* (Linnaeus, 1758) followed by *P. pseudoargentatus* Stephenson, 1961, and *P. sanguinolentus* (Herbst, 1783). The reported DNA match to *P. pseudoargentatus* was of particular interest to Warner et al. (2015) who noted that this species was, at the time, not included in the FDA Guide to Acceptable Market Names for Seafood Sold in Interstate Commerce (The Seafood List) (Food and Drug Administration 2012) nor in the FAO fishery species list (Food and Agriculture Organization of the United Nations 2010–2018), raising concern that this species was unknown to regulators. As a result, in addition to the existing entry for *P. haanii*, *P. pseudoargentatus* was added to The Seafood List in 2015 (Food and Drug Administration 2015). Concerns over an unknown species that was apparently common in the US food supply led us to further investigate the identity of *P. pseudoargentatus* and its close relatives within the *Portunus gladiator* species complex to determine their relationship to what was currently being harvested and marketed as “*P. haanii*” and/or “red swimming crab.”

At the time of the study by Warner et al. (2015), only three sequences from three loci belonging to one specimen identified as “*Portunus pseudoargentatus* (ZMMU Ma 3368),” which was collected from Nhatrang Bay, Vietnam, had been deposited

in GenBank (JX398121, JX398079, JX398094) (Spiridonov et al. 2014). Further complicating matters, there is a set of DNA barcode sequences (BOLD: AAO6694) identified as “*Portunus haanii*” from the Coral Sea in the Barcode of Life Database (BOLD), which led to more uncertainty about the true identity of crabs in the US seafood supply. The specimens in question, which are deposited in the Muséum national d’Histoire naturelle (MNHN) in Paris under the revised name of *P. gladiator* (MNHN-IU-2008-12570-77), have since been examined by one of the present authors (AMW) and confirmed to be neither *P. gladiator* nor *P. pseudoargentatus*, but morphologically and genetically closer to *Monomia lucida* Koch and Āuriš, 2017. *Portunus haanii* was included on the FDA Seafood List prior to this study without knowledge that *P. haanii* is currently an unaccepted species name, having been synonymized under *P. gladiator* Fabricius, 1798 (Stephenson and Cook 1973, Ng et al. 2008).

Within the seafood industry, *Portunus haanii* and/or “red swimming crab” is the species/market name used for crabs harvested extensively from China and Vietnam (Monterey Bay Aquarium Seafood Watch 2013; Fishsource 2016). Photographs identified as *P. haanii* on websites advertising “*Portunus haanii*” (e.g., Fishsource 2016; Alibaba.com 2018) bear a striking resemblance to the color photograph of “*Portunus pseudoargentatus* (ZMMU Ma 3368)” (Chertoprud et al. 2012: pl. 51 fig. H) and Stephenson’s (1961) original description of *P. pseudoargentatus*. The names *Portunus gladiator* and *Portunus pseudoargentatus* were not found associated with any specific crabmeat products in neither our on-line searches nor our discussions with industry representatives.

Taxonomic history

The taxonomy of the *Portunus gladiator* complex is so convoluted that it makes a chronologically arranged taxonomic history difficult to compile. Here, we present the significant taxonomic actions for *P. gladiator*, *P. haanii*, and *P. pseudoargentatus*.

Fabricius (1798) described a swimming crab collected from “Oceano Asiatico Dom. Daldorff” (probably Tranquebar, India; see Ng et al. 2008), which he named *Portunus gladiator*, based on an unspecified quantity of specimens. Fabricius (1798: 368) gave a brief description in Latin, stating: “P. thorace tomentoso utrinque novemdentato: dente postico maiore, minibus sanguineo maculatis. Habitat in Oceano Asiatico Dom. Daldorff. Praecedentibus affinis at minor. Thorax holosericeus, parum inaequalis, hine inde scaber. Chelae sanguineo maculatae digitis apice dentibusque albis.”

Previously, however, Fabricius (1793) had given the same specific epithet to a different species of swimming crab (from “Nova Hollandia” = Australia), *Cancer gladiator*. Fabricius (1793: 449) provided this description: “C. thorace laevi: lateribus octodentatis, postico maximo, minibus angulatis. Cancer hastatus. Mant. Ins. r. 319. 34. Habitat

in Nova Hollandia Mus. Dom. Banks. Minutus pullo modo Cancer hastatus Linnaei. Palmae anticae bidentatae, chelae angulatae. Palmae posticae angulatae.”

Latreille (1825) in his treatment of *Portunus*, considered *Cancer gladiator* Fabricius, 1793, a junior synonym of *Portunus sanguinolentus* (Herbst, 1783), and clearly also considered *Portunus gladiator* Fabricius, 1798, to be a distinct species. From this point on, it appears that the name “*Portunus gladiator*” has been almost exclusively associated with the species described in 1798. Stephenson and Cook (1973) would later accept Latreille’s (1825) synonymization and selected a male specimen of *P. sanguinolentus* from Queensland, Australia (QM W3683) to be the neotype of *C. gladiator* Fabricius, 1793, in order to stabilize its taxonomy.

De Haan (1833) then established the subgenus *Portunus* (*Amphitrite*), to which Miers (1886) subsequently designated *Neptunus* (*Amphitrite*) *gladiator* (Fabricius, 1798), as its type species. Prior to Miers’ action, however, Gistel (1848) had proposed a replacement name, *Portunus* (*Monomia*), for this subgenus as the name *Amphitrite* had been previously used for a polychaete annelid genus, viz. *Amphitrite* Müller, 1771. By virtue of these, *Portunus gladiator* Fabricius, 1798, is the type species of *Portunus* (*Monomia*) Gistel, 1848.

Stimpson (1858) gave the name *Amphitrite haanii* to a species from Tanegashima and Kagoshima, Japan and the ‘China Seas, above 23°N latitude,’ previously identified by De Haan (1833, 1835) as *Portunus* (*Amphitrite*) *gladiator* (Fabricius, 1798). Stimpson (1858: 38) wrote: “Amphitrite Haanii. *A. gladiator*, De Haan; loc. cit. p. 29, pl. i. f. 5. (vix *L. gladiator*, M. Edwards.) In mari Sinensi, lat. bor. 23°; ad insulam ‘Tanegasima’; et in sinu ‘Kagosima’; in fundis arenosis prof. 12–20 org.”

As Stimpson (1858) had material from Japan and also referred to De Haan’s (1833) citations of the species as *P. (A.) gladiator*, all of Stimpson’s and De Haan’s specimens are effectively syntypes. Later, Stimpson (1907: 79) provided a slightly more detailed explanation, clearly opining that De Haan’s (1833, 1835) Japanese *P. gladiator* was a distinct species from H. Milne Edwards’ (1834) *Lupea gladiator* (= *P. gladiator* Fabricius, 1798) from the Indian Ocean. There are 53 extant syntypes of *Amphitrite haanii* in the collection of the Naturalis Biodiversity Center in Leiden as recorded by Yamaguchi and Baba (1993), and among these they selected a male specimen (RMNH 379, CW = 42 mm, CL = 24 mm, C. Fransen pers. comm.) to be the lectotype.

Stephenson (1961) described *Portunus pseudoargentatus* based on one male specimen from the Abrolhos Islands, off the western coast of Australia. He cited differences in the morphology of the male 6th pleomere and male gonopod 1 (G1) between *P. gladiator* and the new species. Crosnier (1962) in his treatment of the swimming crabs of Madagascar commented on the confusion in the identities of specimens from various localities labelled as “*P. gladiator*.” He highlighted the differences in the morphology of the male 6th pleomere and G1, wherein *P. pseudoargentatus* tends to have the male 6th pleomere with less sinuous lateral borders, and the G1 more greatly bent in the middle (midlength) compared to *P. gladiator*. He also referred to Japanese material that he examined to *P. pseudoargentatus*, apparently not considering Stimpson’s (1858, 1907) earlier reports on *P. haanii*.

Stephenson and Cook (1973) did an extensive study on the *Portunus gladiator* complex, and they argued that, with both being then classified in the genus *Portunus*, *P. gladiator* Fabricius, 1798, effectively became a secondary homonym of *P. gladiator* (Fabricius, 1793) (= *P. sanguinolentus*), and a replacement name was needed. They went on to suggest *Amphitrite haanii* Stimpson, 1858, as the earliest available replacement name, eliminating an earlier name, *Cancer menestho* Herbst, 1803, on the basis of a difference in colouration and in the armature of the cheliped merus in the illustration of the latter (viz. Herbst, 1803:pl. 55 fig. 3). They also considered *Portunus pseudoargentatus* Stephenson, 1961, as a junior synonym of *Portunus haanii* (Stimpson, 1858). Additionally, Stephenson and Cook (1973) described aberrant “forms” within *P. haanii*, on the basis of the G1 morphology (e.g., “normal” vs. “unusual” specimens; viz. Stephenson and Cook 1973: figs 6, 7). These “unusual” specimens were two crabs from the Bay of Jeddo, Japan, and the holotype of *P. pseudoargentatus*. They did not, however, take steps to formally distinguish these forms as separate species, subscribing instead to the concept of “*Portunus haanii*” as a morphologically variable species.

Ng et al. (2008: 156, 157) provided a detailed account on the nomenclature of Fabricius’ two species with the epithet *gladiator*, essentially stating that, contrary to Stephenson and Cook (1973), there is no secondary homonymy unless *Cancer gladiator* is considered a distinct species from *Portunus sanguinolentus*: “... as *Cancer gladiator* Fabricius, 1793, is regarded as a junior synonym of *Portunus sanguinolentus* (Herbst, 1783), the name ‘*Portunus gladiator* (Fabricius, 1793)’ has not been recognized or used anywhere. This being the case, there is no homonymy with *Portunus gladiator* Fabricius, 1798, and this name should remain available for use under the Code. The issue of secondary homonymy will only arise if *Cancer gladiator* Fabricius, 1793, is regarded as a valid species of *Portunus* distinct from *Portunus sanguinolentus* (Herbst, 1783). If this were to happen (for example, if the widespread *P. sanguinolentus* was to prove to be a complex of several cryptic species), then the name *Portunus gladiator* Fabricius, 1798, would have to be replaced by the next available name...”. Ng et al. (2008), however, maintained the synonymy of *Portunus gladiator* and *Amphitrite haanii*, while still considering *Portunus pseudoargentatus* a valid species.

Chertoprud et al. (2012) in their report on the commercially valuable brachyuran species of Vietnam, commented that Stephenson and Cook’s (1973) mention of “*Portunus gladiator* (Fabricius, 1793)” was sufficient to activate Article 59 of the Code (ICZN 1999), which therefore necessitated the use of *Portunus haanii* (Stimpson, 1858) as a replacement name for *Portunus gladiator* Fabricius, 1798.

As things stand, the issue on the validity of the names “*Portunus gladiator*” and “*Portunus haanii*” has not been satisfactorily settled. Recent publications on the systematics of Portunidae have bolstered the concept of *Monomia* Gistel, 1848, as a valid genus-level taxon distinct from *Portunus*, but these have also shown that the problem with the taxonomy of the type species, *M. gladiator* (Fabricius, 1798), and its closely related congeners, *M. haanii* (Stimpson, 1859) and *M. pseudoargentata* (Stephenson, 1961), remains unresolved (Chertoprud et al. 2012; Spiridonov et al. 2014; Koch et al. 2017; Koch and Āuriš 2018). This problem, unless addressed,

is expected to have wide-ranging consequences on the taxonomy of *Monomia*. As such, the identities of *M. gladiator*, *M. haanii*, and *M. pseudoargentata* need to be firmly established.

An integrative approach with morphological and molecular phylogenetic analyses was undertaken to resolve and stabilise the taxonomy of the *Portunus gladiator* (= *Monomia gladiator*) complex. The molecular results of the morphologically verified and vouchered reference specimens, which included whole specimens from Asian fish ports and a seafood importer, were then used as standards to identify the contents of cans of pasteurized lump crabmeat labeled as “*Portunus haanii*” and/or “red swimming crab” through comparative analysis of DNA barcode sequences.

Materials and methods

Taxonomic methods

Materials examined are deposited at the US National Museum of Natural History, Smithsonian Institution (USNM); Florida Museum of Natural History, University of Florida (UF); Western Australian Museum (WAM); and Lee Kong Chian Natural History Museum, National University of Singapore (ZRC). These included the holotype of *Portunus pseudoargentatus* (WAM-C7506), as well as crabs purchased at fish ports in India, Thailand, and Taiwan, and whole crabs identified by a US seafood importer as “*Portunus haanii*.” Photographs of the type specimens of *Portunus gladiator* and *Amphitrite haanii*, housed at Zoological Museum at the University of Copenhagen (ZMUC) and Naturalis Biodiversity Center, Leiden (RMNH), respectively, were also examined. Details on all specimens utilized in morphological examinations are provided in the material examined subsection of the taxonomic account below. The morphological terminology largely follows Wee and Ng (1995) and Apel and Spiridonov (1998). The following abbreviations are used:

- CL** carapace length, taken along the dorsal midline from the tips of the frontal teeth to the posterior margin of the carapace;
- CW** carapace maximum width, taken at the level of its widest point;
- P1–P5** first to fifth pereopods, respectively (P1, chelipeds; P2–P5, first to fourth ambulatory legs);
- G1, G2** first and second male pleopods, respectively.

The term, pleomere (first to sixth), here refers to the six somites of the pleon. When possible, DNA was extracted from the specimens utilized for the morphological studies. Details on all specimens utilized in the molecular phylogenetic component of this study are given in Table 1, Nomenclatural decisions are based on the provisions of the International Code of Zoological Nomenclature, here referred to as “the Code” (ICZN 1999).

Table 1. Material examined in molecular analyses with details on voucher identification numbers, sex, country in which the specimen was collected, the fish port or body of water, and pertinent GenBank Accession Numbers. Voucher ID abbreviations: IOM= Institute of Oceanology and Museum, Nha Trang; MNHN= Muséum National d'Histoire Naturelle, Paris; NHMUK= The Natural History Museum, London; UF= University of Florida Natural History Museum, Gainesville; UO= University of Ostrava, Ostrava; USNM= United States National Museum, National Museum of Natural History, Washington, D.C.; WAM= Western Australian Museum, Perth; ZMMU= Zoological Museum of the Moscow University, Moscow; ZRC= Zoological Reference Collection, Lee Kong Chian Natural History Museum, Singapore.

Taxon name	Voucher ID	Sex	Country	Port/Body of Water	12S	16S	COI
<i>Monomia gladiator</i>	ZRC 2016.0145	M	India	Pazhayar Fish Landing, Bay of Bengal	MK270964	—	MK281257
<i>Monomia gladiator</i>	ZRC 2016.0149	F	India	Pondicherry, Bay of Bengal	MK270959	MK271060	MK281259
<i>Monomia gladiator</i>	WAM C61156	F	Australia	Pilbara Shelf, Indian Ocean	MK270957	MK271053	MK281253
<i>Monomia gladiator</i>	WAM C26459	M	Australia	Dampier Archipelago, Cape Brugières	MK270956	MK271047	MK281247
<i>Monomia gladiator</i>	UF 36251	F	Singapore	Singapore Strait	MK270963	MK271029	MK281229
<i>Monomia gladiator</i>	USNM 127068	F	Thailand	Andaman Sea	MK270962	MK271030	MK281230
<i>Monomia gladiator</i>	ZRC 2000.0842	M	Thailand	Pichai Fish Port, Phuket, Andaman Sea	MK270958	—	MK281230
<i>Monomia gladiator</i>	ZRC 2003.0114	M	Thailand	Pattani Fish Port, Gulf of Thailand	MK270960	MK271055	—
<i>Monomia gladiator</i>	ZRC 2003.0197	M	Thailand	Saiburi Crab Landing, Gulf of Thailand	MK270961	MK271056	—
<i>Monomia gladiator</i>	ZRC 2002.0297	F	Thailand	Pichai Fish Port, Phuket, Andaman Sea	—	—	—
<i>Monomia gladiator</i>	MNHN-IU-2014-10087	—	Vietnam	Cù Bè Fishing Port	—	KY524466 ¹	—
<i>Monomia gladiator</i>	ZMMU Ma 3366	—	Vietnam	Cù Bè Fishing Port	—	—	JX398095 ²
<i>Monomia haanii</i>	ZRC 1999.0084	M	Japan	Pacific Ocean	MK270948	MK271054	MK281255
<i>Monomia haanii</i>	WAM C61155	F	Australia	Pilbara Shelf	MK270946	MK271052	MK281252
<i>Monomia haanii</i>	WAM C34767	F	Australia	Exmouth Gulf	MK270944	MK271048	MK281248
<i>Monomia haanii</i>	WAM C34900	M	Australia	Exmouth Gulf	MK271049	MK271049	MK281249
<i>Monomia haanii</i>	WAM C44737	F	Australia	Ningaloo Marine Park	MK271051	MK271051	MK281251
<i>Monomia haanii</i>	WAM C34938	F	Australia	Shark Bay	MK270949	MK271050	MK281250
<i>Monomia haanii</i>	WAM C55510	M	Australia	Shark Bay	MK270943	MK271045	MK281245
<i>Monomia haanii</i>	WAM C7506*	M	Australia	Abrolhos Islands	MK270953	MK271046	MK281246
<i>Monomia haanii</i>	USNM 1421161	M	China	South China Sea, FAO Fishing Area 61	MK270934	MK271033	MK281233
<i>Monomia haanii</i>	USNM 1421181	M	China	South China Sea, FAO Fishing Area 61	MK270935	MK271034	MK281234
<i>Monomia haanii</i>	USNM 1421182	M	China	South China Sea, FAO Fishing Area 61	MK270936	MK271035	MK281235
<i>Monomia haanii</i>	USNM 1421185	M	China	South China Sea, FAO Fishing Area 61	MK270937	MK271036	MK281236
<i>Monomia haanii</i>	USNM 1421187	M	China	South China Sea, FAO Fishing Area 61	MK270950	MK271037	MK281237

Taxon name	Voucher ID	Sex	Country	Port/Body of Water	12S	16S	COI
<i>Monomia haanii</i>	USNM 1421191	M	China	South China Sea, FAO Fishing Area 61	MK270938	MK271038	MK281238
<i>Monomia haanii</i>	USNM 1421194	M	China	South China Sea, FAO Fishing Area 61	MK270951	MK271039	MK281239
<i>Monomia haanii</i>	USNM 1421195	M	China	South China Sea, FAO Fishing Area 61	MK270939	MK271040	MK281240
<i>Monomia haanii</i>	USNM 1421196	M	China	South China Sea, FAO Fishing Area 61	MK270940	MK271041	MK281241
<i>Monomia haanii</i>	USNM 1421202	M	China	South China Sea, FAO Fishing Area 61	MK270941	MK271042	MK281242
<i>Monomia haanii</i>	USNM 1421204	M	China	South China Sea, FAO Fishing Area 61	MK270952	MK271043	MK281243
<i>Monomia haanii</i>	USNM 1421206	M	China	South China Sea, FAO Fishing Area 61	MK270942	MK271044	MK281244
<i>Monomia haanii</i>	USNM 1420827	F	Taiwan	Daxi Fishery Port	MK270933	MK271031	MK281231
<i>Monomia haanii</i>	USNM 1420828	M	Taiwan	Daxi Fishery Port	MK270955	MK271032	MK281232
<i>Monomia haanii</i>	UF 29509	F	Taiwan	Daxi Fishery Port	MK270930	MK271026	MK281227
<i>Monomia haanii</i>	UF 29511	F	Taiwan	Daxi Fishery Port	MK270931	MK271027	MK281228
<i>Monomia haanii</i>	UF 29512	F	Taiwan	Daxi Fishery Port	MK270932	MK271028	—
<i>Monomia haanii</i>	ZRC 1998.0186	M	Taiwan	Daxi Fishery Port	MK270947	MK271059	MK281254
<i>Monomia haanii</i>	MNHN-IU-2014-10086		Vietnam	Cù Bè Fishing Port	—	KY524463 ¹	—
<i>Monomia haanii</i>	UO 12J-Vn12		Vietnam	Cù Bè Fishing Port	—	KY524464 ¹	—
<i>Monomia haanii</i>	ZMMU Ma 3368		Vietnam	Cù Bè Fishing Port	—	KY524480 ¹	JX398094 ²
<i>Monomia argenteata</i>	MNHN-IU-2014-10076		Vietnam	Cù Bè Fishing Port	—	KY524479 ¹	—
<i>Monomia argenteata</i>	MNHN-IU-2014-10075		Vietnam	Cù Bè Fishing Port	—	KY524478 ¹	—
<i>Monomia argenteata</i>	IOM		Vietnam	Cù Bè Fishing Port	—	MK271061	—
<i>Monomia lucida</i>	ZRC 2016.0150	M	Vanuatu	South Pacific Ocean	MK270965	—	JX398096 ²
<i>Monomia lucida</i>	ZMMU Ma 3365		Vietnam	Cù Bè Fishing Port	—	MG563792 ³	—
<i>Monomia lucida</i>	NHMUK 2017.402		Vietnam	Cù Bè Fishing Port	—	MG563793 ³	—
<i>Monomia lucida</i>	MNHN-IU-2014-10083		Vietnam	Cù Bè Fishing Port	—	MG563794 ³	—
<i>Monomia lucida</i>	MNHN-IU-2014-10085		Vietnam	Cù Bè Fishing Port	—	KT365606 ⁴	—
<i>Monomia petrea</i>	UF 188		Guam	Tepungan Channel	—	—	KT365743 ⁴
Outgroup Taxa							
<i>Portunus sanguinolentus</i>	ZRC 2016.0146	M	India	Pazhayar Fish Landing	MK270966	MK271057	—
<i>Portunus pelagicus</i>	ZRC 2016.0147	M	India	Porto Novo	MK270967	MK271058	MK281258

Sequences mined from GenBank are attributed to ¹Koch and ²Duris (2018), ³Spiridonov et al. (2014), ³Koch et al. (2017), and ⁴Evans (2018). *Holotype of *Portunus pseudoargenteatus* Stephenson, 1961

Commercial products

Four cans (454 g each) of pasteurized lump crabmeat labeled as “*Portunus haanii*” were purchased from grocery stores in Maryland and Virginia in 2016 and 2017. Portions of 10 lumps (i.e., single piece of crabmeat reasonably expected to be from an individual crab), five from the top and five from the bottom, from each tub were sampled for DNA extraction (N=40). The DNA barcode region of the cytochrome oxidase subunit I (COI) was amplified and sequenced from samples following the methods described below.

Molecular methods

Genomic DNA was extracted from muscle tissue dissected from ethanol preserved or fresh specimens using the DNeasy Tissue Kit (Qiagen) according to the manufacturer’s animal tissue protocol. Portions of three mitochondrial genes were amplified: a 658 bp barcode region of the cytochrome c oxidase I gene using the primers JgLCO1490 and JgHCO2189 (Geller et al. 2013), a 531 bp region of the 16S ribosomal gene using the primers 16S-ar and 16S-br (Palumbi 1996), and a 375 bp region of the 12S ribosomal gene using the primers 12Sf (Mokady et al. 1994) and 12S1R (Shull et al. 2005). PCR was carried out for 35 cycles with an annealing temperature of 48 °C for COI and 52 °C for 12S and 16S using Promega GoTaq G2 hot start master mix (Promega M7432). PCR products were visualized by agarose gel electrophoresis (1.5% agarose) and purified with ExoSAP-IT (Affymetrix) prior to sequencing. Sequencing reactions were performed using 1 µL of purified PCR product in a 10 µL reaction containing 0.5 µL primer, 1.75 µL Big Dye buffer and 0.5 µL Big Dye (Life Technologies).

Geneious 9.1.7 (Biomatters) was used to visualize, trim, edit, and assemble contigs from forward and reverse sequences. All PCR, sequencing, and analytics were carried out at the Laboratories of Analytical Biology at USNM. Sequences have been deposited in GenBank (NCBI) with accession numbers listed in Table 1.

Partial sequences for each locus were also amplified from *Portunus pelagicus* (ZRC 2016.0147) and *P. sanguinolentus* (ZRC 2016.0146) to serve as outgroup taxa. Multiple sequence alignments were generated using the L-INS-i alignment strategy in MAFFT version 7 (Katoh and Standley 2013). The aligned sequences were then concatenated using Sequence Matrix (Vaidya et al. 2011). In the concatenated data set, positions 1–380 are 12S, 381–914 are 16S, and positions 915–1572 are COI.

A best-fit model of nucleotide sequence evolution compatible with MrBayes and partitioning arrangement for each locus was determined using Partition Finder 2 (Lanfear et al. 2016) with the greedy algorithm selected (Lanfear et al. 2012). The GTR+I+G model was chosen for all three loci. Phylogenetic analyses were performed on the concatenated dataset using maximum likelihood (ML) with RAxML (Stamatakis 2006) and Bayesian Inference (BI) performed with MrBayes 3.1.2 (Ronquist and Huelsenbeck 2003) on the FDA’s Raven2 high performance computing cluster. ML options for RAxML included the GTRCAT model of nucleotide evolution (-m), rapid bootstrap

analysis, and search for best-scoring ML tree (-f a), and 1000 bootstrap replicates. BI analysis was carried out for 10 million generations with two independent runs, each with four chains, and with trees sampled every 1000th generation. Model parameters (tratio, statefreq, shape, pinvar) were unlinked among partitions, and the rate prior (prset ratepr) was set to “variable.” To calculate posterior probabilities, a “burn-in” of 25% of the total trees sampled per run adequately removed trees prior to convergence.

In addition to the concatenated dataset, a COI-only dataset which incorporated sequences from GenBank and BOLD was analysed to identify the species of crab found in four cans of pasteurized lump crabmeat labelled as “*Portunus haanii*”. For visualization purposes, a neighbour joining tree of the 658bp alignment was built using the Jukes-Cantor model in the Geneious Tree Builder. Patristic and K2P distances were calculated for each alignment using MEGA7 (Kumar et al. 2016).

Results

Molecular phylogenetics

Molecular phylogenetic analyses of the concatenated dataset of three mitochondrial loci show that there is a well-supported (98/1) separation between *M. gladiator* and *M. haanii*. *Monomia petrea* (Alcock, 1899) (UF188; KT365743, KT365606) is a strongly supported sister to both (95/1) (Fig. 5). Analysis also confirms that voucher specimens purchased at the Daxi Fishery Port in northern Taiwan as well as those supplied by U.S. seafood importer, Newport International, both under the name “*P. haanii*,” are the same species as the specimen of “*M. pseudoargentata*” (ZMMU 3368; JX398094) from Spiridonov et al. (2014). Furthermore, our specimens with color patterns similar to “*M. haanii*” in Chertoprud et al. (2012) (= *M. gladiator* ZMMU 3366 in Spiridonov et al. 2014; JX398095) are the same species: *M. gladiator s. str.* (Fig. 5). Included in the clade of *M. haanii* is the holotype of *Portunus pseudoargentatus* (WAM-C7056). The topology of the ML phylogram is congruent with the morphological findings.

DNA barcode sequence analyses for species identification of products confirm that crabmeat sold as “*Portunus haanii*” is indeed what we have identified herein as *Monomia haanii s. str.* (Fig. 6). The mean K2P distance between reference *M. haanii* and the 40 product samples is 0.72% compared to the 7.85% between *M. gladiator* and product samples. Similarly, COI sequences from *M. haanii* and *M. gladiator* reference samples have a mean K2P distance of 7.74% (Table 2) and is consistent with congeneric divergences observed in other decapods (Costa et al. 2007).

Table 2. Mean K2P distances between specimens genetically identified as *M. gladiator*, *M. haanii*, and commercial products calculated from a neighbor-joining distance tree built in Geneious.

	<i>Monomia gladiator</i>	<i>Monomia haanii</i>
<i>Monomia haanii</i>	7.74%	
Commercial Products	7.85%	0.72%

Taxonomic accounts

Portunoidea Rafinesque, 1815

Portunidae Rafinesque, 1815

Portuninae Rafinesque, 1815

Monomia Gistel, 1848

Type species. *Portunus gladiator* Fabricius, 1798, type species of *Amphitrite* De Haan, 1833, by subsequent designation (Miers, 1886); pre-occupied by *Amphitrite* Müller, 1771 [Polychaeta]; *Monomia* Gistel, 1848, replacement name for *Amphitrite* De Haan, 1833.

Monomia gladiator s. str. (Fabricius, 1798)

Figs 1A–D, 3A–C, 4A–D

Portunus gladiator Fabricius, 1798: 368; Latreille 1825: 189; Crosnier 1962 (in part): 51, figs 72, 76, 78, 82, 83, pl. 3 fig. 2; Stephenson and Rees 1967a: 14; 1967b: 25; 1968: 293 (in part); Stephenson 1972a: 16, 39 (in part); 1972b: 135 (in part); Bhadra 1998: 410; Dev Roy and Bhadra 2005: 425; 2011: 147.

Cancer menestho Herbst, 1803: 34, pl. 55 fig. 3.

Lupea gladiator, H. Milne Edwards 1834: 456.

Neptunus gladiator, A. Milne-Edwards 1861: 330; Richters 1880: 152; Müller 1887: 475; De Man 1888: 69; Henderson 1893: 367.

Neptunus (Amphitrite) gladiator, Miers, 1886: 177; Alcock 1899: 35, 36; Laurie 1906: 412.

Callinectes gladiator, Stebbing 1915: 58. Non *Callinectes gladiator* Benedict, 1893 (fide Stephenson and Cook 1973).

Monomia gladiator, Barnard 1950: 156; Fourmanoir 1954: 9; Spiridonov et al. 2014: table 1; Trivedi et al. 2018: 66, table 1.

Portunus (Monomia) gladiator, Jeyabaskaran et al. 2000: 51, pl. 36c; Biju Kumar et al. 2007: 286; Ng et al. 2008: 151, 156, 157 (list and discussion).

Portunus (Monomia) gladiator [sic], Krishnamoorthy 2009: 6 (list).

Portunus haanii, Stephenson and Cook 1973: 429 (in part), figs 6A–E, 7A–E, 8A–E, 9A, 10A, C, D, G; Stephenson 1975: 178. Non *Amphitrite haanii* Stimpson, 1858.

Monomia haanii, Chertoprud et al. 2012: 314, pl. 51 fig. G. Non *Amphitrite haanii* Stimpson, 1859.

Non *Cancer gladiator* Fabricius, 1793: 449 (= *Portunus sanguinolentus* (Herbst, 1783), fide Latreille 1825).

Non *Portunus (Amphitrite) gladiator*, De Haan 1833: 65; 1835: pl. 18 fig. 1 (= *Portunus orbitosinus* Rathbun, 1911).

Non *Portunus gladiator*, Stephenson and Campbell 1959: 110, Figs 2J, 3J, pl. 3 fig. 2, pl. 4 fig. J, pl. 5 fig. J (= *Portunus australiensis* Stephenson and Cook, 1973).

Material examined. INDIA: ZRC.2016.0145, 2 males, 1 female, Pazhayar Fish Landing, Nagapattinam District, Tamil Nadu, coll. NK Ng et al., 17 Sep. 2011; ZRC.2016.0149, 1 female, sandy beach, Pondicherry, Union Territory of Puducherry; ZRC.2018.1189, 5 males, Jeppiar Fishing Port, Muttom, Tamil Nadu, coll. PKL Ng et al., 19 Sep. 2016; USNM127069, 1 male, SW of Mumbai, IIOE Anton Bruun, 14 Nov. 1963.

AUSTRALIA: WAM-C26459, 2 males, Dampier Archipelago, Cape Brugières, Western Australia, coll. Slack-Smith and Hewitt, 17 Jul. 1999; WAM-C61155, 2 females, WAM-C61156, 1 female, Pilbara Shelf, Western Australia, coll. E Morello et al. (CSIRO Pilbara Survey), 13 Jun. 2013.

PENINSULAR MALAYSIA: ZRC.2000.1308, 4 males, Perhentian, coll. anon., 16 May 1976.

MYANMAR: ZRC.2016.0030, 1 female, Ayeyarwady Delta, coll. EAF-Nansen Project (Myanmar cruise), 19 May 2015; ZRC.2016.0034, 1 male, Tanintharyi Coast, coll. EAF-Nansen Project (Myanmar cruise), 26 May 2015.

SINGAPORE: ZRC.1965.10.22.1-2, 1 male, 1 female, Siglap, coll. M.W.F. Tweedie, Jul. 1933; ZRC.1984.338-348, 4 males, 7 females, Horsburgh Lighthouse, South China Sea near Singapore, coll. H Huat, 15 Dec. 1982; ZRC.1984.5451-5453, 3 males, Tuas fishery port, coll. WM Lee, 25 Sep. 1982.

THAILAND: USNM127068, 2 females, Andaman Sea, north of Phuket, IIOE Anton Bruun R/V, 31 July. 1963; ZRC.2000.0779, 3 males, 13 females, Phuket, Pichai Fish Port, coll. NK Ng et al., 17–20 Jan. 2000; ZRC.2000.0842, 2 males, Phuket, Pichai Fish Port (Andaman Sea), coll. PKL Ng et al., 3–6 May 2000; ZRC.2002.0297, 4 males, 1 female, Phuket, Pichai Fish Port, coll. JCY. Lai, 2–3 Sep. 2001; ZRC.2002.0298, 3 males, 1 female, Phuket, Pichai Fish Port, coll. JCY Lai, 2–3 Sep. 2001; ZRC.2003.0114, male, Pattani Fishing Port, Pattani Province; ZRC.2003.0197, 1 male, Saiburi Crab Landing, Pattani Province, coll. Z Jaafar et al., 8 Jun. 2003.

Diagnosis. Carapace (Fig. 1A–D) transversally hexagonal, CW/CL ratio 1.79–1.83, with dorsal surface, except patches of granules, densely covered by short tomentum. Regions moderately defined; with discrete patches of granules on gastric, branchial, cardiac and intestinal regions. Front subdivided into four teeth with rounded apices, median pair distinctly smaller than lateral; median sulcus between teeth continuing ventrally to triangular projection appressed to median epistomial tooth. Epistome well defined, median tooth projecting beyond front. Supraorbital margin finely granulate, with two distinct notches; inner orbital angle tooth-like, with glabrous ventromesial ledge. Infraorbital margin with deep, V-shaped notch laterally; in antero-ventral view, mesial part of infraorbital margin granulate, ventrally a large triangular tooth projecting anteriorly, visible from dorsal view. Anterolateral margin armed with nine teeth, including external orbital angle, with granulate margins; first tooth larger than following teeth except 9th, with straight outer margin; 2nd–8th subequal in size, sharp, projecting outward, slightly curved anteriorly; 9th tooth largest, projecting laterally; just underneath anterolateral margin is thick coat of long soft setae which sometimes obscures

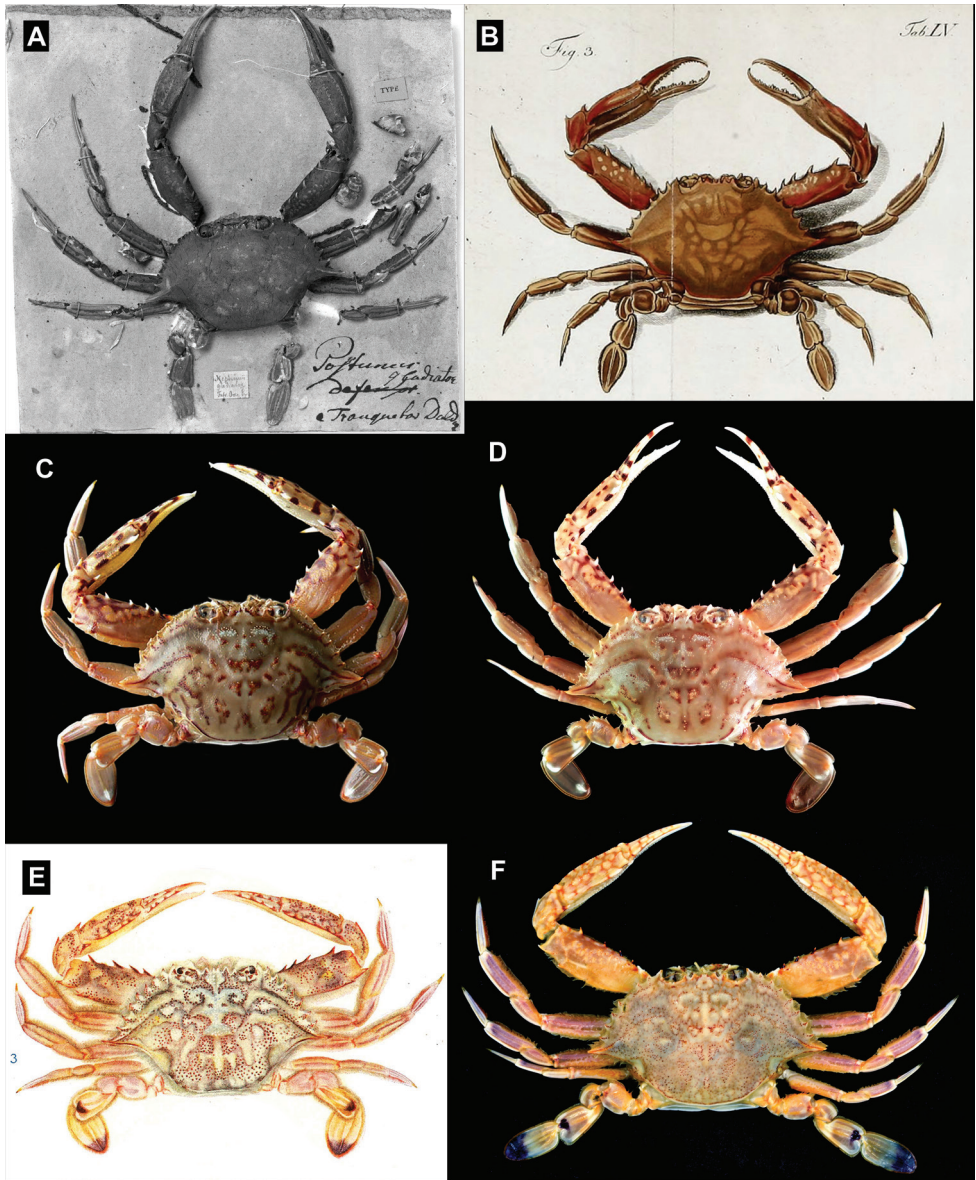


Figure 1. Dorsal habitus of **A** lectotype of *Portunus gladiator* Fabricius, 1798, deposited in Copenhagen Museum (ZMUC-Cru 4705) **B** *Cancer menestho* Herbst, 1803 (= *Monomia gladiator*), probably from Indian Ocean (color print from Herbst, 1803: pl. 55 fig. 3 **C** *Monomia gladiator* (Fabricius, 1798), fresh colouration, Phuket, Thailand (not collected), photo by Rueangrit Promdam **D** *Monomia gladiator* (Fabricius, 1798), fresh coloration, Jeppiar, Tamil Nadu, India (ZRC), photograph by PKL Ng; **E** “*Neptunus (Amphitrite) gladiator*” [sic] (= *Monomia haanii*) from Sagami Bay, Japan (color print from Sakai, 1939: pl. 47 fig. 3) **F** *Monomia haanii* (Stimpson, 1858), fresh colouration, South China Sea (USNM 1421161) shipped frozen by US seafood importer.

teeth. Posterolateral margin concave, posterolateral angle rounded; posterior margin lined with small granules, straight to slightly convex; posterior margin with ventrally directed, smooth flange, lateral extremities of which coincide with posterolateral angle of carapace, each armed with small, lamelliform tooth.

Basal article of antennule completely filling antennular fossa, subsequent two articles slender. Basal article of antenna short, with broad, lateral projection entering but not obstructing orbital hiatus; flagellum long, exceeding well beyond orbit. Eyes with well-developed corneas, short, thick peduncles. Proepistome well developed, anterior tip with projecting conical tooth; epistome not extensively projecting posteriorly. Endostome with well-developed lateral ridges.

Third maxillipeds finely granulate on ischium, merus and exopod, setose on external surfaces, extensively pilose on mesial margins; ischium longer than wide, subrectangular, with deep, submesial sulcus; merus longer than wide, rhomboidal, anterolateral angle strongly projecting laterally; palp articles subcylindrical. Exopod stout, with subdistal triangular projection on inner medial border; flagellum well developed.

Male thoracic sternites covered with thick tomentum, thickest on exposed surfaces of sternites 5–8 (Fig. 3A–C); sternites 1–4 distinctly granulate, with granules becoming larger, coarser in large specimens; sternites 1, 2 fused, separated from sternite 3 by straight suture; sternites 3, 4 almost completely fused except for remnants of suture at lateral extremities, replaced mesially by smooth, setose groove; sternite 4 with narrow median groove on exposed surface, continuing posteriorly into sterno-pleonal cavity. Sutures 4/5, 5/6, and 6/7 present on exposed surface of thoracic sternum but interrupted within sterno-pleonal cavity; suture 7/8 present on most of exposed thoracic sternal surface, but disappearing just before sternopleonal cavity. Median longitudinal line evident at level of sternites 6, 7, 8, absent elsewhere. Press-button tubercle of sternopleonal locking mechanism located on posteromesial projection of sternite 5.

Chelipeds (P1), long, robust, surfaces tomentose; slightly heterochelous, major chela usually with modified cutting/crushing tooth proximally on cutting margin of dactylus. Merus long, with 4, sometimes 5, curved spines along flexor margin, and 2 distal spines on extensor margin; both margins densely setose. Carpus with sharp spine on inner angle, and flattened spine on external surface continuing as a strong carina, with additional, shorter carina above it. Dorsal surface of palm (propodus) with two straight, longitudinal granular crests, inner one distally ending distally in strong spine; small proximal spiniform tooth at articulation with carpus; two additional, curved granular crests on external surface of palm, first ending at level of articulation with dactylus, second, lower, ending near gape, creating cristate, proximo-ventral margin of palm; inner surface of palm with two wide, distinct rows of granules. Fingers generally straight except for curved, pointed tips; subequal in length to palm; with two granulate crests each on external and internal surfaces; lowest carina on fixed finger extending into palm; numerous teeth on cutting margins, arranged in groups so that each group has large central tooth flanked by smaller teeth of decreas-

ing size, giving the cutting margins appearance of having three or more denticulate, triangular lobes.

First to third ambulatory legs (P2–P4), long, slender; decreasing in length and size, with P2 largest, P4 smallest; flexor margins of meri, carpi, propodi and dactyli heavily setose. Fifth ambulatory (natatory) leg (P5) with quadrate merus, pentagonal carpus, flat, subrectangular propodus, and flat, oval dactylus; margins of articles regularly setose; propodus with four raised glabrous longitudinal bands, including flexor and extensor margins, interspersed with tomentum; dactylus with five raised glabrous bands, including flexor and extensor margins, interspersed with tomentum, distal third with low median crest continuing proximally as narrow tomentose stripe; in fresh specimens, P5 propodus with white band on postero-distal margin, no purple spot, P5 dactylus with small white spot on distal end.

Male pleon (Fig. 3A–C) ‘inverted T’-shaped, external surfaces mostly tomentose, 3rd–5th pleomeres fused. First pleomere very thin, less wide, mostly obscured by flange of posterior margin of carapace. Second pleomere much wider than first, lateral edges resting on P5 coxa, with prominent transverse keel running along entire width. Third pleomere widest, formed like a wedge, visible from both dorsal and ventral view; strong transverse crest somewhat forming posterior margin of cephalothorax, with shallow notch medially, posterolateral angles sharp, acute; sulcus between fused 3rd and 4th pleomere moderately deep, glabrous. Fourth pleomere subrectangular, wider than long, lateral margins convex, central region with low transverse crest. Fifth pleomere subtrapezoidal, basal margin wider than anterior. Sixth pleomere subrectangular; median length about 1.2 times maximum width; width at maximum lateral convexity greater than basal width; anterior margin concave, basal margin straight, lateral margins convex anteriorly, concave posteriorly. Telson subtriangular, apex rounded, lateral margins slightly concave, basal margin convex; median length 1.2 times basal width, with rounded tip, broadly rounded posterior margin.

G1 (Figs 4A–D) with proximal half relatively wide, somewhat flattened, strongly bent medially, by at least 45° but much less than 90°, distal half very slender, filiform; apically slightly recurved, rounded; distal tip much narrowed to small unarmed aperture. G2 about half length of G1, slender, distal tip minutely bilobed. Penis long, slender, uncalcified; emerging from sternocoxal condyle of P5.

Remarks. Following the recognition of *Monomia* Gistel, 1848, as a genus distinct from *Portunus* Weber, 1795 (see Mantelatto and Robles 2007; Mantelatto et al. 2009; Schubart and Reuschel 2009; Chertoprud et al. 2012; Spiridonov et al. 2014), the taxonomy of its type species, *Monomia gladiator* (Fabricius, 1798), needs to be assessed.

Firstly, there are five specimens identified as syntypes of *Portunus gladiator* Fabricius, 1798, in the Zoological Museum of the University of Copenhagen (ZMUC) (see <http://www.zmuc.dk/inverweb/invertebrater/Crustacea%20databases/Fabricius%20collection.htm>). All are dry specimens with the catalog numbers ZMUC-CRU 4704 through 4708 (see Ng et al. 2008). Examination of their photographs (available online) show that one of them is clearly not conspecific nor even congeneric (viz. ZMUC-CRU 4707). Accompanying this specimen in the photograph is a hand-

written label identifying it as “*Achelous Whitei* A. M. Edw.,” a name now considered a junior synonym of *Lupocycloporus gracilimanus* (Stimpson, 1858) (viz. Ng et al. 2008), a widespread species known from several localities in the Indo-West Pacific region; and a check with the available literature (e.g., Stimpson, 1907: pl. 10 fig. 3) confirms this identification. To stabilize the taxonomy of *Portunus gladiator* Fabricius, 1798, one of the other four syntypes, a male ZMUC-CRU 4705 (Fig 1A), is hereby designated as the lectotype, and the other three specimens (ZMUC-CRU 4704, 4706 and 4708) become paralectotypes.

Secondly, we agree with Ng et al. (2008) that there is no secondary homonymy between *Portunus gladiator* Fabricius, 1798, and *Cancer gladiator* Fabricius, 1793, because the two species were originally described in different genera, and the name “*Cancer gladiator* Fabricius, 1793” is no longer available by virtue of its synonymization under *Portunus sanguinolentus* (Herbst, 1783) (viz. Latreille 1825; Stephenson and Cook 1973). Although Chertoprud et al. (2012) cite the use of the name “*Portunus gladiator* (Fabricius, 1793)” by Stephenson and Cook (1973) as meeting the requirement of Article 59.1 of the Code, we view this as a misinterpretation of the Code’s provisions for homonymy. Furthermore, with the recognition of *Monomia* Gistel, 1848, as a full genus, thus absolutely eliminating any reservations concerning secondary homonymy, the species name *Monomia gladiator* (Fabricius, 1798) is, therefore, valid.

Thirdly, there is the matter of the confusion between *M. gladiator* and *M. haanii*. Stephenson and Cook (1973) proposed *Amphitrite haanii* Stimpson, 1858, as a replacement name for *P. gladiator* Fabricius, 1798, and this has led to the current confusion between the two names. As the following discussion will show, it is evident that *Monomia gladiator* and *M. haanii* (Stimpson, 1858) are distinct and valid species, and, therefore, the latter cannot be used as a replacement name for the former.

Monomia gladiator differs from *M. haanii* primarily in these three morphological characters: (1) in the fresh specimens of *M. gladiator*, there is a white band on the postero-distal margin of the P5 propodus, but no purple spot, and a small white spot on the distal tip of the P5 dactylus (Fig. 1C, D; also Chertoprud et al. 2012: pl. 51 fig. G) (vs. in *M. haanii*, there is a large purple spot on the distal tip of the P5 propodus, and the distal one-third of the P5 dactylus is colored purple; Fig. 1E, F; also Sakai, 1939: pl. 47 fig. 3; 1965: pl. 57, fig. 1; Chertoprud et al. 2012: pl. 51 fig. H); (2) the anterolateral margins of the male 6th pleomere are more flared out and rounded in *M. gladiator*, and at their widest extent exceed the basal width of the 6th pleomere (Fig. 3A–C; also Stephenson and Cook, 1973: fig.10A) (vs. straight, convergent anteriorly, and separated from concave posterolateral margin by an angular convexity in *M. haanii*, widest at base, Figs 2B, 3D–F); and (3) the G1 is bent at an angle of about 45° at mid-length and the distal tip is slender and relatively narrower in *M. gladiator* (Fig. 4A–D) (vs. G1 bent at or almost at 90°, and distal tip is somewhat wider than the immediate subdistal region in *M. haanii* (Figs 2C, 3E–H). Aside from these are some minor differences; for instance, the mesial part of the infraorbital margin of *M. gladiator* is granulate and there is no tooth on the mesial end as it abuts the basal article of

the antenna, there is, however, a large tooth immediately ventral to this margin, which projects outward and is visible from dorsal view as if it were part of the infraorbital margin. In *M. haanii*, the mesial end of the infraorbital margin has a large tooth which abuts against the basal article of the antenna. Also, the spines on the flexor margin of the P1 merus of *M. gladiator* tend to be more projecting and recurved than those of *M. haanii*, which are relatively smaller and less curved. Examination of available fresh-color photographs (viz. Chertoprud et al. 2012: pl. 51 Figs G, H) also show that these meral spines tend to be entirely white in *M. gladiator* while they are red at the base and white at the tips in *M. haanii*.

Furthermore, the molecular phylogenetic analysis corroborates the morphological evidence, clearly showing two distinct and well-supported clades corresponding to the two species. Specimens identified as *M. gladiator* based on the characters described above, including a topotypic specimen from India (ZRC.2016.0149), form a well-supported clade distinct from another clade containing specimens with the morphological attributes of *M. haanii*, which also includes a topotypic specimen from Japan (ZRC.2000.0084). Furthermore, the same molecular phylogenetic analysis shows that the specimen referred to by Chertoprud et al. (2012) as “*Monomia haanii*” (JX398095) falls within the *M. gladiator* clade, whereas the specimen referred to by Chertoprud et al. (2012) as “*Monomia pseudoargentata*” (JX398094) falls within the *M. haanii* clade. These specimens should now be identified as *M. gladiator* and *M. haanii*, respectively.

Finally, *Lupea gladiator* H. Milne Edwards, 1834 (Indian Ocean), is re-included in the synonymy of *Monomia gladiator* (Fabricius, 1898), and *Cancer menestho* Herbst, 1803 (probably from Indian Ocean) is hereby considered a junior subjective synonym of *Monomia gladiator* (Fabricius, 1798). We believe that Stephenson and Cook (1973) erred in removing these two names from the synonymy of *gladiator sensu* Fabricius, 1798. In the case of H. Milne Edwards’ specimen, the error is due simply to their conflation of *haanii* with *gladiator*. In the case of Herbst’s species, we disagree with them that the illustration of *Cancer menestho* does not show 2 spines on the posterior margin of the cheliped merus. What they call a “non-protruding” spine is an artifact of perspective. We have seen in our photographs of *M. gladiator*, that this second spine can appear non-protruding when the merus is viewed from directly above and if the marginal setae obscure its entire outline. Once this so-called difference is eliminated, there is no compelling reason why *C. menestho* should also not be treated as a synonym of *M. gladiator*.

***Monomia haanii* s. str. (Stimpson, 1858)**

Figs 1E, F, 2, 3D–F, 4E–H

Portunus (Amphitrite) gladiator: De Haan 1833: 39; 1835: pl. 1 fig. 5, pl. A. Non *Portunus gladiator* Fabricius, 1798.

Amphitrite haanii Stimpson, 1858: 38; 1907: 79; Ng et al. 2008: 151 (synonymy), 156 (discussion).

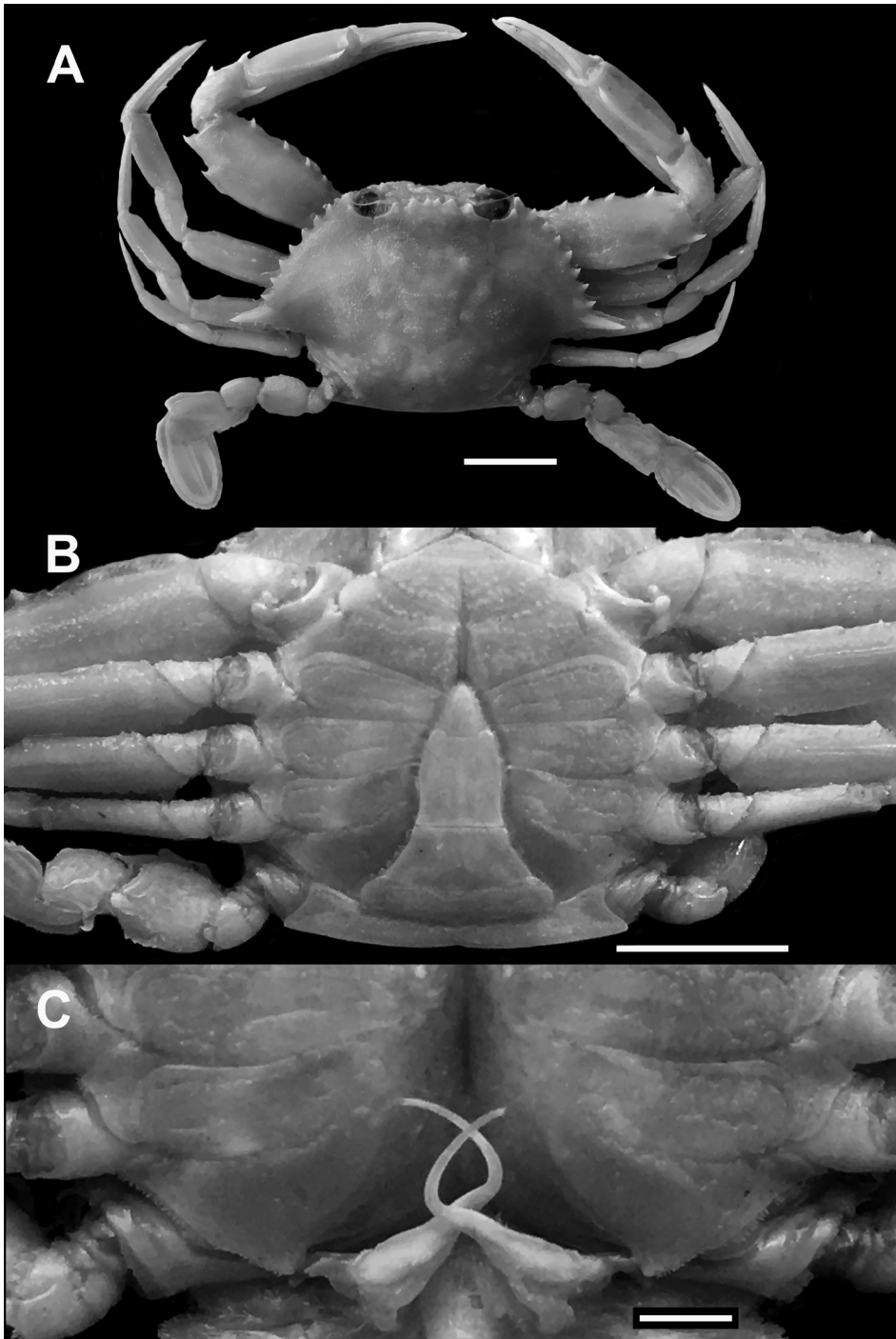


Figure 2. *Monomia haanii* (Stimpson, 1858), lectotype, male (RMNH 379), Japan. **A** dorsal habitus **B** thoracic sternum and pleon, ventral view **C** sterno-pleonal cavity and G1s, ventral view. Photographs by CHIJ Fransen (RMNH). Scale bars: A, B 10 mm; C 3 mm.

- Neptunus (Amphitrite) gladiator*: Ortmann 1893: 73; Lanchester 1902: 544; Parisi 1916: 173; Balss 1922: 107; Sakai 1934: 303; 1936: 129, pl. 36 fig. 3; 1939: 390, fig. 5a, pl. 47 fig. 3; Shen 1937: 101, fig. 2; Lin 1949: 19. Non *Portunus gladiator* Fabricius, 1798.
- Portunus gladiator*: Rathbun 1902: 26, Miyake 1961: 172; Sakai 1965: 118, pl. 57 fig. 1; 1976: 341, fig. 180a, pl. 120 fig. 1; Stephenson and Rees 1967: 24; 1968: 293; Takeda and Miyake 1968: 551; Stephenson 1972a: 16, 39; 1972b: 135; Ko and Lee 2013: 40, pl.-fig. 34. Non *Portunus gladiator* Fabricius, 1798.
- Portunus (Achelous) gladiator*; Rathbun 1910: 36 (from Gulf of Siam). Non *Portunus gladiator* Fabricius, 1798.
- Portunus pseudoargentatus* Stephenson, 1961: 109, Figs 2A, 3E, pl. 2 fig. 4, pl. 4 fig. F, pl. 5 fig. D; Stephenson and Rees 1967b: 25; 1968: 294; Yang and Dai 1994: 137, fig. 13.
- Portunus haanii*: Stephenson and Cook 1973 (in part): 429, figs 6F–H, 7F–H, 8F–H, 9B; Dai and Yang 1991: 223, fig. 120(2), pl. 27(4); Moosa 1996: 521; Apel and Spiridonov 1998 (in part): 291; Ng et al. 2001: 16 (list).
- Portunus (Monomia) gladiator*: Sakai 1976: 341, fig. 180, pl. 120 fig. 1; Kim and Chang 1985: 52. Not *Portunus gladiator* Fabricius, 1798.
- Portunus haanii*: Miyake 1983: 85, pl. 29 fig. 2; Takeda 1989: 152
- Portunus (Monomia) haanii* Yamaguchi & Baba, 1993: 396, figs 137A–C.
- Monomia pseudoargentata*: Chertoprud et al. 2012: 315, pl. 51 fig. H; Spiridonov et al. 2014: 412, 427, fig. 3I, tab. 1.
- Monomia haanii*: Ng et al., 2017: 68 (list).
- Not *Amphitrite media* Stimpson, 1858: 39; 1907: 79, pl. 10 fig. 1.
- Not *Monomia haanii*, Chertoprud et al. 2012: 314, pl. 51 fig. G (= *Monomia gladiator* (Fabricius, 1798)).

Material examined. JAPAN: RMNH 379, lectotype, male, Japan, coll. P.F. von Siebold, 1823–1829 (photographs only); ZRC.1999.0084, 1 male, 1 female, off Hota, Uchibo, coast of Boso Peninsula, coll. T Komai, 22 Aug. 1997; USNM5255, 4 male, 1 female, coll. FC Dale and PL Jouy, Palos R/V; USNM26254, 1 male, 1 female, off Wakanoura, coll. DS Jordan and JO Snyder, 1900; USNM45882, 1 female, off Wakanoura, Kishu; USNM54519, 1 female, Yamagata Prefecture, coll. M Sasaki, Aug. 1917; USNM72540, 2 males, Enoshima, Bay of Jeddo, coll. ES Morse; USNM60250, 1 male, Toyama Bay, coll. M Sasaki, 1925; USNM112423, 1 male, Shimizu, Sugura, Albatross R/V, 14 Oct. 1906.

AUSTRALIA: WAM-C7506, 1 male (holotype of *Portunus pseudoargentatus* Stephenson, 1961), Abrolhos Islands, Western Australia, coll. RW George, 11 May 1960; WAM-C34767, 1 female, Exmouth Gulf, Western Australia, coll. S Morrison and P Unsworth, 5 Nov. 2004; WAMC34900, 1 female, Exmouth Gulf, Western Australia, coll. S Morrison and P Unsworth, 12 Mar. 2004; WAM-C34938, 1 gravid female, Shark Bay, Western Australia, coll. S Morrison et al., 4 Oct. 2002; WAM-C5510, 1 male, Shark Bay, Western Australia, coll. S Morrison et al., 5 Mar. 2003; WAM-C44737, 1 female, Ningaloo Marine Park, Western Australia, coll. MP Salotti and SM Slack-Smith, 1 Feb. 2008.

SOUTH CHINA SEA (FAO Area 61): USNM1421161, 1 male, USNM1421181, 1 male, USNM1421182, 1 male, USNM1421185, 1 male, USNM1421187, 1 male, USNM1421191, 1 male, USNM1421194, 1 male, USNM1421195, 1 male, USNM1421186, 1 male, USNM121202, 1 male, USNM1421204, 1 male, USNM1421206, 1 male.

TAIWAN: UF29509, 1 female, Daxi Fishery Port; UF29511, 1 female, Daxi Fishery Port; UF29512, 1 female, Daxi Fishery Port; USNM1420827, 1 female, Daxi Fishery Port; USNM1420828, 1 male, Daxi Fishery Port; ZRC.1998.0186, 2 males, 1 female, Daxi Fishery Port, coll. PKL Ng, 3–4 Aug. 1996; ZRC.2016.0408, 4 males, 3 females, Daxi Fishery Port, coll. PKL Ng, 1 Jul. 2016.

Diagnosis. Similar to *Monomia gladiator* except in the following morphological characters. Infraorbital margin granulate, terminating mesially in small triangular tooth, in line with rest of margin. Sixth pleomere (Fig. 2B, 3D–F) longer than wide, maximum width at base; anterior half of lateral margins convergent anteriorly, posterior half concave to straight, anterior and posterior halves separated by angular convexity. G1 (Fig. 3C, 4E–H) bent at 90° at midlength, tapering distally toward a minutely spatulate tip, slightly broader than immediate subdistal area. In fresh specimens, P5 propodus with dark purple spot on distal tip, distal one-third of P5 dactylus colored dark purple as well (Fig. 1E, F).

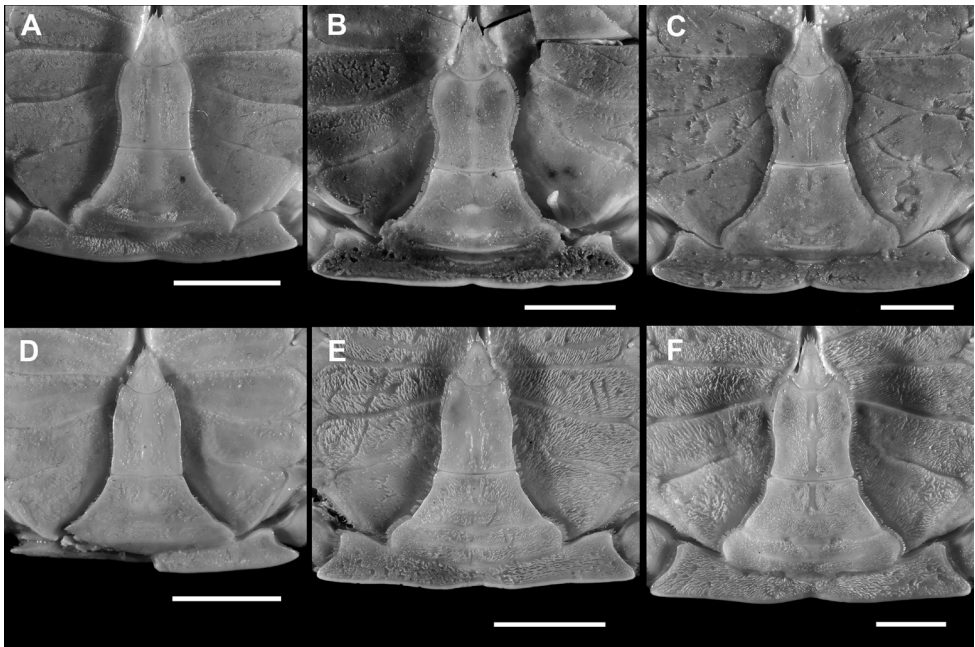


Figure 3. Male thoracic sternum and pleon. **A–C** *Monomia gladiator* (Fabricius, 1798), **A** ZRC 2018.1189, Jeppiar, Tamil Nadu, India **B** ZRC 2016.0145, Pazhayar Tamil Nadu, India; **C**) ZRC 2000.0842, Phuket, Thailand. **D–F** *Monomia haanii* (Stimpson, 1858) **D** WAM-C7506, holotype of *Portunus pseudoargentatus* Stephenson, 1961, Abrolhos Is., Western Australia **E** ZRC 1999.0084, Boso Peninsula, Japan **F** ZRC 2016.0408, Daxi Fishery Port, Taiwan. Scale bar: 10 mm.

Remarks. The primary morphological differences between *Monomia haanii* (Stimpson, 1858) and *M. gladiator* (Fabricius, 1798) have already been discussed in the Remarks for the latter species. Yamaguchi and Baba (1993) had previously reported on the type material of *Amphitrite haanii* Stimpson, 1858, listing several syntypes col-

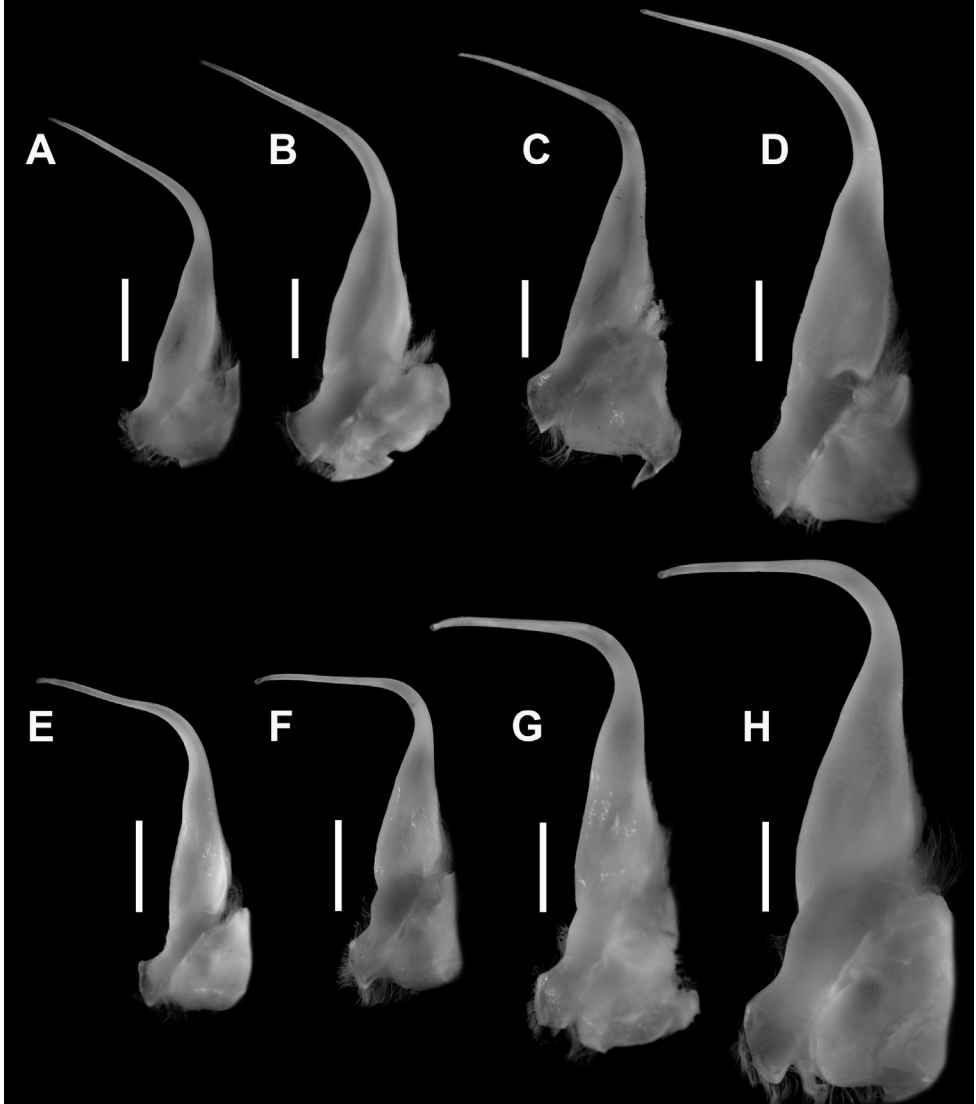


Figure 4. Left G1, sternal view (except D and E). **A–D** *Monomia gladiator* (Fabricius, 1798): **A**) ZRC 2018.1189, Jeppiar, Tamil Nadu, India **B**) ZRC 2016.0034, Tanintharyi coast, Myanmar **C**) ZRC 2016.0145, Pazhayar, Tamil Nadu, India **D**) ZRC 2000.0842, Phuket, Thailand (flipped right G1). **E–H** *Monomia haanii* (Stimpson, 1858) **E**) WAM-C7506, holotype of *Portunus pseudoargentatus* Stephenson, 1961, Abrolhos Is., Western Australia (flipped right G1) **F**) ZRC 1999.0084, Boso Peninsula, Japan **G**) ZRC 2016.0408, smaller male, Daxi Fishery Port, Taiwan **H**) ZRC 2016.0408, larger male, Daxi Fishery Port, Taiwan. Scale bar: 3 mm.

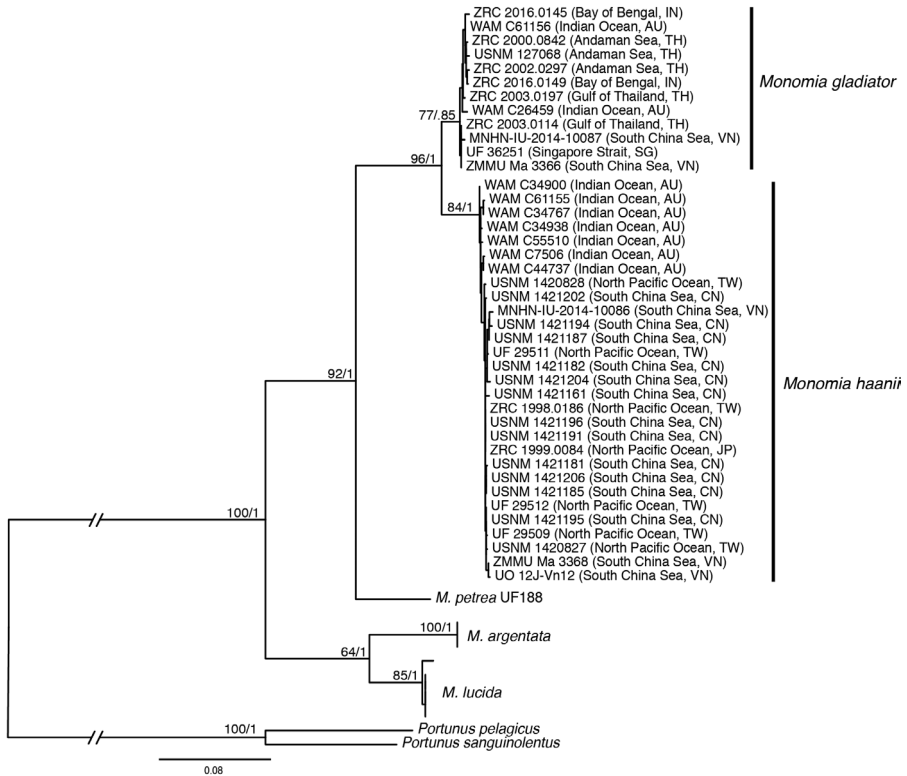


Figure 5. Maximum likelihood phylogram of three mitochondrial loci showing genetic distinction between *M. gladiator* and *M. haanii* along with three other members of the genus *Monomia*. Abbreviations: IOM = Institute of Oceanology and Museum, Nha Trang; MNHN = Muséum National d'Histoire Naturelle, Paris; NHM = The Natural History Museum, London; UF = University of Florida Natural History Museum, Gainesville; UO = University of Ostrava, Ostrava; USNM = United States National Museum, National Museum of Natural History, Washington, D.C.; WAM = Western Australian Museum, Perth; ZMMU = Zoological Museum of the Moscow University, Moscow; ZRC = Zoological Reference Collection, Lee Kong Chian Natural History Museum, Singapore; CN = China; IN = India; JP = Japan; SG = Singapore; TH = Thailand; TW = Taiwan; VN = Vietnam.

lected by P.F. von Siebold from Japan in the years 1823 to 1829, deposited in the then Rijkmuseum van Natuurlijke Historie (RMNH) in Leiden, and preserved either dry or in alcohol. From these they selected a lectotype, a young male (42 by 24 mm; RMNH 379) preserved in alcohol. No further or detailed descriptions of the material were provided. Although they did provide photographs of several specimens, the lectotype was photographed still inside the bottle (viz. Yamaguchi and Baba, 1993: fig. 137A) and no definitive or diagnostic features of its morphology could be discerned. The collections of the RMNH are now housed in the Naturalis Biodiversity Center, Leiden, and photographs of the lectotype (Fig. 2) were kindly provided to the authors by Dr. Charles Fransen. From these photographs, the diagnostic morphology of the 6th pleomere (Fig. 2B) and the 90-degree bend of the G1 (Fig. 2C) can already be observed even in such

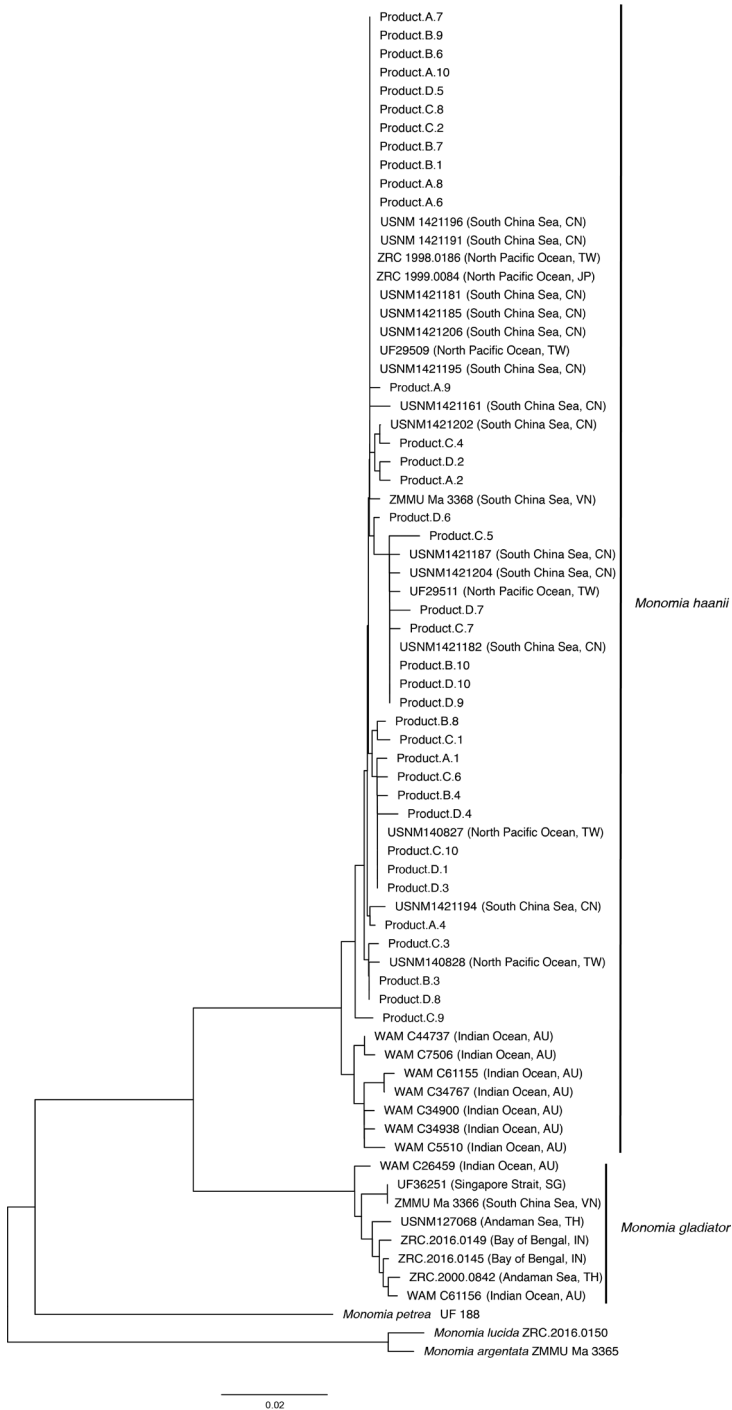


Figure 6. Neighbor joining phylogram of the barcode region of COI to visualize relationships between museum-vouchered reference specimens and 40 samples from four cans of pasteurized lump crabmeat (Product A–D) labeled “*Portunus haanii*” and/or “red swimming crab”

a young specimen. Furthermore, although the fresh coloration of this species was not recorded by either De Haan (1833) or Stimpson (1858), later observations of topotypic material show that the purple spots on the P5 propodus and dactylus, as well as the more profuse but scattered spotting on the dorsal carapace, and the dark colored spines on the anterior margin of the P1 merus are consistently observed (Fig. 1E, F; also Sakai, 1976: pl. 120 fig. 1; Miyake, 1983: pl. 29 fig. 1).

Stephenson and Cook (1973) had also previously synonymized *Portunus pseudoargentatus* Stephenson, 1961, under *P. haanii* on the basis of their similar morphology. We confirm that *P. pseudoargentatus* is a junior subjective synonym of *Monomia haanii* (Stimpson, 1858) *sensu stricto* on the basis of similarities in the dentition of the infraorbital margin, the shape of the male 6th pleomere (Fig. 3D), and the ~90° angle of the bend of the G1 at midlength (Fig. 4E; also Stephenson, 1961: fig. 2A). In a photograph of the presumably newly preserved holotype, the pigmented spots on the left P5 propodus and dactylus are still there (Stephenson, 1961: pl. 2 fig. 4), although these have since faded and can no longer be seen during the present examination of the holotype.

Specimens morphologically identifiable as *M. haanii* comprise a highly-supported clade that includes specimens from Japan, the type locality of *M. haanii*, as well as the holotype of *Portunus pseudoargentatus* Stephenson, 1961, and the Vietnamese specimens referred to by Chertoprud et al. (2012) (JX398094) and by Koch et al. (2017) and Koch and Duris (2018) (KY524463, KY524464) as "*Monomia pseudoargentata*". Both morphological and molecular phylogenetic analyses support the recognition of *Monomia haanii* as a full species. Molecular results also support a morphological basis for synonymy of *P. pseudoargentatus* Stephenson, 1961, under *M. haanii*.

Discussion

The need to address and resolve the *Portunus gladiator* species complex was brought about largely because of the incongruence in taxon names utilized by the scientific community and the seafood industry. This incongruence was highlighted by Warner et al. (2015) who matched sequences from crab cakes to a single sequence of "*P. pseudoargentatus*" and raised concerns that this was a species in the US food supply unknown to regulators. Subsequently, "*P. pseudoargentatus*" was added to The Seafood List (Food and Drug Administration 2015), based on that report. Pasteurized, lump crab meat labeled as "Portunus haanii" or "red swimming crab" is routinely imported into the United States; however, the species name *P. haanii* has been synonymised under *P. gladiator* for some time (Ng et al., 2008). As it turned out, this complex had a very complicated taxonomic past, but here we have used traditional morphological methods in concert with molecular phylogenetic analyses to establish the morphological and molecular boundaries between the two species we recognize herein, *Monomia gladiator* and *Monomia haanii*.

Revising and describing the morphological differences between these two species was necessary to verify the identity of the individual specimens used to generate reference DNA sequences for identification of picked crab meat samples labelled as "*Portunus haanii*." The morphological findings of two distinct species were corroborated

in our multi-locus phylogenetic analysis that showed complete congruence between morphologically derived identity and genetic clade membership. This reciprocally informative approach has enabled us to confirm that the commercial products that we tested, that were labelled and sold as *Portunus haanii*, were in fact *Monomia haanii* and should be labelled as such. Based on our findings, the FDA's seafood labeling guidance to industry, The Seafood List, can be emended to reflect this current understanding of the species in question.

Commercially important species are often presumed to be well understood because they have tangible value, but in the case of decapod crustaceans molecular phylogenetic analyses are re-writing much of what carcinologists thought they knew about species-, genus-, and family-level relationships (Keenan et al. 1998; Ma et al. 2009; Lai et al. 2010; Bracken-Grissom et al. 2013; de Carvalho et al. 2013; Windsor and Felder 2014; Evans 2018; Tavares and Santana 2018). For example, Lai et al. (2010) determined that what was then known as *Portunus pelagicus* was actually a complex of four species; and the status of the genus *Penaeus* Fabricius, 1798, has been highly debated in the literature (Pérez Farfante and Kensley 1997; Lavery et al. 2004; Dall 2007; Flegel 2007; 2008; McLaughlin et al. 2008; Ma et al. 2009; Ma et al. 2011). These name changes are to be expected as new methods of DNA analysis are developed and applied to illuminate and clarify evolutionary relationships.

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Appendix I

The types of *Monomia gladiator* (Fabricius, 1798) are currently deposited in the Natural History Museum of Denmark (NHMD) in Copenhagen. The lectotype (ZMUC-Cru 4705) has the corresponding catalog number, NHMD-82551. The paralectotypes (ZMUC-Cru 4704, 4706, and 4708) have the corresponding catalog numbers, NHMD-82550, -82552, and -82554, respectively (J. Olesen pers. comm.)