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The taxonomic status of the headshield slug genus *Nakamigawaia* Kuroda and Habe, 1961 (Gastropoda: Cephalaspidea: Aglajidae), with the description of a new species from the Western Pacific

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ABSTRACT

Nakamigawaia is a poorly understood genus of Aglajidae sea slugs with only two species formally ascribed. In this paper we explore new morpho-anatomical characters using stereo and scanning electron microscopy and employ different molecular approaches (a cytochrome *c* oxidase sub-unit I gene phylogeny, the Automatic Barcode Gap Discovery species delimitation method, and genetic distances) to compare specimens across the geographical span of the genus and from two distinct chromatic morphotypes occurring in the Western Pacific (blackish morph and white-dotted morph). Our results support the conspecificity of these two morphs and show they belong to an undescribed species here named *Nakamigawaia nakanoae* sp. nov. The species differs from the type species of the genus, *N. spiralis*, by the presence of a distinct open-dilated shell and differs from its Western Atlantic congener *N. felis* by subtle differences in the shell, male reproductive system and caudal lobes. Genetically (COI uncorrected *p*-distance) the two species (*N. nakanoae* and *N. felis*) are 18.8–20.1% distinct. The definition of the genus *Nakamigawaia* is discussed and the current assignment to the latter of lineages other than the type species is questioned.

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biodiversity; DNA barcoding; Mollusca; morphology; phylogeny; Western Pacific

Introduction

The family Aglajidae is the second most diverse of the gastropod Heterobranchia order Cephalaspidea, with an estimated 85 valid species worldwide divided into 15 genera (Zamora-Silva and Malaquias 2018). These marine slugs inhabit tropical and temperate waters predominantly at shallow depths where they are found crawling on sandy bottoms, coral rubble and algae (Burn and Thompson 1998; Gosliner et al. 2008; Camacho-García et al. 2014; Zamora-Silva and Malaquias 2018), yet at least three species have been reported from deeper bathymetries including abyssal depths (Zamora-Silva and Malaquias 2018; Chaban et al. 2019). Aglajids are predators feeding mostly upon vagile prey such as other sea slugs, nematodes, polychaetes and small fish (Zamora-Silva and Malaquias 2016).

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One of the genera of Aglajidae is the enigmatic *Nakamigawaia* Kuroda and Habe, 1961. The genus was introduced for the species *N. spiralis* Kuroda and Habe, 1961 from Japan (type locality: Kasajima beach, Sagami Bay) and originally included in the family Chelidonuridae Kuroda and Habe 1961 (presently regarded as a synonym of Aglajidae; MolluscaBase 2021). Kuroda and Habe (1961) provided only a brief description, in Japanese, of the colouration of the animal and an illustration of the shell. The definition of the genus and species remained largely elusive, and, for example, Rudman (1978) and Gosliner (1980) did not consider this genus in their dedicated accounts on the systematics of the aglajid slugs.

Baba (1985) was the first author to provide a detailed description of the type species *N. spiralis*, recognising at the same time similarities to representatives of the Aglajidae genus *Melanochlamys*, which led the author to firmly suggest the inclusion of the genus *Nakamigawaia* in the family. The validity of *Nakamigawaia* and its inclusion in the Aglajidae have been corroborated by Camacho-García et al. (2014) and Zamora-Silva and Malaquias (2018) based on molecular phylogenetics. In addition, Camacho-García et al. (2014) first suggested a close relationship between *N. spiralis* (based on specimens from Papua New Guinea and the Philippines) and the morphologically similar tropical western Atlantic species '*Aglaja*' *felis* described by Marcus and Marcus (1970) from Puerto Rico, which led the authors to reassign the latter species to the genus *Nakamigawaia*.

Despite the putative phylogenetic evidence for a common generic affiliation of the Atlantic and Pacific species, the occurrence of substantial differences between their shells (convex, dilated with an open whorl in '*A. felis*', and spiralled in *N. spiralis*), led Ortea et al. (2014) to argue these species warranted a different generic assignment, and introduced the genus name *Migaya* for the Atlantic lineage. However, Zamora-Silva and Malaquias (2018), based on extended genetic and taxon sampling, supported the validity of *Nakamigawaia* for both Atlantic *N. felis* and Pacific *N. spiralis* and argued that different shell types could be present in the same genus, and considered *Migaya* a junior synonym of *Nakamigawaia*.

Specimens from the western Atlantic have consistently been ascribed to the species '*felis*', but interestingly those from the Pacific have been named erratically by authors and attributed either to the genus *Chelidonura* or to *Nakamigawaia*, resulting in pervasive taxonomic confusion and a still unsettled nomenclature, with the name '*spiralis*', in fact, being hardly used. For example, Susuki (2000, p. 16) and Ono (1999, p. 14, 15) identified specimens from the Izu Peninsula and Kerama Islands (Japan) as *Chelidonura* sp.; and Gosliner et al. (2008, p. 39) labelled specimens from the Philippines and Papua New Guinea as *Nakamigawaia felis*, whereas the same authors later used the name *Nakamigawaia spiralis* (Gosliner et al. 2015, p. 45) and *Nakamigawaia* sp. (Gosliner et al. 2018, p. 367). In 2018, Nakano (2018, p. 55; 2019, p. 55) identified specimens from Japan as *Nakamigawaia felis*.

In addition, two distinct chromatic morphs occur in the western Pacific, namely one that is consistent with the original description and having a uniform blackish pattern (Kuroda and Habe 1961; Baba 1985), and the other characterised by a white-dotted pattern on a brownish background (Ono 1999; Gosliner et al. 2008, 2015, 2018; Nakano 2018, 2019). In this work we use morpho-anatomical characters and a molecular phylogenetic framework to test the taxonomic affiliation of these two morphs and of specimens from the western Atlantic in order to shed light on the diversity and systematics of the genus *Nakamigawaia*.

Material and methods

Taxon sampling and literature review

Samples were obtained from the collections of the Department of Natural History, University Museum, University of Bergen, Norway (ZMBN). Specimens of *Nakamigawaia* from Taiwan (two black morph specimens and two white-dotted morph specimens), Japan (black morph), Venezuela and the Bahamas were used for morphological studies, for a total of 11 specimens (six of those sequenced for their DNA; [Table 1](#)). In addition, we obtained from GenBank 16 sequences of the cytochrome *c* oxidase subunit I gene (COI), representing the genera *Chelidonura*, *Nakamigawaia*, *Philinopsis* and *Tubulophilinopsis*. The tree was rooted with the species *Philine quadripartita* ([Table 1](#)).

Table 1. List of specimens used for molecular analysis, including locality, voucher numbers and GenBank accession numbers. *Listed as *Aglaja felis* in GeneBank; ** listed as *Nakamigawaia spiralis* in GenBank. Novel sequences generated for this study are depicted in bold font.

Taxa	Sample ID	Voucher number	Locality	COI gene GenBank Accession number
* <i>Nakamigawaia felis</i>		ZMBN84913a	Venezuela Isla: Tortuga	MF036535
* <i>Nakamigawaia felis</i>		CASIZ175655	The Bahamas: Great Exuma, Stocking Island	JN825179
* <i>Nakamigawaia felis</i>		CASIZ175653	The Bahamas: Great Exuma, Stocking Island	JN825178
<i>Nakamigawaia felis</i>	EH05	ZMBN84913.1	Venezuela: Isla Tortuga	OK094481
<i>Nakamigawaia felis</i>	EH06	ZMBN84913.2	Venezuela Isla Tortuga	OK094482
* <i>Nakamigawaia felis</i>		CPIC-00684	The Bahamas: Great Exuma, Stocking Island	JN825180
* <i>Nakamigawaia felis</i>		CPIC-00685	The Bahamas: Great Exuma, Stocking Island	JN825181
** <i>Nakamigawaia nakanoae</i> sp. nov.		ZMBN95949	Australia: Lizard Island	MF036536
* <i>Nakamigawaia</i> sp.		CASIZ173495	The Philippines: Bohol Island, Panglao	JN825177
<i>Nakamigawaia nakanoae</i> sp. nov.	EH01	ZMBN116778.1	Taiwan: Shadao, Kenting National Park, Pingtung County	OK094484
** <i>Nakamigawaia nakanoae</i> sp. nov.		ZMBN95960	Australia: Lizard Island	MF036537
<i>Nakamigawaia nakanoae</i> sp. nov.	EH03	ZMBN116777.1	Taiwan: Shadao, Kenting National Park, Pingtung Count	OK094486
<i>Nakamigawaia nakanoae</i> sp. nov.	EH02	ZMBN116778.2	Taiwan: Shadao, Kenting National Park, Pingtung County	OK094485
<i>Nakamigawaia nakanoae</i> sp. nov.	EH04	ZMBN116777.2	Taiwan: Shadao, Kenting National Park, Pingtung County	OK094483
<i>Philinopsis speciosa</i>		ZMBN95993	Japan: Kyoda Beach, Okinawa	MF036556
<i>Philinopsis pusa</i>		ZMBN95958	The Bahamas: Great Exuma, Stocking Island	MF036552
<i>Chelidonura hirundinina</i>		ZMBN95971	Australia: Lizard Island	MF036524
<i>Chelidonura varians</i>		ZMBN95978	Australia: Lizard Island	MF036530
<i>Chelidonura alisonae</i>		ZMBN95991	Hawaii: Anini Beach, Kauai	MF036523
<i>Tubulophilinopsis pilsbryi</i>		ZMBN95948	Australia: Lizard Island	MF036569
<i>Tubulophilinopsis lineolata</i>		ZMBN95941	Australia: Lizard Island	MF036565
<i>Philine quadripartita</i>		-	?Mediterranean Sea	KU557520

DNA extraction, amplification and sequencing

DNA was extracted from tissue clipped from the parapodial lobes of the animals following the protocol of the Qiagen DNeasy Blood and Tissue Kit. Sequences of the gene COI were amplified and sequenced using the universal primers by Folmer et al. (1994). Polymerase chain reactions (PCRs) were performed in 25 μL Eppendorf tubes containing 1 μL of DNA, 2.5 μL Qiagen Buffer, 2.5 μL dNTPs, 5 μL Qiagen Q-Solution, 2 μL forward and reverse primers (1 μL for each primer direction), 0.15 μL of Qiagen HotStar TAQ DNA polymerase, 8.35 μL of Sigma water, and 3.5 μL of magnesium chloride. The PCR thermal cycle included an initial denaturation at 95°C for 5 min, followed by 39 cycles of 45 s at 94°C, 45 s at 45°C (annealing temperature), extension at 72°C for 2 min, and a final extension at 72°C for 10 min.

PCRs that did not yield results were redone with the restriction enzyme Takara and the primers GasF1_t1 (TGTA AACGACGGCCAGTTTCAACAAACCATAARGATATTGG) and GasR1_t1 (CAGGAAACAGCTATGACACTTCWGGRTGHCCRAARAATCARAA) (Stein et al. 2013). In those cases the PCR was started with an initial denaturation step of 5 min at 95°C, followed by 5 cycles of 40 s at 94°C, 40 s at 45°C and 1 min at 72°C, then 35 cycles of 40 s at 94°C, 40 s at 51°C and 1 min at 72°C. This was followed by a final extension step of 5 min at 72°C.

Gel electrophoresis was performed on a 1% agarose/buffer to assess the quality and quantity of the amplified DNA. The successful PCR products were purified with Exonuclease I Shrimp Alkaline Phosphatase (ExoSAP). The total volume of each purification reaction was 10 μL , which consisted of 0.1 μL of Exo I (10 U/ μL), 1 μL SAP (1 U/ μL), 0.9 μL of Sigma water and 8 μL of PCR product. The mixtures were incubated at 37°C for 30 min, followed by 85°C for 15 min, and finally kept cool at 4°C.

Sanger sequencing reactions were prepared using primers diluted to 3.2 μM . Each sequencing reaction contained 6.5 μL Sigma water, 1 μL buffer, 1 μL BigDye, 1 μL primer and 0.5 μL PCR product (or 1 μL PCR product for samples with a low amount of DNA, with no changes made to the amount of Sigma water). Prior to sequencing the mixtures were run in a thermal cycle including an initial step at 96°C for 5 min, followed by 25 cycles at 96°C for 10 sec, 5 sec at an annealing temperature of 50°C, and 60°C for 4 min, and finally kept cool at 4°C. After the thermal cycle 10 μL Sigma water was added to each sample. Samples were sequenced with an ABI 3730XL DNA Analyser (Applied Biosystems) at the DNA Sequencing Facility, Department of Biological Sciences, University of Bergen, Norway.

Phylogenetic and species delimitation analyses

Both forward and reverse DNA chromatograms were edited and assembled using Geneious R11 (Biomatters, Auckland, New Zealand; Kearse et al. 2012). All sequences were blasted using GenBank to test for contamination. Sequences were aligned with the programme Muscle (Edgar 2004) implemented in Geneious using default parameters. The alignment was trimmed at both ends where at least 50% of the individual sequences had nucleotides and was translated into amino acids to test for the presence of stop codons. The best-fit model of evolution was estimated with JModelTest 2.1.10 (Darriba et al. 2012).

Bayesian molecular phylogenetic inference was done in the program MrBayes 3.2.7 (Ronquist and Huelsenbeck 2003) through the platform CIPRES Science Gateway V.3.3 (Phylo.org) with three parallel runs of five million generations with sampling every 100 generations. Node support was assessed using Bayesian posterior probabilities (Felsenstein 1985). Convergence of runs was assessed using Tracer 1.7.1 (Rambaut and Drummond 2007) with the burn-in set to 25%. The phylogenetic tree was visualised in FigTree 1.4.4 (Rambaut and Drummond 2009) and edited in Adobe Illustrator.

Uncorrected p -distances between and within species and colour morphs were estimated with the programme MEGA X (Kumar et al. 2018). The Automatic Barcode Gap Discovery method (ABGD; Puillandre et al. 2012), implementing the Simple Distance algorithm and default settings, was used to aid in defining candidate species.

Anatomical work and scanning electron microscopy

A total of 11 specimens (three representing the black morph and two the dotted-white morph of *N. 'spiralis'*, and six representing *N. felis*) were dissected for structures of taxonomic relevance, namely the male reproductive system and the shell. Drawings of soft anatomical structures were made under a stereomicroscope with the aid of a *camera lucida*. Shells were cleansed with diluted commercial bleach, rinsed in distilled water and mounted on metallic stubs covered with carbon sticky tabs, coated with gold–palladium and later studied and imaged with a Fei Quanta 450 scanning electron microscope at the Electron Microscopy Laboratory, Faculty of Mathematics and Natural Sciences, University of Bergen, Norway.

Results

Phylogenetic analyses

The monophyly of the genus *Nakamigawia* received marginal support in the COI phylogenetic analysis. Three sub-clades were rendered within *Nakamigawaia*: a clade with Caribbean specimens compatible with the species *N. felis*, a clade with western Pacific specimens (Taiwan and Lizard Island, Australia) with a mix of black and dotted-white animals that we provisionally assigned to *N. 'spiralis'* (= *N. nakanoae* sp. nov.; name adopted hereafter; see Taxonomic section), and a single specimen from the Philippines (*Nakamigawaia* sp.; based on GenBank sequences) that split off alone and probably corresponds to an unnamed species.

The COI genetic uncorrected p -distance between black and white-dotted morphs of *N. nakanoae* sp. nov. was estimated at 0–0.4%, which – like the phylogenetic results – suggests these two morphs are conspecific. The genetic distance within specimens of *N. felis* was 0–0.7%, and within specimens of *N. nakanoae* sp. nov. it was 0–1.2%. The genetic distance between *N. felis* and *N. nakanoae* sp. nov. was 18.8–20.1%. The ABGD analysis rendered 11 candidate species including *N. felis* and *N. nakanoae* sp. nov. (see Supplementary material).

Taxonomic section**Genus NAKAMIGAWAIA Kuroda and Habe, 1961*****Nakamigawaia felis* (Er. Marcus and Ev. Marcus, 1970)**

(Figure 1(b), 3(e), (f), 4(b))

Diagnosis

External colouration plain black. Internal shell dilated with an open whorl extending half a turn, protoconch dorsally covered by teleoconch, a plate-like structure can project ventrally near protoconch. Caudal lobes of nearly equal length. Male reproductive system with pyriform penial chamber and tubular, long, folded prostate. Prostate about 3 times longer than penial chamber.

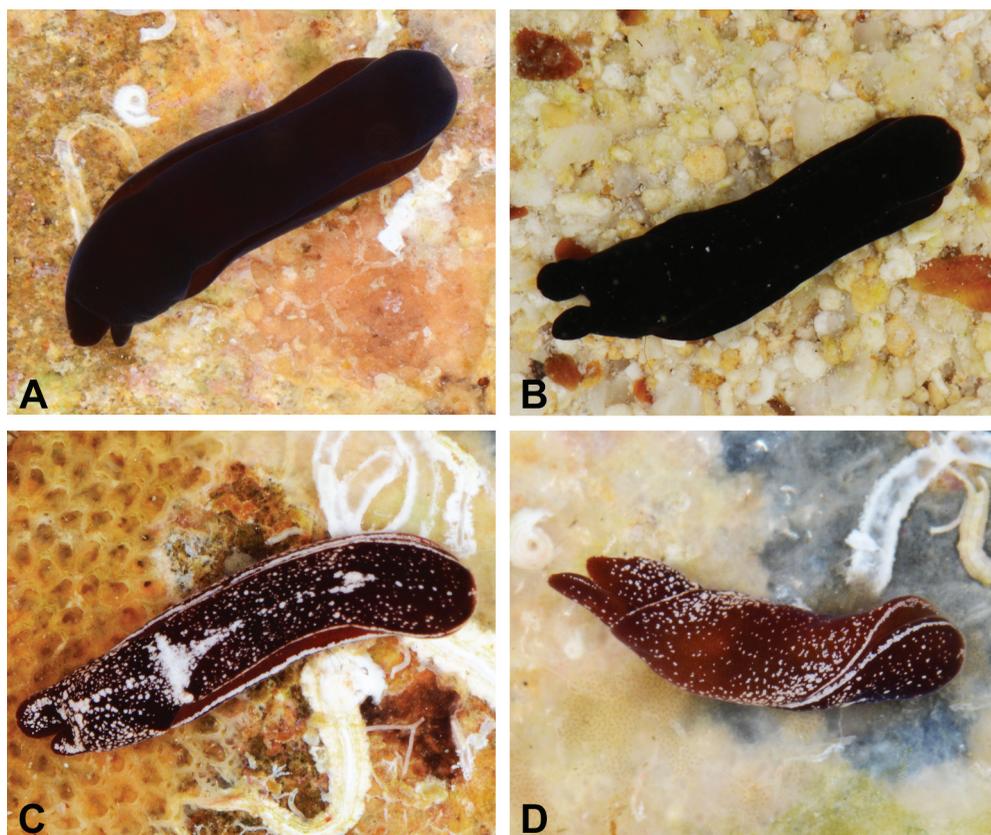


Figure 1. Live images of *Nakamigawaia nakanoae* sp. nov. (A, C, D) and *Nakamigawaia felis* (B). A, black morph, Taiwan, ZMBN 116778, animal length (L) = 10 mm (paratype). B, the Bahamas, ZMBN 91108, L = 13 mm. C–D, white dotted morph, Taiwan, ZMBN 116777, L = 8 mm (paratype).

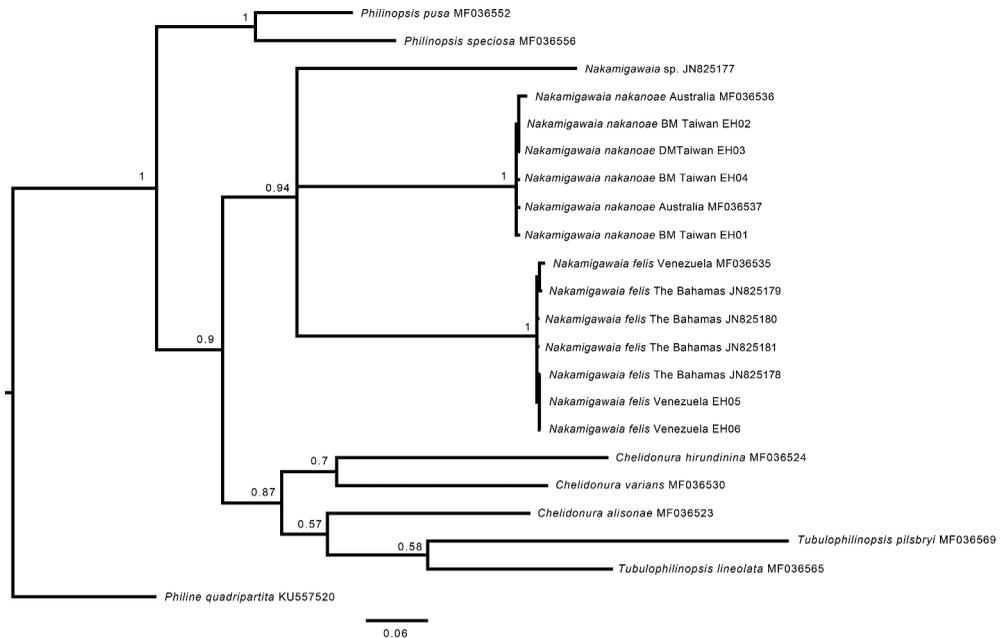


Figure 2. Bayesian phylogeny of Aglajidae species with a focus on the genus *Nakamigawaia*, based on the mitochondrial Cytochrome *c* oxidase subunit I (COI) gene. Numbers on branches are posterior probabilities. Tree rooted with *Philine quadripartita*. BM = black morph. DM = white dotted morph.

Examined material

Venezuela, Isla Tortuga, Playa El Yaque's Lagoon, 2 specimens dissected and sequenced, ZMBN 84913, animal length (L) = 13 mm, coll. Manuel Malaquias, 16 March 2010. The Bahamas, Eleuthera Island, Savannah Sound, 4 specimens dissected, ZMBN 91108, L = 10–13 mm, coll. Manuel Malaquias, 14 April 2013.

External morphology (Figure 1(b)): Plain black including foot. Caudal lobes about the same length.

Shell (Figure 3(e), (f)): Convex, dilated, with an open whorl extending half a turn. Protoconch smooth, partly visible ventrally; dorsally fully covered by extension of teleoconch; a plate-like structure can project ventrally near protoconch (fragile, breaks off easily).

Male reproductive system (Figure 4(b)): Penial chamber pyriform, rounded distally and funnel-like proximally. Length between genital opening and insertion of prostate around 1 mm. Prostate tubular, cylindrical, folded; length around 3 mm.

Ecology

Found crawling on sand between 0.2 and 1 m depth.

Distribution

Caribbean Sea, between the Bahamas in the north and Venezuela in the south (Malaquias 2014; Caballer et al. 2015).

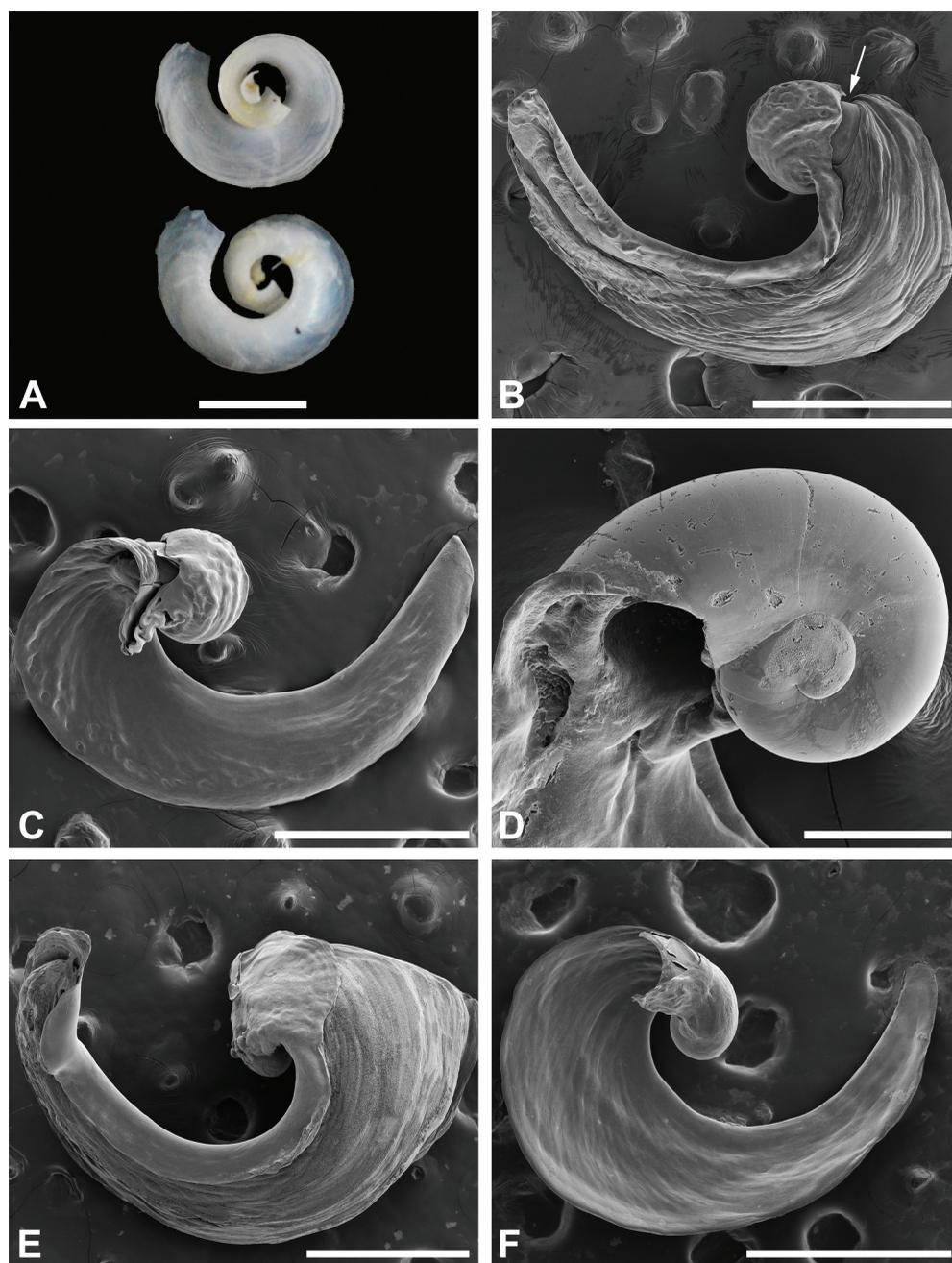


Figure 3. Shells of *Nakamigawaia* species. A, *N. spiralis*, holotype, NSMT-39805, ventral (upper) and dorsal (lower) views (macrophotograph), Japan. B, *N. nakanoae* sp. nov., micrograph of dorsal view; arrow pointing to protoconch, ZMBN 132073, animal length (L) = 7 mm (fixed length), Japan. C, *N. nakanoae* sp. nov., micrograph of ventral view, ZMBN 132073, L = 7 mm (fixed length), Japan. D, *N. nakanoae* sp. nov., micrograph of protoconch observed from ventral side of shell, ZMBN 116778, L = 10 mm, Taiwan. E, *N. felis*, micrograph of dorsal view, ZMBN 91108, L = 13 mm, the Bahamas. F, *N. felis*, micrograph of ventral view, ZMBN 91108, L = 12 mm, the Bahamas. Scale bars: a = 200 μ m; b, c, e, f = 500 μ m; d = 100 μ m.

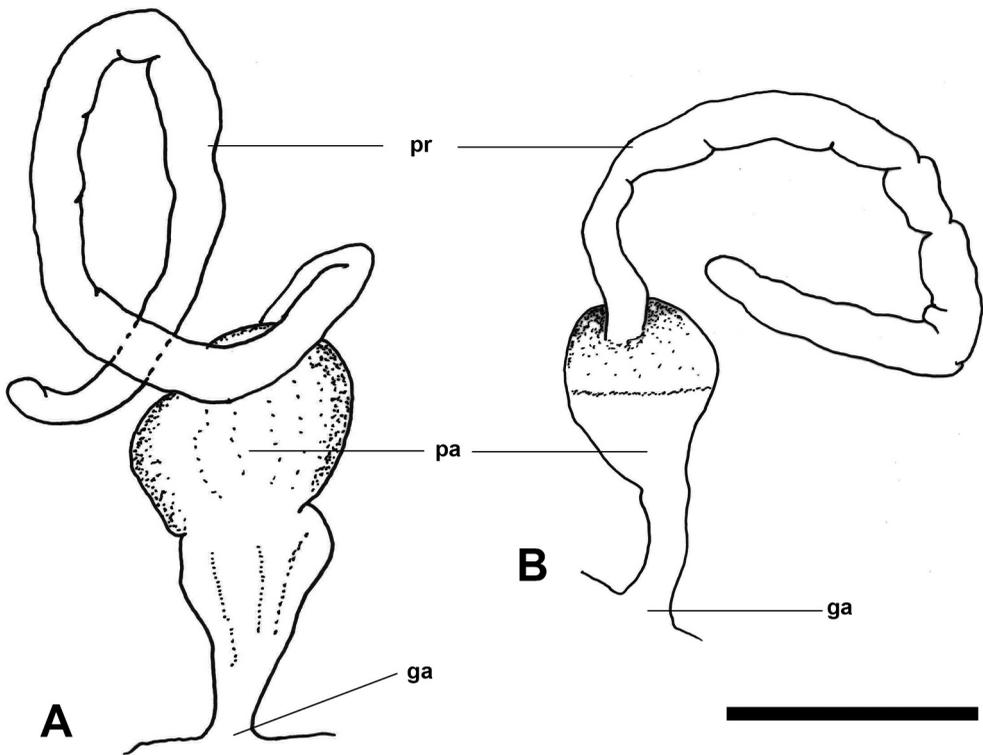


Figure 4. Male reproductive system of (A) *Nakamigawaia nakanoae* sp. nov. and (B) *N. felis*. A: Taiwan, ZMBN 116778, animal length (L) = 10 mm. B: the Bahamas, ZMBN 91108, L = 13 mm. ga, genital aperture; pa, penial atrium; pr, prostate. Scale bar: 1 mm.

Nakamigawaia nakanoae Hellem and Malaquias sp. nov.
(Figures 1(a), (c), (d), 3(b)–(d), 4(a))

Zoobank: [lsid:zoobank.org:act:EB686586-9EFD-448A-B743-15C0FDC583DF](https://doi.org/10.21203/rs.3.rs-1000000/v1)

Diagnosis

External colouration plain black or brownish dotted in white. Internal shell dilated with an open whorl extending half a turn, protoconch dorsally visible near transition to teleoconch, otherwise covered by teleoconch layer; ventrally a funnel-like structure can be present near protoconch. Right caudal lobe nearly half length of left lobe. Male reproductive system with pyriform penial chamber and tubular, long, folded prostate. Prostate about 5 times longer than penial chamber.

Type locality

Taiwan, Pingtung County, Kenting National Park, Shadao (21.912233N, 120.846961E).

Etymology

The species is named after Dr Rie Nakano, for her passion and contributions to the study of the sea slug molluscs of Japan and for her continuous support throughout the years of our research work.

Examined material

Taiwan, Pingtung County, Kenting National Park, Shadao, 1 specimen, ZMBN 130245 (black morph), L = 10 mm, holotype, coll. Manuel Malaquias, 7 May 2017. Taiwan, Pingtung County, Kenting National Park, Shadao, 2 specimens, ZMBN 116814 (black morph), L = 10 mm, paratypes, coll. Manuel Malaquias, 7 May 2017. Taiwan, Pingtung County, Kenting National Park, Shadao, 2 specimens dissected and sequenced plus 46 observed, ZMBN 116778 (black morph), L = 8–10 mm, paratypes, coll. Manuel Malaquias, 7 May 2017. Taiwan, Pingtung County, Kenting National Park, Shadao, 2 specimens dissected and sequenced plus 5 observed, ZMBN 116777 (white dotted morph), L = 8 mm, paratypes, coll. Manuel Malaquias, 7 May 2017. Japan, Kagoshima, Amami-Oshima Island, Ayamaru, 1 specimen dissected, ZMBN 132073 (black morph), L = 7 mm (fixed length), coll. Rie Nakano, 19 May 2015.

External morphology (Figure 1(a), (c), (d)): Plain black including foot, or brownish background dotted in white, with rim of head-shield and parapodial lobes white or dashed-white; foot sole brownish, white-dotted. Caudal lobes asymmetrical, with right lobe nearly half length of left lobe.

Shell (Figure 3(b)–(d)): Convex, dilated, with an open whorl extending half a turn. Protoconch smooth, entirely or partly visible ventrally; dorsally visible only near transition to teleoconch; a funnel-like structure (fragile, breaks off easily) can be present ventrally near protoconch.

Male reproductive system (Figure 4(a)): Penial chamber pyriform, rounded distally and funnel-like proximally. Length between genital opening and insertion of prostate around 1 mm. Prostate tubular, cylindrical, folded; length around 5 mm. One specimen from Japan (ZMBN 132073) has a 7.5 mm prostate.

Ecology

Specimens were found at depths between 1 and 6 m crawling on white sandy bottoms.

Distribution

Kenting in Taiwan (current study), Kerama Islands, Amami-Oshima Island and Izu Peninsula in Japan (Ono 1999; Susuki 2000), the Philippines and Papua New Guinea (Gosliner et al. 2008), and in Lizard Island, eastern Australia (current study).

Discussion

There is widespread confusion and inconsistency about the naming of slugs with black-coloured headshields occurring in the western Pacific (Kuroda and Habe 1961; Baba 1985; Ono 1999; Gosliner et al. 2008, 2015, 2018; Nakano 2018, 2019) and controversy regarding the definition of the genus *Nakamigawaia* (Ortea et al. 2014; Zamora-Silva and Malaquias 2018). The original description in Japanese of the type species *N. spiralis* is brief and includes a single illustration of the shell (Figure 3A). The description refers to animals with

black body colour and a shell coiling in three turns, opening downwards. Specimens were collected by Mr Korokuro Nakamigawa at 'Zushi beach' [= Kasajima], Kanagawa Prefecture, in a tide pool (Kuroda and Habe 1961; Baba 1985).

The first comprehensive study of *N. spiralis* was performed by Baba (1985). The author provided detailed data on the morphology and anatomy and sound evidence of its taxonomic affinity with the family Aglajidae. Striking features of the description by Baba (1985) are the comparatively large size of the animals (ranging between 25 and 35 mm) and the long prostate (about 20 mm for a specimen 25 mm in length overall; measurements based on [fig. 4C](#) in Baba 1985).

In our preliminary identification of black headshield slugs collected in southern Taiwan and southern Japan we assigned these specimens to *N. spiralis*. However, detailed comparative studies of the shell and reproductive system revealed substantial differences between what was observed and what is known about these structures in *N. spiralis*. The shells in our specimens were dilated, with an open whorl extending half a turn ([Figure 3](#)), and there were differences in the soft anatomy: prostates were shorter, ranging between 3.5 and 6.5 mm in specimens that measured about 10 mm in total length when alive. However, the difference in the size of the reproductive system might be explained by the unique length of the animals. Interestingly, such slugs with large black headshields have not been observed or reported during at least the last decade (Rie Nakano, personal communication). Therefore, based on differences in the shell shape and apparent dissimilarities in parts of the male reproductive system, we assign our specimens to a new species, named *Nakamigawaia nakanoae* sp. nov. The known distribution of *N. spiralis* from Sagami Bay to the Kyusyu region (Kuroda and Habe 1961; Baba 1985) suggests a species of temperate affinity, whereas *N. nakanoae* sp. nov. is confined to tropical waters.

This study confirms the validity of the species *N. felis* and its tropical western Atlantic geography, and suggests the occurrence of an additional species of *Nakamigawaia* present in the Philippines at least. No phylogenetic or genetic differentiation was detected between the white-dotted and black morphs of *N. nakanoae* sp. nov. (COI uncorrected *p*-distance ranged between 0% and 0.4%), whereas the large genetic difference between the latter species and *N. felis* (18.8–20.1%) suggests an old speciation event or fast evolutionary rates in these species ([Figure 2](#)).

Externally, *N. felis* and the black morph of *N. nakanoae* sp. nov. are nearly indistinguishable, but there is one subtle difference: namely, the right and left caudal lobes in the latter species are of different lengths, whereas in *N. felis* they are of similar size ([Figure 1](#)). Also, the internal shells of these two species are similar, but in *N. felis* the dorsal part of the protoconch is covered by a teleoconch layer, whereas in *N. nakanoae* sp. nov. part of the protoconch is dorsally visible ([Figure 3](#)). The male reproductive systems are also similar in morphology, yet the ratio between the length of the prostate and the length of the penial chamber (from the genital aperture to the insertion point of the prostate) is about 5× in *N. nakanoae* sp. nov. and only 3× in *N. felis* ([Figure 4](#)).

The systematics of *Nakamigawaia* and its recognised lineages (*N. spiralis*, *N. felis*, *N. nakanoae* sp. nov., *Nakamigawaia* sp. from the Philippines) admittedly remains elusive after this work. Our COI gene tree barely recovers support for the monophyly of the genus, but this is not surprising considering that it is a single-locus phylogeny. On the other hand, the works of Camacho-García et al. (2014) and Zamora-Silva and Malaquias (2018), based on

a broader character set, strongly support the monophyly of *Nakamigawaia*. However, if the genus *Nakamigawaia* is to be defined by the presence of a spiralled internal shell and a long and convoluted prostate, then perhaps *N. felis*, *N. nakanoae* sp. nov. and eventually *Nakamigawaia* sp. will have to be assigned to a distinct genus. In this case, the junior synonym name *Migaya* Ortea, Caballer and Espinosa, 2014 is available. Nevertheless, answering this question warrants the study of new specimens of the true *N. spiralis* with spiralled shells and its comparison in a phylogenetic framework with other aglajids including the species currently assigned to *Nakamigawaia*. Until additional evidence is available, we believe that the use of the generic name *Nakamigawaia* for these lineages better promotes taxonomic stability.

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