

Revision of the genus *Cuvierina* Boas, 1886 based on integrative taxonomic data, including the description of a new species from the Pacific Ocean (Gastropoda, Thecosomata)

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Abstract

Shelled pteropods (Gastropoda, Thecosomata, Euthecosomata) are a group of holoplanktonic gastropods that occur predominantly in the surface layers of the world's oceans. Accurate species identifications are essential for tracking changes in species assemblages of planktonic gastropods, because different species are expected to have different sensitivities to ocean changes. The genus *Cuvierina* has a worldwide warm water distribution pattern between $\sim 36^{\circ}\text{N}$ and $\sim 39^{\circ}\text{S}$. Based on an integrative taxonomic approach combining morphometric, genetic, and biogeographic information, the two subgenera of *Cuvierina*, *Cuvierina s. str.* and *Urceolarica*, are rejected. A new species is introduced: *Cuvierina tsudai* **sp. n.**, which has to date been considered the same species as *Cuvierina pacifica*. *Cuvierina tsudai* **sp. n.** is endemic to the Pacific Ocean and is characterised by a shell height of 7.2–8.0 mm, a moderately cylindrical shell shape, the absence of micro-ornamentation and a triangular aperture. *Cuvierina pacifica* is restricted to the centre of the oligotrophic southern Pacific gyre, has a shell height of 6.6–8.5 mm, a more cylindrical shell shape, no micro-ornamentation and a less triangular aperture than *C. tsudai* **sp. n.**

Keywords

Integrative taxonomy, DNA barcoding, geometric morphometrics, pteropods, biogeography

Introduction

Pteropods are holoplanktonic heterobranch gastropods classified in a superorder comprised of the orders Thecosomata and Gymnosomata, commonly referred to as “sea butterflies” and “sea angels”, respectively (Lalli and Gilmer 1989, Pierrot-Bults and Peijnenburg 2015). The order Thecosomata consists of Euthecosomata that have sinistrally coiled or straight, bilaterally symmetrical shells, and Pseudothecosomata that have either sinistrally coiled shells, an internal gelatinous pseudoconch, or are shell-less in the adult stage (Meisenheimer 1905, Tesch 1913). Pteropods play an important role in marine food webs (Jörger et al. 2010), and although most species occur in warm tropical and subtropical waters, the highest abundances have been observed for some (sub)polar cold water species (Bé and Gilmer 1977, Van der Spoel and Heyman 1983, Bednaršek et al. 2012, Burridge et al. 2016). Because of their thin-walled, aragonite shells, euthecosomes are exceptionally vulnerable to the effects of ocean acidification (e.g., Fabry et al. 2008, Bednaršek and Ohman 2015, Gattuso et al. 2015, Moya et al. 2016).

The genus *Cuvierina* is a remarkable group of shelled pteropods with relatively large (5.1–11.1 mm), straight, bottle-shaped shells (Janssen 2005). Ever since *Cuvierina* was described as a mollusc genus (as *Cuvieria* Rang, 1827, emended by Boas 1886), it has often been considered to consist of a single species, *C. columnella* (Rang, 1827), the type species of the genus by monotypy. The first taxonomic division within the genus came with the description of a second *Cuvierina* species, introduced as *Cuvieria urceolaris* (Mörch, 1850), but in later literature it was often interpreted as a form or subspecies of *C. columnella* (e.g., Tesch 1913, Van der Spoel 1967, Rampal 1975). A third form, *Cuvierina columnella* f. *atlantica*, was described by Van der Spoel (1970), and validated as a taxon of the species group by Bé et al. (1972). Bé and Gilmer (1977) interpreted the morphological differences between the three taxa as infraspecific variability. Contrarily, Rampal (2002) distinguished these taxa as independent species but introduced the taxon *C. spoeli* to replace the taxonomically invalid *Cuvierina columnella* f. *atlantica*. Because the holotype of *C. spoeli* was from the Indian Ocean, where *C. atlantica* is absent, it rather represented *C. columnella* and was rejected as a valid species by Janssen (2005). Two further extant species, *C. cancapae* and *C. pacifica*, were described by Janssen (2005).

According to the most recent taxonomic revision of *Cuvierina*, five extant species were assigned to two subgenera based on shell morphology and supposed lineages of fossil occurrences since the early Miocene (Janssen 2005, 2006). The subgenus *Cuvierina* s. str. consisted of *C. atlantica*, *C. columnella*, and *C. pacifica*, which are characterised by relatively slender, cylindrical shells, triangular rather than kidney-shaped apertures and the presence (*C. columnella*) or absence (*C. atlantica*, *C. pacifica*) of micro-ornamentation. Two geographical varieties were recognised within *C. pacifica*, one from the North Pacific and the other from the South Pacific, but were not formally introduced as new species. The subgenus *Urceolarica*, containing *C. cancapae* and *C. urceolaris*, is characterised by more inflated, bottle-shaped rather than

cylindrical shells, pronounced micro-ornamentation, and kidney-shaped rather than triangular apertures.

All extant *Cuvierina* species are restricted to the surface layers of tropical and subtropical waters from -45°N to -40°S . In the Atlantic Ocean, *C. atlantica* occurs in the subtropical gyres and *C. cancapae* is found in tropical waters. In the Indian Ocean, *C. columnella* is found in the southern subtropical zone and *C. urceolaris* occurs in tropical waters and further south along Madagascar towards South Africa. *Cuvierina columnella* and *C. urceolaris* also occur in the Pacific Ocean along with *C. pacifica* (Janssen 2005, BurrIDGE et al. 2015).

BurrIDGE et al. (2015) examined the diversity, distribution, and evolution of *Cuvierina* taxa using integrative geometric morphometric, molecular, and biogeographic methods. They confirmed that the five species described for *Cuvierina* species have significantly different shell shapes and that *C. pacifica* consists of two disjunct morphometric groups, registered as *C. pacifica* N and *C. pacifica* S in their study. Three genetic lineages were distinguished based on mitochondrial Cytochrome Oxidase I DNA: the Atlantic lineage with *C. atlantica* and *C. cancapae*, the Indo-Pacific lineage with *C. columnella*, *C. urceolaris*, and *C. pacifica* N, and the South Pacific lineage with *C. pacifica* S. A new taxonomic description of *C. pacifica* N is required because the holotype of *C. pacifica* has the shell shape of *C. pacifica* S.

Based on the findings of Janssen (2005) and the integrative approach of BurrIDGE et al. (2015) the taxonomy of the genus *Cuvierina* is revised. The subgenera *Cuvierina* s. str. and *Urceolarica* are rejected, a new species, *C. tsudai*, is described from the Pacific Ocean, and the species description of *C. pacifica* is restricted to the South Pacific lineage. A taxonomic key is provided for the identification of *Cuvierina* species.

Methods

Two approaches were used to distinguish between *C. tsudai* and *C. pacifica* based on differences in shell shape. First, simple measurements of shell height and width, aperture diameters, and position of maximum shell width as applied to museum specimens by Janssen (2005) were used to distinguish between *C. tsudai* and *C. pacifica*. Second, geometric morphometric data of shell shapes in ventral and apertural orientations were used for 168 adult specimens of *Cuvierina* that were registered as *C. pacifica* N or *C. pacifica* S in BurrIDGE et al. (2015). The specimens corresponded to museum specimens as identified and measured by Janssen (2005, N = 92), additional museum specimens (N = 24), and recently collected fresh specimens (N = 52). Geometric morphometric methods consisted of digitising shell outlines using tpsDig and tpsUtil (Rohlf 2006) to contain 76 ventral and 37 apertural semi-landmarks per shell, after which a generalised least square Procrustes superimposition was applied (GLS, Kendall 1977 in Zelditch et al. 2004) to rotate, translate, and scale the semi-landmark coordinates. A subsequent thin-plate spline (TPS) analysis (e.g., Zelditch et al. 2004) provided centroid sizes, a size measure depending on surface area, and multiple relative

warp axes per specimen, containing information on shape. To describe the new species *C. tsudai* as well as to reject the validity of the *Cuvierina* subgenera, Cytochrome Oxidase I mitochondrial (COI) DNA and 28S ribosomal DNA sequence data from Burridge et al. (2015) were used.

Results and discussion

Distinction between *Cuvierina tsudai* and *C. pacifica*

Cuvierina tsudai and *C. pacifica* are similar in size but have different shell shapes, COI mtDNA and 28S rDNA. Because of their Pacific distributions and similarities in shell size, *C. tsudai* and *C. pacifica* have to date been considered the same species. Although Janssen (2005) demonstrated their presence as morphological varieties within *C. pacifica*, the congruence between morphometric and genetic differentiation supports the separation into two species (Figs 1A–J and 2, fig. 4 in Burridge et al. 2015). Shell heights of *C. tsudai* specimens are between 7.2 and 8.8 mm, showing a large overlap with *C. pacifica*, which measures between 6.6 and 8.5 mm (Janssen 2005). However, in terms of shell shape, *C. pacifica* and *C. tsudai* are significantly different (fig. 29 lower left in Janssen 2005, Burridge et al. 2015). The shell of *C. tsudai* is wider (more inflated) than the slender and more cylindrical *C. pacifica* (Fig. 2). *Cuvierina pacifica* has a larger height/width-ratio between 3.25 and 3.96 (mean 3.50) compared to *C. tsudai*, which has a ratio between 2.77 and 3.46 (mean 3.14). The position of maximum shell width is located at 34–45% (mean 40%) of the shell height from the septum upwards for *C. pacifica* and at 33–42% (mean 37%) for *C. tsudai* (Janssen 2005). The aperture of *C. tsudai* is wider, more triangular and more concave on the ventral side than in *C. pacifica*. The overall shape variation is larger for *C. tsudai* than for *C. pacifica* (Fig. 2). The average pairwise genetic distance of COI mtDNA (658bp fragment) between *C. tsudai* (N = 16) and *C. pacifica* (N = 43) is 4.5%. The genetic variation of COI within *C. tsudai* is 1.6% compared to 0.8% within *C. pacifica*. The 28S rDNA fragment (965bp) of *C. tsudai* differs at least at one position compared with other *Cuvierina* species, except for *C. columnella* (Burridge et al. 2015).

The larger genetic and shell shape variation for *C. tsudai* compared to *C. pacifica* coincides with a much larger Pacific distribution and lower ecological specificity of *C. tsudai*. *Cuvierina pacifica* is restricted to the centre of the oligotrophic southern Pacific gyre and occupies a more specialised ecological niche based on ecological niche modelling (ENM) than *C. tsudai* (Burridge et al. 2015). This study used presence-only data and six uncorrelated environmental parameters, of which ocean surface temperature and chlorophyll *a* concentration were the most important. The distribution of *C. tsudai* was mostly explained by maximum monthly sea surface temperatures (SST) and near-surface chlorophyll *a* concentrations (both 30.8%). The distribution of *C. pacifica* was best explained by low maximum monthly chlorophyll *a* concentrations (57.1%).

Description of *Cuvierina tsudai* sp. n.**Superfamily Cavolinioidea Gray, 1840****Family Cuvierinidae Gray, 1840****Genus *Cuvierina* Boas, 1886 (= replacement name for *Cuvieria* Rang, 1827 non Lesueur & Petit, 1807, pl. 30 (Coelenterata))****Type species.** *Cuvieria columnella* Rang, 1827, p. 323, pl. 45 figs 1–3, by monotypy.***Cuvierina tsudai* sp. n.**<http://zoobank.org/B33A28E9-BCDE-4F2B-9349-F3E18CCD87BE>*Cuvieria columnella* Rang, 1827: 323 (partim).*Cuvierina columnella*: Boas 1886: 132, 217, pl. 6 fig. 95g (partim, non Rang); Rampal 2002: 214 (partim, non Rang).*Cuvierina columnella* (Rang, 1827) forma *columnella* (Rang, 1827) – Van der Spoel 1967: 79 (partim, non Rang); Van der Spoel 1970: 120, fig. 19 (partim, non Rang).*Cuvierina* (*Cuvierina*) *pacifica* Janssen, 2005: 46 figs. 18–20 (partim, northern Pacific specimens only, non figs. 14–17 = *C. pacifica*).*Cuvierina pacifica* N (Janssen, 2005): BurrIDGE et al. 2015: 5, fig. 2.**Holotype.** RMNH.5004167, also see Fig. 1A and Table 1.**Type locality.** 8°47'N, 158°49'W.**Paratypes.** See Fig. 1B–I and Table 1 for all specimen information. Three specimens from the type locality (RMNH.5004168); three specimens from the Zoological Museum of the University of Copenhagen, Denmark (ZMUC, not registered) illustrated by Janssen (2005, figs. 18–20); five specimens from four locations (RMNH.5004169–72) studied by BurrIDGE et al. (2015, referred to as *C. pacifica* N therein). The latter five specimens have COI mtDNA and 28S rDNA sequences available at GenBank (see Table 1).**Additional material examined.** Specimens recorded as *C. pacifica* from the North Pacific Ocean in Janssen (2005: 49, 71), housed in the Muséum National d'Histoire Naturelle (MNHN, Paris, France) and ZMUC (Copenhagen, Denmark). Specimens from BurrIDGE et al. (2015), referred to as *C. pacifica* N in Table S1 therein, with photographs deposited at the Dryad repository (<http://dx.doi.org/10.5061/dryad.7n1q4>) and COI mtDNA (KP292730–72) and 28S rDNA sequences (KP292636–42) deposited at GenBank. These specimens are housed in Naturalis Biodiversity Center (Leiden, The Netherlands) and ZMUC (Copenhagen, Denmark). Registration numbers, if available, from Janssen (2005).**Diagnosis.** Shell moderately small, adult specimens 7.2–8.8 mm high, height/width-ratio 2.77–3.46 (mean 3.14), position of maximum shell width 33–42% (mean 37%) of shell height from septum upwards. Aperture triangular. No longitudinal micro-ornamentation.

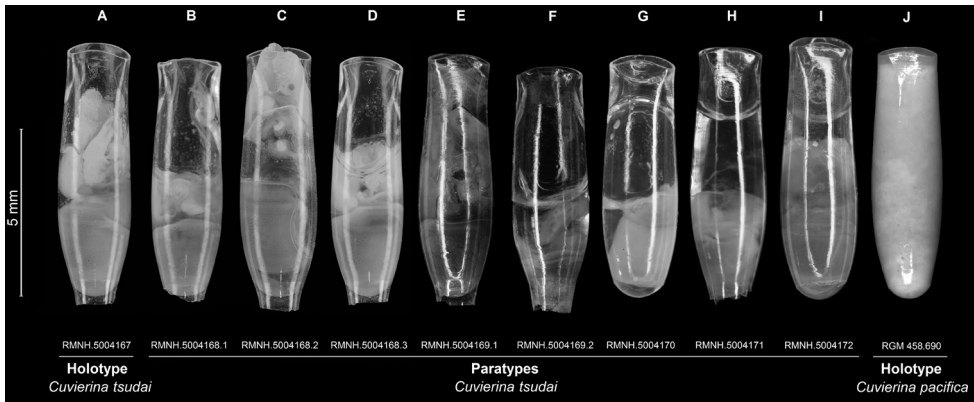


Figure 1. Holotype and paratypes of *C. tsudai* and holotype of *C. pacifica*. **A** Holotype (RMNH.5004167) and **B–I** paratypes (RMNH.5004168–72) of *C. tsudai* and **J** holotype of *C. pacifica* (RGM 458.690) photographed in a ventral view. Photographs of RMNH.5004169–72 from Burridge et al. (2015); RMNH.5004167–68 taken by R. van der Hulst and RGM 458.692 taken by E.F. de Vogel, this study. RMNH = Naturalis Biodiversity Center, mollusc collection and RGM = Naturalis Biodiversity Center, fossil planktonic mollusc collection, Leiden.

Description. The shell shape of *Cuvierina tsudai* differs from other *Cuvierina* species. Its shell height is smaller than in *C. columnella*, *C. cancapae*, and *C. atlantica*, but larger than in *C. urceolaris*, and of similar size compared to *C. pacifica*. The position of maximum shell width is distinctly higher than for *C. columnella* and *C. atlantica* and lower than for *C. pacifica*. It is more cylindrical in shape than the inflated (bottle-shaped) *C. urceolaris* but less cylindrical than *C. atlantica* and *C. pacifica*. It differs from *C. urceolaris* and *C. cancapae* by the absence of micro-ornamentation. It has a more triangular and wider aperture than *C. urceolaris* and *C. pacifica* (Fig. 3, Janssen 2005, Burridge et al. 2015).

Distribution. *Cuvierina tsudai* has a wide, exclusively Pacific distribution between 36°N and 39°S, in which it co-exists with *C. columnella*, *C. urceolaris*, and *C. pacifica*. It has been found most often in the North Pacific, but also occurs in the South Pacific. It has not been found thus far in the central, oligotrophic parts of the South Pacific subtropical gyre, the southeast Pacific, the coral triangle west of the Philippines or southwest of Papua New Guinea.

Etymology. Named after Atsushi Tsuda, professor in biological oceanography at the University of Tokyo, Japan, for sending us pteropod samples from the Pacific Ocean and in recognition of his services to the zooplankton research community.

Rejection of the subgenera in *Cuvierina*

Two subgenera of *Cuvierina* were described that supposedly evolved since the early Miocene (Aquitian, 23 million years ago): *Cuvierina s. str.*, with extant species *C.*

Table 1. Voucher and sampling information of type specimens of *C. tsudai* including the holotype of *C. pacifica*.

Museum voucher	Image voucher	Collection date	Latitude	Longitude	Cruise	Station	COI GenBank	28S GenBank	First studied
Holotype of <i>C. tsudai</i>									
RMINH.5004167	C_PNE_SE1201_21_01	2012-05-15	8°47'N	158°49'W	SE1201	21			This study
Paratypes of <i>C. tsudai</i>									
RMINH.5004168.1	C_PNE_SE1201_21_02	2012-05-15	8°47'N	158°49'W	SE1201	21			This study
RMINH.5004168.2	C_PNE_SE1201_21_03	2012-05-15	8°47'N	158°49'W	SE1201	21			This study
RMINH.5004168.3	C_PNE_SE1201_21_04	2012-05-15	8°47'N	158°49'W	SE1201	21			This study
RMINH.5004169.1	C_PNE_KH1110_08_01	2011-12-19	22°47'N	158°06'W	KH-11-10	8	KP292730	KP292636	Burridge et al. 2015
RMINH.5004169.2	C_PNE_KH1110_08_20	2011-12-19	22°47'N	158°06'W	KH-11-10	8	KP292748	KP292637	Burridge et al. 2015
RMINH.5004170	C_PNE_KM1109_02_02	2011-03-04	21°14'N	158°11'W	Kilo Moana 1109	2	KP292755	KP292639	Burridge et al. 2015
RMINH.5004171	C_PNE_KM1109_08_01	2011-03-06	21°20'N	158°22'W	Kilo Moana 1109	8	KP292759	KP292640	Burridge et al. 2015
RMINH.5004172	C_PNW_TMKT1020_05_01	2010-09-29	27°08'N	125°33'E	R/V Tansai-Maru KT-10-20	5	KP292766	KP292642	Burridge et al. 2015
ZMUC, not registered	figure 18	1933-08-21	33°45'N	137°30'W	DANA	4794			Janssen 2005
ZMUC, not registered	figure 19	1934-02-12	32°56'N	131°50'W	DANA	4807			Janssen 2005
ZMUC, not registered	figure 20	1929-05-25	20°04'N	125°59'E	DANA	3718 V			Janssen 2005
Holotype of <i>C. pacifica</i>									
RGM 458.692	figure 15	1986-04/05	18°39'S	172°12'W	Manihiki Plateau Expedition	U351a			Janssen 2005

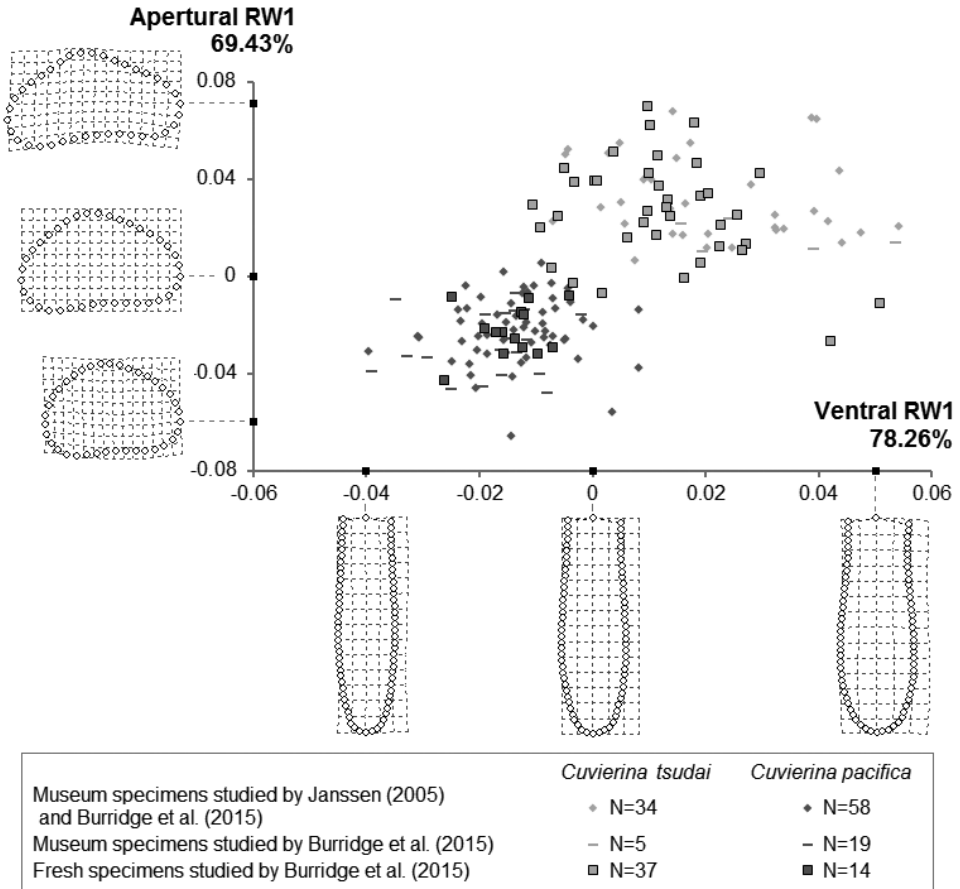


Figure 2. Shape variation in *C. tsudai* and *C. pacifica* by means of Relative Warp (RW) data. Ordination of RW data of *C. tsudai* and *C. pacifica* for the first ventral and apertural RWs (N = 167 excluding 1 specimen with only one orientation). On the X-axis, RW1 depicts 78.26% of the total ventral shape variation. On the Y-axis, 69.43% of the apertural shape variation is explained by its RW1. Shape variations depicted by ventral and apertural RW1 (with subsequent RWs = 0) are shown.

atlantica, *C. columnella*, and *C. pacifica*, and *Urceolarica* with extant species *C. cancapae* and *C. urceolaris* (see Janssen 2005, 2006). They were based on distinguishing shell characteristics in fossil species such as the position of maximum shell width, aperture shape and presence or absence of micro-ornamentation. However, the morphology and molecular phylogenetic information of recent species are in conflict with this separation. *Cuvierina columnella*, typically a *Cuvierina s. str.* species, has distinct micro-ornamentation, which was considered one of the distinguishing characters of the subgenus *Urceolarica*. It was shown that there are three divergent and well-supported lineages based on genetic data: the Atlantic (*C. atlantica* and *C. cancapae*), Indo-Pacific (*C. columnella*, *C. urceolaris* and *C. tsudai*), and South Pacific (*C. pacifica*) lineages (fig. 4 in Burridge et al. 2015). Hence, we reject the two subgenera within *Cuvierina*.



Figure 3. Typical specimens of six *Cuvierina* species.

Taxonomic key to *Cuvierina* pteropods

The following taxonomic key identifies adult *Cuvierina* pteropod species based on distinctive shell shape characteristics and shell sizes. Photographs of typical adult shells are shown in Fig. 3.

- 1 Micro-ornamentation present 2
- Micro-ornamentation absent 4
- 2 Strongly inflated shell shape, shell height 5.1–6.7 mm..... *C. urceolaris*
- Moderately inflated or cylindrical shell shape, shell height 7.5–11.1 mm.... 3
- 3 Cylindrical shell shape, shell height 8.8–11.1 mm *C. columnella*
- Moderately inflated shell shape, shell height 7.5–9.3 mm *C. cancapae*
- 4 Cylindrical shell shape and triangular aperture, shell height 6.7–10.5 mm....
..... *C. atlantica*
- Moderately inflated or cylindrical shell shape, triangular to kidney-shaped
aperture, shell height 6.6–8.8 mm 5
- 5 Cylindrical shell shape and kidney-shaped aperture, shell height 6.6–8.5 mm....
..... *C. pacifica*
- Moderately inflated shell shape and triangular aperture, shell height 7.2–
8.8 mm..... *C. tsudai*

Conclusions

Morphometric, genetic, and biogeographic information has led to the introduction of a new species of the warm water pteropod genus *Cuvierina* and the rejection of its

subgenera. We encourage a combined evidence approach of taxonomy to more accurately identify species boundaries and higher taxonomic relationships in planktonic gastropods. Accurate taxonomic identification is a prerequisite to assess to what extent species are affected by ocean changes and to potentially use them as bioindicators.

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