

RESEARCH NOTE

**Asymmetry of eye peduncle length in *Euprotomus iredalei* (Abbott, 1960)
(Mollusca: Neostromboidae: Strombidae)**

Stephen J. Maxwell^{1,*}, Jessica L. Watt², Mark A. Payne³

¹College of Business Law and Governance, James Cook University, Cairns, QLD 4878, Australia.

²College of Public Health, Medical and Veterinary Sciences, James Cook University, Smithfield, QLD 4878, Australia.

³College of Science and Engineering, James Cook University, Cairns, QLD 4870, Australia.

*Corresponding author: stephen.maxwell@my.jcu.edu.au

Maxwell S.J., Watt J.L., Payne M.A. (2026) Asymmetry of eye peduncle length in *Euprotomus iredalei* (Abbott, 1960) (Mollusca: Neostromboidae: Strombidae). *Strombus* 30(1-2): 1-4.

Stromboideans are a complex of gregarious herbivorous marine gastropods found in tropical and subtropical waters globally. This study examined the Australian endemic species, *Euprotomus iredalei* (Abbott, 1960), for which no previous morphological studies have been conducted to date (Fig. 1). This study examined the ommatophores of 39 specimens that were collected in July 2018 by commercial aquarium fishermen supplying Cairns Marine off the Western Australian coast near Karratha (± 20 km). The ommatophores, which in the case of Strombidae, consist of the eye stalk with a cephalic tentacle attached towards the distal end.

Recent studies have indicated that stromboideans are sensitive to the pollutants tributyltin (TBT) and triphenyltin (TPT), and this has been found to cause imposex in affected populations *Laevistrombus turturilla* (Röding, 1798) in Johor Straits, Malaysia (Horiguchi, 2009; Cob et al., 2011). One of the purported indicators of imposex is the asymmetry in eyestalk length found in Philippine *Canarium urceus* (Linné, 1758) (= *Canarium esculentum* Maxwell, Rymer, Congdon & Dekkers, 2020) (Ruaza, 2019). Historical studies into the stromboidean eyestalk have centred on its regeneration following amputation (Gillary, 1972; Hughes, 1976) or the behaviour of the eyes during burrowing (Abbott, 1962). This is the first study to investigate further the length of the right and left ommatophore using *Euprotomus iredalei* from a pristine location (DPIRD Fisheries, 2020).

A sample ($n = 39$, Males = 21, Females = 18) of *Euprotomus iredalei* (Abbott, 1960) adult animals were obtained originating in Karratha, Western Australia, collected by commercial aquarium fishers diving in 10 m on clean sand near a coral reef, and supplied to Carin Marine, Queensland, in late 2018. This area is considered a pristine marine environment (DPIRD Fisheries, 2020). The maturity of this species was confirmed by the researchers, evidenced by a thickened outer lip and a formed anterior siphonal canal: this indicates that the shell had achieved terminal growth. Following collection, animals were preserved in ethanol for 6 months until the time of examination for this study, with shells deposited in the BlueSky Research Foundation Collection (BSRF 5.016). Sex was determined by the presence of a verge in males, and the animals were examined for evidence of imposex, particularly the development of a malformed verge on females (Cob et al., 2011). The eye peduncles were measured at the base where it was attached to the body of the animal (Ruaza, 2019). Each of the peduncles was classified as left and right relative to the position of the proboscis. Peduncles were

straightened with forceps and their length measured using a digital calliper with an accuracy of +/- 0.01 mm. A two-sided paired samples t-test was carried out using SPSS on the data to determine if there was a significant difference within the species between the lengths of the right and left peduncles. Additionally, the effect size of this difference was determined. The proportional sizes for each animal between left and right peduncles were calculated, and a two-sided independent samples T-test was conducted to determine if there was a significant difference in proportional lengths of the peduncles within the population and between males and females.

There was no evidence for imposex, with all animals having normal external sex organs. In all organisms examined, the left peduncle ($7.32 \text{ mm} \pm 0.19$) was significantly shorter ($t_{(38)} = 10.315, p < 0.001$) than the right ($8.86 \text{ mm} \pm 0.23 \text{ SE}$). The effect size was large ($d = 1.05$). There were no differences in the proportional length of peduncles between females and males ($t_{(38)} = 0.056, p = 0.955$).



Figure 1. A sample of the *Euprotomus iredalei* (Abbott, 1960) from Karratha used in this study (photo courtesy of Uwe Weinreich).

The findings of this study showed that all animals examined had two peduncles of differing lengths. In previous studies, this difference was considered an abnormality (Ruaza, 2019), but this study provides contradictory results, indicating it is a natural anatomical trait in the target species. Stromboideans have three distinct shell growth phases: the first twelve to thirty-five days are spent in a pre-metamorphic larval form that is followed by a growth phase where the shell coils and, finally, the flaring lip growth phase, which indicates the commencement of sexual maturation (Berg, 1976; Mitton et al., 1989; Appeldoorn, 2005). The verge in males and gonads only develop once the maturing phase has commenced after the shell has reached terminal growth, defined as the formation of the out lip (Stevely, 1979; Avila-Poveda et al., 2006). Therefore, studies that do not exclude juveniles are prone to false positive imposex results with respect to undertaking comparative physiology, as males have yet to develop their mature verges. This may offer an explanation for this inconsistency, as the body form, such that verges and gonads are not fully developed until the animal has reached sexual maturity.

Stromboideans have evolved to have a shell with two anterior sinuses that enable the eye peduncles to extend past the outer lip while maintaining the aperture face down or allowing the animal to see while partially buried (Savazzi, 1991). The differing peduncle length reflects this, with the right peduncle being longer, as it is more anterior than the left, and the eye stalk has further to reach the more anterior sinus of the outer lip. Therefore, we contend that the differing lengths of eye peduncles have a relationship with the shape and form of the basal sinuses on the outer lip in the species under consideration.

In conclusion, there is widespread sexual dimorphism in the stromboidean complex in shell size, with larger females than males (Abbott, 1949, 1960; Reed, 1993; Maxwell et al., 2017, 2020, 2021a, 2021b, 2022). Furthermore, sexual dimorphism is known in relative body size, some organs and the radula (Reed, 1993, 1995; Mutlu, 2004), and more generally, Physiological morphological and molecular differences observed in non-sexual organs of male and female animals that can affect many aspects of their biological traits (Niksirat et al., 2021). However, this study determined that sexual dimorphism in peduncle length was not present in the target species.

REFERENCES

Abbott D.P. (1962) Observations on the Gastropod *Terebellum terebellum* (Linnaeus) with particular reference to the behavior of the eyes during burrowing. *The Veliger* 5: 1–3.

Abbott R.T. (1949) Sexual dimorphism in Indo-Pacific *Strombus*. *The Nautilus* 63: 58–61.

Abbott R.T. (1960) The genus *Strombus* in the Indo-Pacific. *Indo-Pacific Mollusca* 1: 33–146.

Appeldoorn R.S. (2005) *Development of a combined model of growth in weight for juvenile and adult queen conch (Strombus gigas) and its application to the population off La Pargurea*. NOAA Miami Regional Library, Florida.

Avila-Poveda O.H. & Baqueiro-Cárdenas E.R. (2006) Size at sexual maturity in the queen conch *Strombus gigas* from Columbia. *Boletín de Investigaciones Marinas y Costeras* 35: 223–233.

Berg C.J. (1976) Growth of the queen conch *Strombus gigas*, with a discussion of the practicality of its mariculture. *Marine Biology* 34: 191–199.

Cob Z.C., Arshad A., Bujang J.S., Abd Ghaffar M. (2011) Description and evaluation of imposex in *Strombus canarium* Linnaeus, 1758 (Gastropoda, Strombidae): a potential bio-indicator of tributyltin pollution. *Environmental Monitoring and Assessment* 178: 393–400.

Department of Primary Industries and Region Development Fisheries (DPIRD). (2020) *Rock oyster research moves to the Pilbara*. Western Australian Government, Perth.

Gillary H.L. (1972) The regenerating eye of *Strombus*: anatomy and electrophysiology. *American Zoologist* 12: 691.

Hughes H.P.I. (1976) Structure and regeneration of the eyes of strombid gastropods. *Cell Tissue Research* 171: 259–271.

Horiguchi, T. (2009) Mechanism of imposex induced by organotins in gastropods. In: Arai T., Harino H., Ohji M., Langston W.J. (Eds.) *Ecotoxicology of Antifouling Biocides*. Springer, Tokyo. Pp. 111–123.

Maxwell S.J., Rowell M.K., Hernandez Duran L.C., Rymer T.L. (2020) Population structure of *Canarium labiatum* (Röding, 1798) (Mollusca: Neostrombidae: Strombidae) on Green Island, Great Barrier Reef, Queensland. *Proceedings of the Royal Society of Queensland* 128: 13–20.

Maxwell S.J., Rymer T.L., Congdon B.C. (2017) Sex-ratio bias in *Laevistrombus canarium* Linné, 1758 (Gastropoda: Strombidae) from Far North Queensland, Australia. *Memoirs of the Queensland Museum* 60: 91–96.

Maxwell S.J., Rymer T.L., Congdon B.C. (2021b) A theoretical composite model for population sex-specific shell size dynamics in Strombidae (Gastropoda, Neostrombidae). *Journal of Natural History* 55: 2661–2672.

Maxwell S.J., Rymer T.L., Watt J. (2021) Field notes on sex-bias in *Gibberulus dekkersi* Maxwell, Hernandez Duran, Rowell and Rymer, 2021 (Gastropoda: Neostrombidae: Strombidae) on the Great Barrier Reef. *Pacific Science* 75(4): 525–530.

Maxwell S.J., Todd S.J., Rymer T.L. (2022) Population structure and morphology of *Canarium (Canarium) incisum* and *Canarium (Canarium) esculentum* (Mollusca: Neostrombidae: Strombidae) from the Philippines with preliminary notes on aperture colouration based on DArTseq data. *Acta Zoologica* 104: 1–7.

Mitton J.B., Berg C.J., Orr K.S. (1989) Population structure, larval dispersal, gene flow in the queen conch, *Strombus gigas*, of the Caribbean. *Biological Bulletin* 177: 356–362.

Mutlu E. (2004) Sexual dimorphisms in radula of *Conomurex persicus* (Gastropoda: Strombidae) in the Mediterranean Sea. *Marine Biology* 145: 693–698.

Niksirat H., Silino V., Steinback C., Levander F. (2021) High-resolution proteomic profiling shows sexual dimorphism in zebrafish heart-associated proteins. *Journal of Proteome Research* 20: 4075–4088.

Reed S.E. (1993a) Gonadal comparison of masculinized females and androgynous males to nor-mal males and females in *Strombus* (Mesogastropoda: Strombidae). *Journal of Shellfish Research* 12: 71–75.

Reed S.E. (1993b) Size differences between sexes (including masculinized females) in *Strombus pugilis* (Mesogastropoda: Strombidae). *Journal of Shellfish Research* 12: 77–79.

Reed S.E. (1995) Sexual trimorphism in *Strombus luhuanus* Linné, 1758 (Mollusca: Gastropoda) at Shirahama, Japan. *Journal of Shellfish Research* 14: 159–160.

Ruaza F.C. Jr. (2019) Imposex Incidence, morphological and histological description of gonad in *Canarium urceus urceus* Linnaeus, 1758 (Mollusca: Gastropoda) in Caraga Region, Philippines. *Journal of Aquatic Science* 5: 1–6.

Savazzi E. (1991) Constructional morphology of strombid gastropods. *Lethaia* 24: 311–331.

Stevely J.M. (1979) The biology and fishery of the queen conch, *Strombus gigas*: a review. *Proceedings of the Fourth Annual Tropical and Subtropical Fisheries Technology Congress of the Americas*: 203–210.