RESEARCH ARTICLE



Morphology and phylogenetic analysis of two new deep-sea species of Chrysogorgia (Cnidaria, Octocorallia, Chrysogorgiidae) from Kocebu Guyot (Magellan seamounts) in the Pacific Ocean

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Abstract

Two new species of *Chrysogorgia* Duchassaing & Michelotti, 1864 collected from Kocebu Guyot in the Magellan seamounts of the Pacific Ocean are described and illustrated: *Chrysogorgia ramificans* **sp. nov.** collected from a depth of 1831 m and *Chrysogorgia binata* **sp. nov.** collected from a depth of 1831 m and *Chrysogorgia binata* **sp. nov.** collected from a depth of 1669 m. *Chrysogorgia ramificans* **sp. nov.** belongs to the *Chrysogorgia* "group A, Spiculosae" with rods distributed in body wall and tentacles, and *C. binata* **sp. nov.** belongs to the "group C, Squamosae typicae" with rods and/or spindles not present but only scales. *Chrysogorgia ramificans* **sp. nov.** differs from congeners by its main stem with 2/5R branching sequence at the bottom forming two large bottlebrush-shaped branches with 1/3R branching sequence at the top. *Chrysogorgia binata* **sp. nov.** is similar to *C. scintillans* Bayer & Stefani, 1988, but differs by its larger polyps, larger sclerites in the body wall, and different scales in the upper part of polyps. The mtMutS genetic distances between *C. ramificans* **sp. nov.** and *C. binata* **sp. nov.** and 0.33%–2.28% and 0.33%–2.94%, respectively, while the intraspecific distances are in the range of 0–0.16%. Molecular phylogenetic analysis indicates that *C. ramificans* **sp. nov.** is clustered with *C. chryseis* Bayer & Stefani, 1988, both with high support indicating close relationships.

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Keywords

Anthozoa, Chrysogorgia ramificans sp. nov., Chrysogorgia binata sp. nov., gorgonian, phylogeny, taxonomy

Introduction

Within the gorgonian family Chrysogorgiidae, the genus *Chrysogorgia* Duchassaing & Michelotti, 1864 is the largest and most common group, distributed worldwide including the Antarctic, ranging from 100 m to 3860 m water depth. In some colonies it is characterized by a spiralling main axis that branches sympodially giving off secondary branches that subdivide dichotomously, resulting in a bottle-brush colony shape. In others, the sympodial main axis does not spiral, resulting in fan-like planar or bi-flabellate colonies (Pante and Watling 2011; Cordeiro et al. 2015). To date, *Chrysogorgia* contains 70 species (Cairns 2018). Among them, 45 species are found only from the Pacific, 17 species only from the Atlantic and 7 only from the Indian Ocean (Cairns 2001, 2007, 2018; Pante and Watling 2011; Cordeiro et al. 2015). *Chrysogorgia flexilis* Wright & Studer, 1889 occurs in both the Pacific and Indian Oceans (Wright and Studer 1889; Cairns 2001).

Based on the presence of rods or scales in the body wall and tentacles, Versluys (1902) divided *Chrysogorgia* species into three groups, which were summarized by Cairns (2001) as following: "group A, Spiculosae" (rods and/or spindles in body wall and tentacles) with 38 species, "group B, Squamosae aberrantes" (rods and/or spindles in tentacles but not in body wall) with 13 species, and "group C, Squamosae typicae" (rods and/or spindles not present; only scales) with 18 species. More recently, Cordeiro et al. (2015) described the species *C. upsilonia*, which possesses spindles in the body wall but not in the tentacles, and classified it as "group D, Spiculosae aberrantes". The separation of four groups was further recognized by Cairns (2018).

During the investigation of the Magellan seamount benthic diversity in the tropical Western Pacific, we obtained two golden gorgonians from the Kocebu Guyot using a remotely operated vehicle (ROV). Based on morphological and phylogenetic analyses, both species proved to be new species of *Chrysogorgia* and are described as *C. ramificans* sp. nov. and *C. binata* sp. nov., respectively. Their genetic distances and phylogenetic relationships within *Chrysogorgia* are discussed.

Materials and methods

Specimen collection and morphological examination

Specimens were obtained by the ROV *FaXian* (Discovery) from the Kocebu Guyot in the Magellan seamounts in the tropical Western Pacific during the cruises of the R/V *KeXue* (Science) in 2018 (Fig. 1). The specimens were photographed in situ before sampled, photographed onboard, and then stored in 75% ethanol after collection. Small branches were cut off and stored at -80 °C for molecular study.

The morphological terminology follows Bayer et al. (1983). A stereo dissecting microscope was used to examine the general morphology and anatomy. The sclerites of the polyps



Figure 1. Sampling site in the Kocebu Guyot in the Western Pacific Ocean.

and branches were isolated by digestion of the tissues in sodium hypochlorite, and then were washed with deionized water repeatedly. To investigate the structure of polyps and sclerites, they were air-dried and mounted on carbon double adhesive tape and coated for scanning electron microscopy (SEM) observation. SEM scans were obtained and the optimum magnification was chosen for each kind of sclerites by using TM3030Plus SEM at 5 kV.

The type specimens (registration numbers: MBM286307 and MBM286346) of the two new species have been deposited in the Marine Biological Museum of Chinese Academy of Sciences (MBMCAS) at Institute of Oceanology, Chinese Academy of Sciences, Qingdao, China.

DNA extraction and sequencing

Total genomic DNA was extracted from the polyps of each specimen using the TIANamp Marine Animal DNA Kit (Tiangen Bio. Co., Beijing, China) following the manufacturer's instructions. PCR amplification for the mitochondrial genomic region 5'-end of the DNA mismatch repair protein – *mutS* – homolog (mtMutS) was conducted using primers AnthoCorMSH (5'-AGGAGAATTATTCTAAGTATGG-3'; Herrera et al. 2010) and Mut-3458R (5'-TSGAGCAAAAGCCACTCC-3'; Sánchez et al. 2003). PCR reactions were performed using I-5 2 × High-Fidelity Master Mix DNA polymerase (TsingKe Biotech, Beijing, China). The amplification cycle conditions were as follow: denaturation at 98 °C for 2 min, followed

by 32 cycles of denaturation at 98 °C for 20 s, annealing at 50 °C for 20 s, extension at 72 °C for 15 s, and a final extension step at 72 °C for 2 min. PCR purification and sequencing were performed by TsingKe Biological Technology (TsingKe Biotech, Beijing, China).

Genetic distance and phylogenetic analyses

All the available mtMutS sequences of *Chrysogorgia* spp. and the out-group species from related chrysogorgiidid genera were downloaded from GenBank, and those without associated publications or named *Chrysogorgia* sp. were omitted from the molecular analyses (see Table 2 and Fig. 8). The sequences were aligned using MAFFT v.7 (Katoh and Standley 2013) with the G-INS-i algorithm. Genetic distances, calculated as uncorrected "*p*" distances within each species and among species, were estimated using v.6 (Tamura et al. 2013).

For the phylogenetic analyses, only one sequence was randomly selected from the conspecific sequences without genetic divergence (see Table 2). The evolutionary model GTR+G was the best-fit model for mtMutS, selected by AIC as implemented in jModeltest2 (Darriba et al. 2012). Maximum likelihood (ML) analysis was carried out using PhyML-3.1 (Guindon et al. 2010). For the ML bootstraps, we consider values < 70% as low, 70–94% as moderate and \geq 95% as high following Hillis and Bull (1993). Node support came from a majority-rule consensus tree of 1000 bootstrap replicates.

Bayesian inference (BI) analysis was carried out using MrBayes v3.2.3 (Ronquist and Huelsenbeck 2003) on CIPRES Science Gateway. Posterior probability was estimated using four chains running 10,000,000 generations sampling every 1000 generations. The first 25% of sampled trees were considered burn-in trees. For the Bayesian posterior probabilities, we consider values < 0.95 as low and \ge 0.95 as high following Alfaro et al. (2003). The accession numbers of the mtMutS sequences are listed next to the species names in the phylogenetic tree (Fig. 8).

Results

Class Anthozoa Ehrenberg, 1834 Subclass Octocorallia Haeckel, 1866 Order Alcyonacea Lamouroux, 1812 Suborder Calcaxonia Grasshoff, 1999 Family Chrysogorgiidae Verrill, 1883 Genus *Chrysogorgia* Duchassaing & Michelotti, 1864

Chrysogorgia ramificans sp. nov. http://zoobank.org/DF4284E7-CC5E-4AE7-94C8-4E84366387E9 Figs 2, 3

Holotype. MBM286307, station FX-Dive 174 (17°29.93'N, 153°14.69'E), Kocebu Guyot, depth 1831 m, 8 April 2018. GenBank accession number: MK431863.



Figure 2. External morphology and polyps of *Chrysogorgia ramificans* sp. nov.: **A** The holotype in situ **B** Close-up of branches and polyps in situ **C** The colony after collection **D** A single polyp under light microscope **E** Single polyp under SEM. Scale bars: 20 cm (**C**); 1 mm (**D**, **E**).

Diagnosis. *Chrysogorgia* (tertiary "group A, Spiculosae" – rods or spindles in the tentacles and the body wall) with a short basal stem leading to a bottlebrush-shaped main stem, giving of a single major branch also bottlebrush-shaped. Minor branches subdivided dichotomously, up to fourth order, with the first branch internode 20–30 mm long. Branching sequence 1/3R in two large branches and 2/5R in the basal stem. Polyps 2–4 mm tall with a thin neck. Sclerites of polyp body of large and thick rods and spindles with many warts. Small scales and rods in tentacles with many warts. Scales in coenenchyme elongate with irregular edges and a few warts.

Description. Specimen about 73 cm long with the holdfast not recovered. Main stem forming two large bottlebrush-shaped branches whose axis has a brown metallic luster. The larger branch is 49 cm long and the other 45 cm long. The basal stem about 24 cm long and 4 mm in diameter (Fig. 2C). Branching sequences differing from bottom to top, 2/5R in the basal stem and 1/3R in the two large branches. Branches subdivided dichotomously, up to fourth order, the first branch internodes 20–30 mm long, with the terminal branchlets up to 41 mm. Distance between adjacent branches 8–12 mm, and orthostiche intervals about 30 mm in the two large branches and 50 mm at the bottom. Polyps thin, about 2–4 mm long and 1.0–1.5 mm wide at bases, with a long neck less than 1 mm wide. Two to four polyps on the first internodes, up to eight on terminal branchlets (Fig. 2D). No polyps on main axis internodes.

Rods and spindles of base of polyp body wall large and thick, rarely branched, with many warts on surface, and measuring $247-628 \times 109-180 \mu m$, with an average of $430 \times 136 \mu m$ (Figs 2D, 3C). Rods and spindles longitudinally arranged in the polyp neck extending to the rachis of tentacles, all covered with many warts, and measuring $95-520 \times 25-96 \mu m$, with an average of $304 \times 54 \mu m$ (Fig. 3A). Scales of pinnules small with coarse edges and many warts on surface, and measuring $114-214 \times 29-49 \mu m$, with an average of $146 \times 36 \mu m$ (Fig. 3B). Scales elongated and flat in coenenchyme with dentate edges and a few warts, and measuring $139-553 \times 35-87 \mu m$, with an average of $267 \times 61 \mu m$ (Fig. 3D).

Etymology. The Latin adjective *ramificans* (branching) refers to the ramous structure of the stem.

Distribution. Found only from the Kocebu Guyot with water depth of 1831 m.

Habitat. Colony attached to a rocky substrate with a small, oval-shaped holdfast (Fig. 2A).

Remarks. Chrysogorgia ramificans sp. nov. differs from all known congeners by its main stem, with 2/5R branching sequence, forming two large bottlebrush-shaped branches with 1/3R branching sequence (Cairns 2001, 2018; Pante and Watling 2011). The new species belongs to the Chrysogorgia "group A, Spiculosae", in which it mostly resembles C. monticola Cairns, 2007. However, C. ramificans sp. nov. differs from C. monticola by the much longer interval of adjacent branches (8–12 mm vs. 4–5 mm), longer first internode of branch (20–30 mm vs. 12 mm), larger polyps (2–4 mm vs. 1.1 mm in height), much wider rods (109–180 µm vs. 50–80 µm) with thick ends and warty surface in the body walls (vs. with thin ends and usually smooth surface), and small and unbranched rods at the tentacle base (vs. large and lobed) (Cairns 2007).

Within the group A, *Chrysogorgia ramificans* sp. nov. is also similar to *C. arborescens* Nutting, 1908, *C. tuberculata* Cordeiro et al., 2015 and *C. terasticha* Versluys, 1902. However, the new species differs from *C. arborescens* by its much longer interval of adjacent branches (8–12 mm vs. 3 mm), the higher number of polyps in the distal branchlets (up to 8 vs. 2), and usually regular sclerites (vs. irregular) (Nutting 1908). It differs from *C. tuberculata* by the larger orthostiche intervals (30–50 mm vs. 8–23 mm), rods with numerous coarse warts (vs. spindles with acute and sparse



Figure 3. Sclerites of *Chrysogorgia ramificans* sp. nov. **A** Sclerites of polyp neck extending to the rachis of tentacles **B** Sclerites in the pinnules **C** Sclerites at the body base **D** Sclerites of coenenchyme. Scale bars: 200 μ m (**A**, **C**, **D**); 50 μ m (**B**).

warts), rods present in tentacles (vs. only scales), and scales in coenenchyme with inconspicuous warts (vs. prominent) (Cordeiro et al. 2015). The new species can be easily distinguished from *C. terasticha* by its branching sequence (1/3R at top and 2/5R at bottom vs. 1/4L), larger orthostiche intervals (30–50 mm vs. no more than 8 mm), no nematozooids in coenenchyme (vs. many), larger polyps (2–4 mm vs. no more than 1.6 mm), larger rods with various ends (vs. smaller with rounded ends), and the absence of scales at the polyp base (vs. presence) (Versluys 1902).

Chrysogorgia binata sp. nov.

http://zoobank.org/D9FCB01F-49B7-4BBA-B3F0-D40026DB6519 Figs 4–7; Table 1

Holotype. MBM286346, station FX-Dive 173 (17°28.69'N, 153°09.95'E), Kocebu Guyot, depth 1669 m, 7 April 2018. GenBank accession number: MK431862.

Diagnosis. *Chrysogorgia* ("group C, Squamosae typicae") with a biflabellate colony and a short main stem. Polyps 3–5 mm tall. Scales smooth and thin in the basal part of polyps body with various shape, up to 1 mm long. Scales in the upper part of polyps of various shapes, converged to form an inconspicuous and blunt point at the base of a naked tract below each tentacle. Scales bluntly lancet-shaped, often with numerous coarse granules, longitudinally arranged around both sides of each naked tract. Scales irregular and coarse, usually with lobed edges in the back of tentacles. Scales of coenenchyme slipper-shaped with a medial contraction. Nematozooids absent from coenenchyme.

Description. Specimen with two attached individuals of the crustacean genus *Galathea* Fabricius, 1793 (Fig. 4C). Main stem short with a principal bifurcation, forming two roughly parallel, fans (Fig. 4A, F). Calcareous holdfast small and white, about 7 mm in diameter (Fig. 4F). Major branches of each fan subdivided dichotomously or sympodially. Specimen about 16 cm long and 15 cm wide with a brilliant golden axis, and the stem about 1.5 mm in diameter at base (Fig. 4C). Internodes about 5–9 mm long, each with a single polyp except the terminal twigs, which may sometimes have two polyps. Polyps large and orange in situ, about 3–5 mm tall by 1–2 mm wide, with sclerites forming an inconspicuous blunt point at the base of a naked tract below each tentacle (Figs 4D, 5A). Terminal polyps usually with a long and narrow body (Fig. 4E).

In the basal part of the polyp body, the sclerites comprise transversally arranged, large, smooth scales. They represent a variety of shapes, a few with broad marginal lobes, length by width measuring $216-936 \times 58-283 \mu m$, with an average of 549 \times 166 µm, (Figs 5B, 6A). There are also scales in the upper part of polyps of various shapes, sometimes thick and with a medial contraction, often sharp at one end, broad and lobed at the other, which combine to form an inconspicuous and blunt point at the base of a naked tract below each tentacle; measuring $275-635 \times 77-254 \mu m$, with an average of $451 \times 151 \,\mu\text{m}$ (Figs 5B, 6B). Above these points are irregular, elongate or lancet-shaped scales mostly with coarse granules on surface, that are arranged longitudinally around the sides of each naked tract. The scales measure $337-650 \times 45-85 \ \mu m$ with an average $431 \times 70 \ \mu m$ (Figs 5B, 7A). The scales in the back of tentacles are coarse, of various shapes, mostly flat and lobed, and densely and transversally arranged, measuring $88-352 \times 19-149 \mu m$, with an average of $183 \times 55 \mu m$ (Figs 5A, 7B). The scales in the pinnules are curved at an obtuse angle, and are sometimes thick with a few lobes on their edges; measuring $87-196 \times 19-34 \mu m$, with an average of $152 \times 19-34 \mu m$ $27 \ \mu m$ (Fig. 7D). The scales of the coenenchyme are generally slipper-shaped, some elongate elliptical, occasionally with indentations in their edges, and measure 138-361 \times 40–87 µm, with an average of 222 \times 56 µm (Fig. 7C). The coenenchyme is thin with no nematozooids.



Figure 4. External morphology of the holotype and polyps of *Chrysogorgia binata* sp. nov. **A** Two planar fans of the colony after fixation **B** The holotype (arrow) in situ. Laser dots spaced at 33 cm used for measuring dimensions **C** The colony after collection **D** A single polyp under SEM **E** Single terminal polyp under light microscope **F** Short trunk with the first bifurcation of branches after fixation. Scale bars: 10 cm (**A**); 5 cm (**C**); 1 mm (**D**, **E**); 1 cm (**F**).

Etymology. The Latin adjective *binatus* (binate) refers to the biflabellate structure of the species.

Distribution. Found only from the Kocebu Guyot in the Magellan seamounts with water depth of 1669 m.

Characters/ Species	C. binata sp. nov.	C. chryseis	C. desbonni	C. electra	C. pinnata	C. scintillans	C. stellata	C. upsilonia	
Group type	С	В	А	С	А	С	В	D	
Colony shape	biflabellate	biflabellate	biflabellate	biflabellate	flabellate	biflabellate or multiflabellate	multiflabel- late	flabellate	
Internode length (mm)	5–9	5	3-4	6–12	3–3.5	6–7	8–10	4–30	
Polyp height (mm)	3–5	up to 2	up to 2.8	1.75–2.00	up to 2.8	up to 2.75	2–4	up to 4	
Eight points beneath the tentacles	short and blunt	long and sharp	inconspicuous	inconspicuous	inconspicuous	short and blunt	long and sharp	inconspicu- ous	
Sclerites in body wall	scales various shape with low and broad marginal lobes	scales terete, tapering smoothly toward pointed ends with weak and broad marginal lobes	spindles often curved, somewhat flattened	scales elongate rods elong with narrow with flatter prominent tips marginal lobes		scales various shape with low and broad marginal lobes	scales terete with broad marginal lobes	spindles tuberculate	
Maximum length of scale in body wall (mm)	0.93	0.7	0.75	0.6	0.56	0.65	1.1	0.67	
Sclerites in tentacles	scales	rods and scales	rods and scales	scales	rods	scales	rods and scales	scales	
Maximum length of rods in tentacles (mm)	0.65	0.3	0.24	-	0.21	-	0.5	0.16	
Scale shape in coenenchyme	mainly slipper shape	various shape with prominent marginal lobes	elongate, warty with irregular margins	elongate, tapered with prominent marginal lobes	relatively smooth with finely serrate edges	mainly slipper shape	elongate with more or less marginal lobes	with serrate margins	
Nematozooids on stem and large branches	absent	absent	-	absent	-	absent	conspicuous	_	
Distribution	Western Pacific	Western Pacific	Western Atlantic	Western Pacific	Eastern Pacific	Central and Eastern Pacific	Central Pacific	South Atlantic	
References	Present Bayer and Cairns 2001 study Stefani 1988		Cairns 2001	Bayer and Stefani 1988		Bayer and Stefani 1988, Cairns 2018	Bayer and efani 1988, airns 2018Nutting 1908, Bayer and Stefani 1988		

Table 1. Comparison of Chrysogorgia species with planar structure. "-"means missing data.

Habitat. Colony attached to a rocky substrate with a small holdfast (Fig. 4B).

Remarks. Within the known species of *Chrysogorgia*, seven species mainly possess a planar structure (Table 1). Among these, including our specimen, *C. desbonni* Duchassaing & Michelotti, 1864 and *C. pinnata* Cairns, 2007 belong to the *Chrysogorgia* "group A, Spiculosae"; *Chrysogorgia chryseis* Bayer & Stefani, 1988 and *C. stellata* Nutting, 1908 belong to the "group B, Squamosae aberrantes". *Chrysogorgia binata* sp. nov.; *C. electra* Bayer & Stefani, 1988 and *C. scintillans* Bayer & Stefani, 1988 belong to the "group C,



Figure 5. Polyps of *Chrysogorgia binata* sp. nov. **A** Three polyps under SEM **B** Head and body of one polyp under SEM. Scale bars: 2 mm (**A**); 1 mm (**B**).

Squamosae typicae". The species *C. upsilonia* Cordeiro, Castro & Pérez, 2015 belongs to the "group D, Spiculosae aberrantes". Based on the arrangement of the sclerites, *Chrysogorgia binata* sp. nov. can easily be distinguished from the species in groups A, B and D.

Both *Chrysogorgia binata* sp. nov. and *C. electra* have a biflabellate colony. However, the new species differs from the latter by its larger polyps (3–5 mm vs. generally 1.75–2.00 mm in height), eight short and blunt points beneath the tentacles (vs. inconspicuous), scales of various shapes with low and broad marginal lobes in the body wall (vs. elongate with narrow prominent marginal lobes), scales mainly slipper-shaped in coenenchyme (vs. elongate tapered with prominent marginal lobes) (Bayer and Stefani 1988). *Chrysogorgia binata* sp. nov. differs from *C. scintillans* by its larger polyps (3–5 mm vs. up to 2.8 mm in height), larger sclerites in the body wall (maximum length 0.93 mm vs. 0.65 mm), scales in the upper part of polyps (irregular and usually with sharp end vs. regular and usually with smooth end), scales around the sides of each naked tract (lancet-shaped and usually with coarse granules vs. twisted, flat and often lobed) (Bayer and Stefani 1988, Cairns 2018).



Figure 6. Sclerites of *Chrysogorgia binata* sp. nov. **A** Sclerites in the basal part of the polyp body **B** Sclerites in the point at the base of a naked tract below each tentacle. Scale bar: $300 \mu m$ (all at the same scale).

Genetic distance and phylogenetic analyses

Two mtMutS sequences of the two new species were obtained and deposited in Gen-Bank, and the accession number and the length are as follows: MK431863, 695 bp for *C. ramificans* sp. nov.; and MK431862, 690 bp for *C. binata* sp. nov. The alignment datasets each comprised 649 nucleotide positions. The mtMutS genetic distances among the species of *Chrysogorgia* range from 0.16% to 2.94%, while the intraspecific distances within *C. binata* sp. nov., *C. tricaulis, C. artospira, C. averta, C. abludo* and

	Species/ populations	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1	Chrysogorgia ramifican <i>s</i> sp. nov. MK431863	-													
2	<i>C. binata</i> sp. nov. MK431862	2.28%	_												
3	<i>C.</i> cf. <i>stellata</i> JN2279201	2.12%	0.16%	-											
4	C. tricaulis JN227990, JN227991, JN227998, GQ180123–31, EU268056	0.82%	1.79%	1.63%	0										
5	<i>C. artospira</i> GQ180132-5, GQ353317	0.65%	1.63%	1.47%	0.16%	0									
6	<i>C. artospira</i> GQ868346	0.82%	1.79%	1.63%	0.33%	0.16%	-								
7	<i>C. averta</i> KC788265, GQ180136	0.98%	1.96%	1.79%	0.49%	0.33%	0.49%	0							
8	<i>C. abludo</i> GQ180139, JN227999	1.47%	2.45%	2.28%	0.98%	0.82%	0.98%	1.14%	-						
9	<i>C. abludo</i> GQ180138	1.96%	2.94%	2.77%	1.47%	1.31%	1.47%	1.63%	0.49%	-					
10	C. chryseis DQ297421, JN227992	2.28%	0.49%	0.33%	1.79%	1.63%	1.79%	1.96%	2.45%	2.94%	-				
11	<i>C. pinnata</i> JN227988	0.65%	1.63%	1.47%	0.16%	0.00%	0.16%	0.33%	0.82%	1.31%	1.63%	-			
12	<i>C. monticola</i> JN227989	0.33%	2.28%	2.12%	0.82%	0.65%	0.82%	0.98%	1.47%	1.96%	2.28%	0.65%	-		
13	<i>Radicipes stonei</i> MG986912	2.28%	2.61%	2.45%	1.79%	1.63%	1.79%	1.96%	2.45%	2.94%	2.61%	1.63%	2.28%	-	
14	<i>Radicipes gracilis</i> JN227987	1.79%	2.12%	1.96%	1.31%	1.14%	1.31%	1.47%	1.96%	2.45%	2.12%	1.14%	1.79%	1.14%	-

Table 2. Interspecific and intraspecific uncorrected pairwise distances at mtMutS of species of*Chrysogorgia* and *Radicipes*.

C. chryseis are in the range 0–0.16% (Table 2). Thus, there is no distinct barcoding gap between the intra- and interspecific distances. The genetic distances between the new species *C. ramificans* sp. nov. and the known sequences of the congeners range from 0.33%-2.28%, and those between *C. binata* sp. nov. and the congeners are in the range of 0.33%-2.94% (Table 2).

The ML and BI phylogenetic trees of the mtMutS gene were nearly identical in topology and thus were combined into a consensus tree with both support values (Fig. 8). All the *Chrysogorgia* species were separated into two main groups (Fig. 8). Group I includes *C. binata* sp. nov., *C. cf. stellata* and *C. chryseis*, and Group II contains the subclades *C. ramificans* sp. nov. + *C. monticola*, *C. artospira*, *C. pinnata*, *C. averta*, *C. abludo*, *C. tricaulis* and *C. monticola*.



Figure 7. Sclerites of *Chrysogorgia binata* sp. nov. **A** Sclerites around the sides of each naked tract **B** Sclerites in the back of tentacles **C** Sclerites of the coenenchyme **D** Sclerites in pinnules. Scale bar: 100 μ m (all at the same scale).



Figure 8. Unrooted maximum likelihood (ML) tree inferred from the mtMutS sequences of *Chrysogorgia* and related genera and species. Numbers at the nodes represent ML and Bayesian inference (BI) support values, respectively. Newly sequenced species are in bold.

Discussion

Chrysogorgia ramificans sp. nov. mostly resembles C. monticola Cairns, 2007, which is also strongly supported by the phylogenetic tree and their genetic distance. However, the two species can be easily separated, as discussed above. In the phylogenetic trees, C. binata sp. nov., C. cf. stellata Bayer & Stefani, 1988 and C. chryseis Bayer & Stefani, 1988 formed a single clade with high support, indicating their close relationships (Fig. 8). However, C. binata sp. nov. belongs to the Chrysogorgia "group C, Squamosae typicae", while C. chryseis belongs to "group B, Squamosae aberrantes" (Table 1; Bayer and Stefani 1988). Bayer and Stefani (1988) also reported a specimen they recorded as C. cf. stellata Nutting, 1908, which was based only on some detached branches. Chrysogorgia binata sp. nov. differs from C. cf. stellata by its larger polyps (3–5 mm vs. about 2 mm), many elongate or lancet-shaped scales below the tentacle base (vs. one or two rods with coarse granules), short, squarish or polygonal scales in the polyp body wall (vs. narrow and long), regular slipper-shaped scales in coenenchyme (vs. relatively irregular) (Bayer and Stefani 1988). No sequences are available for Chrysogorgia stellata Nutting, 1908, a species possessing a multiflabellate colony form, while C. binata sp. nov. is biflabellate. The new species differs from C. stellata also in the shorter and more blunt points beneath the tentacles (vs. long and sharp), various shapes of scales in the upper part of the body wall (vs. only a single slenderly elongate shape), and generally slipper-shaped scales in coenenchyme (vs. elongate with more lobed margin).

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References

- Alfaro ME, Zoller S, Lutzoni F (2003) Bayes or Bootstrap? A simulation study comparing the performance of Bayesian Markov chain Monte Carlo sampling and bootstrapping in assessing phylogenetic confidence. Molecular Biology and Evolution 20(2): 255–266. https:// doi.org/10.1093/molbev/msg028
- Bayer FM, Grasshoff M, Verseveldt J (1983) Illustrated Trilingual Glossary of Morphological and Anatomical Terms Applied to Octocorallia. E. J. Brill/Dr. W. Backhuys, Leiden, 75 pp.
- Bayer FM, Stefani J (1988) A new species of *Chrysogorgia* (Octocorallia: Gorgonacea) from New Caledonia, with descriptions of some other species from the western Pacific. Proceedings of the Biological Society of Washington 101(2): 257–279.
- Cairns SD (2001) Studies on western Atlantic Octocorallia (Gorgonacea: Ellisellidae). Part 1: The genus *Chrysogorgia* Duchassaing & Michelotti, 1864. Proceedings of the Biological Society of Washington 114(3): 746–787. https://doi.org/10.2988/0006-324X(2007)120[243:SOWAOG]2.0.CO;2
- Cairns SD (2007) Calcaxonian Octocorals (Cnidaria: Anthozoa) from the Eastern Pacific seamounts. Proceedings of the California Academy of Sciences 58(25): 511–541.
- Cairns SD (2018) Deep-Water Octocorals (Cnidaria, Anthozoa) from the Galápagos and Cocos Islands. Part 1: Suborder Calcaxonia. ZooKeys 729: 1–46. https://doi.org/10.3897/ zookeys.729.21779
- Cordeiro RTS, Castro CB, Pérez CD (2015) Deep-water octocorals (Cnidaria: Octocorallia) from Brazil: family Chrysogorgiidae Verrill, 1883. Zootaxa 4058(1): 81–100. https://doi. org/10.11646/zootaxa.4058.1.4
- Darriba D, Taboada GL, Doallo R, Posada D. (2012) jModelTest 2: more models, new heuristics and parallel computing. Nature Methods 9: 772. https://doi.org/10.1038/nmeth.2109
- Duchassaing P, Michelotti J (1864) Supplément au mémoire sur les coralliaires des Antilles. Memorie della Reale Accademia delle Scienze di Torino 2(23): 97–206. https://doi. org/10.5962/bhl.title.105196
- Ehrenberg CG (1834) Beiträge zur physiologischen Kenntniss der Corallenthiere im allgemeinen, und besonders des rothen Meeres, nebst einem Versuche zur physiologischen Systematik derselben. Abhandlungen der Königlichen Akademie der Wissenschaften, Berlin 1: 225–380.
- Grasshoff M (1999) The shallow-water gorgonians of New Caledonia and adjacent islands (Coelenterata, Octocorallia). Senckenbergiana biologica 78: 1–121.

- Guindon S, Dufayard JF, Lefort V, Anisimova M, Hordijk W, Gascuel O (2010) New algorithms and methods to estimate maximum-likelihood phylogenies: assessing the performance of PhyML 3.0. Systematic Biology 59: 307–321. https://doi.org/10.1093/sysbio/syq010
- Haeckel E (1866) Generelle morphologie der Organismen, vol. 2. Verlag von Georg Reimer, Berlin. https://doi.org/10.1515/9783110848281
- Herrera S, Baco A, Sánchez JA (2010) Molecular systematics of the bubblegum coral genera (Paragorgiidae, Octocorallia) and description of a new deep-sea species. Molecular Phylogenetics and Evolution 55(1): 123–135. https://doi.org/10.1016/j.ympev.2009.12.007
- Hillis DM, Bull JJ (1993) An empirical test of bootstrapping as a method for assessing confidence in phylogenetic analysis. Systematic Biology 42(2): 182–192. https://doi.org/10.1093/sysbio/42.2.182
- Katoh K, Standley DM (2013) MAFFT Multiple Sequence Alignment Software version 7: improvements in performance and usability. Molecular Biology and Evolution 30: 772–780. https://doi.org/10.1093/molbev/mst010
- Lamouroux JVF (1812) Extrait d'un mémoire sur la classification des polypiers coralligènes non entièrement piérreux. Nouveau Bulletin des Sciences, Société Philomathique de Paris 3(63): 181–188.
- Nutting CC (1908) Descriptions of the Alcyonaria collected by the U.S. Bureau of Fisheries steamer Albatross in the vicinity of the Hawaiian Islands in 1902. Proceedings of the United States National Museum 34: 543–601. https://doi.org/10.5479/si.00963801.341624.543
- Pante E, Watling L (2011) Chrysogorgia from the New England and Corner Seamounts: Atlantic-Pacific connections. Journal of the Marine Biological Association of the United Kingdom 92(5): 911–927. http://doi.org/10.1017/S0025315411001354
- Ronquist FR, Huelsenbeck JP (2003) Mrbayes 3: Bayesian phylogenetic inference under mixed models. Bioinformatics 19: 1572–1574. https://doi.org/10.1093/bioinformatics/btg180
- Sánchez JA, Lasker HR, Taylor DJ (2003) Phylogenetic analyses among octocorals (Cnidaria): mitochondrial and nuclear DNA sequences (lsu-rRNA, 16S and ssu-rRNA, 18S) support two convergent clades of branching gorgonians. Molecular Biology and Evolution 29: 31–42. https://doi.org/10.1016/S1055-7903(03)00090-3
- Tamura K, Stecher G, Peterson D, Filipski A, Kumar S (2013) MEGA6: Molecular Evolutionary Genetics Analysis Version 6.0. Molecular Biology and Evolution 30(12): 2725–2729. https://doi.org/10.1093/molbev/mst197
- Verrill AE (1883) Report on the Anthozoa, and on some additional species dredged by the "Blake" in 1877–1879, and by the U.S. Fish Commission steamer "Fish 126 Hawk" in 1880–82. Bulletin of the Museum of Comparative Zoology 11: 1–72.
- Versluys J (1902) Die Gorgoniden der Siboga-Expedition. 1. Die Chrysogorgiiden. Siboga Expeditie 13: 1–120.
- Wright EP, Studer T (1889) Report on the Alcyonaria collected by H.M.S. Challenger during the years 1873–76. Report on the Scientific Results of the Voyage of H.M.S. Challenger during the years 1873–76, Zoology 31(64): 1–314.