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# Size Structure of the Macrofuna Community in La Azufrada Coral Reef (Isla Gorgona, Colombian Pacific) <sup>+</sup>

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Abstract: Fauna associated with coral reefs is mainly comprised of macroinvertebrates. In order to 13 predict the impact that disturbances such as coral reef degradation could have on these macrofauna 14 communities, it is essential to characterize their community attributes, and particularly their size 15 distributions. In this research, biomass, and community structure of macrofauna assemblages asso-16 ciated with La Azufrada coral reef (Colombian Pacific), were compared between live and inert cor-17 alline substrates, and among size groups of the same coralline substrate. Transitions from live to 18 inert coralline substrates could lead to macrofaunal assemblages with remarkably lower organic 19 matter contributions, and higher proportions of smaller crustacean macroinvertebrates (< 1mm). 20

**Keywords:** macrofauna; coralline substrates; community attributes; size groups; size distribution; 21 Colombian Pacific 22

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# 1. Introduction

Coral reefs are considered one of the most diverse ecosystems in the world, hosting 25 a similar number of species and habitat complexity than tropical forests [1]. Coral reef 26 services rely largely on their associated macrofauna because they represent the highest 27 abundance and diversity, as well as play a key role in the food webs of reefs by linking 28 primary producers to secondary and tertiary consumers [1-3]; contributing to organic 29 matter and nutrient recycling [4]; or promoting the erosion of the coralline substrate [4]. 30 However, due to their cryptic behavior, small size, and polemic taxonomic classifications, 31 research of macrofauna communities associated with coral formations are still very scarce, 32 particularly when compared with research targeting fishes and larger animals [5–7]. In 33 the Eastern Tropical Pacific (ETP), early studies of macrofauna communities associated 34 with coral reefs were conducted during the seventies in Panama and Colombia, focusing 35 mainly on the description of species of Decapoda intimately associated with the branching 36 coral of the genera *Pocillopora* Lamarck, 1816 [8]. 37

Awareness regarding the degradation of coral reefs has risen, as it is a worldwide38and growing phenomenon [9]. In the ETP, coral degradation is linked with suboptimal39climatic and oceanographic conditions, particularly with the warming events during El40Niño Southern Oscillation (ENSO) [10], which are also intensified due to climate change41[11]. Coral mortality is followed by changes in the dominance of the type of coralline sub-42strate from live corals to dead coral fragments covered mainly by turf algae [10,12]. Hence,43

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understanding the impact that coralline substrate transformation has on these macrofauna assemblages is essential.

Another attribute that is important to evaluate is body size, considering that it affects 3 basic ecological processes such as metabolic rates, generation times, or predator-prey in-4 teractions [13]. Therefore, changes in the community size spectrum can affect the direction 5 and efficiency of energy and organic matter flows through the food webs [13,14]. In order 6 to make predictions of the impacts linked to size distribution changes, it is crucial to un-7 derstand how attributes, such as density, biomass, or taxonomic composition are allocated 8 among size groups. However, only recently, Fraser et al. (2021a,b) characterized the size 9 spectrum of these community attributes in macrofauna assemblages associated with dif-10 ferent coralline substrates in the Great Barrier Reef of Australia [15,16]. 11

This research documents differences in macrofauna assemblages between live and 12 inert coralline substrates in a Colombian Pacific reef, through the evaluation of two community attributes, biomass and community structure. More notable, this research highlights how these attributes are distributed among size groups, and how each size group 15 contributes to the overall biomass and community structure in each coralline substrate. 16

### 2. Materials and Methods

## 2.1. Study Area

Gorgona is a continental island located approximately 35 km off the coast (Figure 19 1A). The area around Isla Gorgona is characterized by high precipitations (mean annual 20 >6600 mm), high sea surface temperatures (26–29 °C), high turbidity (transparency < 6–12 21 m), salinities ranging 25–33, and a semidiurnal tidal regime with high fluctuations (>5 m) 22 [17].

Despite the suboptimal conditions for corals to grow, coral formations with different 24 degrees of development can be found in Gorgona [10] (Figure 1B). La Azufrada is indeed 25 one of the most developed coral reefs in the Colombian Pacific [18] (Figure 1B,C). Coral 26 framework in La Azufrada is dominated by *Pocillopora* branching corals [10,18]. 27



**Figure 1.** Map of Isla Gorgona and La Azufrada coral reef in the Colombian Pacific: (**A**) Map of the Colombian Pacific coast indicating Isla Gorgona (black square). Scale: 100 km; (**B**) Map of Isla Gorgona showing the distribution of coral formations around the island according to their type. Legend: Patchy coral communities (green); Incipient reefs (yellow); Fringing reefs (red) [18]-modified. Scale: 1 km; (**C**) Map of La Azufrada coral reef showing the 20 sampling points. Scale: 100 m.

#### 2.2. Sampling

To characterize the macrofauna community inhabiting the coralline substrate in La 35 Azufrada, 40 samples were collected at 20 points randomly distributed along the reef (Figure 1C). Two samples were collected at each point within a 1 m<sup>2</sup> area: (1) live coralline 37 substrate, defined as entire Pocillopora spp. colonies, and (2) inert coralline substrate, defined as the coral skeleton without live tissue. Live colonies were firstly covered with a 39 zip lock bag, detached from the floor, and then sealed underwater. The inert coralline 40

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substrate was collected in 3.5 L containers in a 30 cm quadrat. In the laboratory, each sample unit was defaunated, poured through a 0.5 mm sieve, and preserved in 96% ethanol.

#### 2.3. Laboratory Processing

Samples were filtered through a sieving tower composed of five sieves. Hence, ac-4 cording to their size, macroinvertebrates were retained within the different meshes, and 5 sorted into five size groups: 0.5–1 mm, 1–2 mm, 2–4 mm, 4–8 mm, >8 mm. Wet weight 6 biomass of each size group was measured in an analytical balance (0.001 g resolution) 7 after removing the excess water from invertebrates by lightly blotting the sample in a pa-8 per towel. Afterwards, macroinvertebrates of each size group were counted, classified at 9 high taxonomic levels, and preserved in 96% ethanol. 10

#### 2.4. Data Analysis

All data were firstly standardized to 100 cm<sup>2</sup>. Biomass differences between coralline 12 substrates were tested using a paired t-test. Furthermore, an ANOVA of the Linear Mixed-Effect Model was run to test statistical differences between the size groups within each 14 coralline substrate. Biomass data were previously transformed to meet data symmetry 15 (ln+1). Afterwards, the Tukey's Honestly-Significant Difference post-hoc test was per-16 formed to evaluate pairwise differences between size groups. 17

Densities per taxonomic group were used to test differences in macrofauna commu-18 nity composition between coralline substrates using an ANOSIM test. Hence, a dissimi-19 larity matrix was previously built using the Manhattan distance. Afterwards, the SIMPER 20 analysis was used to evaluate the contribution of each taxon to these dissimilarities. 21

# 3. Results

### 3.1. Biomass

Wet weight biomass of the macrofauna assemblage was significantly different be-24 tween coralline substrates (p < 2.058e-09) (Figure 2A). Significant differences in biomass 25 were also found between size groups in both, live and inert coralline substrates (p < p26 0.0001). All pairwise comparisons were statistically significant (p < 0.05), except for the 27 size groups 0.5–1 mm and 1–2 mm in inert coralline substrates. Moreover, in both coralline 28 substrates, biomass values increased as the size of the organisms increased, and biomass 29 in the size group >8 mm were largely responsible for total biomass (Figure 2B,C and Table 30 1). The dominance of the largest size group was, however, considerably more outstanding 31 in live than in inert coralline substrates (Figure 2B,C and Table 1). 32



Figure 2. Boxplots of wet weight biomass (g/100cm<sup>2</sup>) of macrofauna showing the median (horizontal 33 bar), the interquartile range (boxes), and the outliers (black points): (A) Biomass between live and 34 inert coralline substrates; (B) Biomass among size groups in live coralline substrates. The 0.5–1 mm 35 group was not included in the analysis because none of the obtained measurements was >0.004 g 36 37 (error of the analytical balance); (C) Biomass among size groups in inert coralline substrates.

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(0.5-1  mm, 1-2  mm, 2-4  mm, 4-8  mm, >8  mm). Values are standardized to 100 cm <sup>2</sup> .						
Substrate	0.5–1 mm	1–2 mm	2–4 mm	4–8 mm	>8 mm	Total
Live	-	0.028 (0.009)	0.051 (0.015)	0.209 (0.067)	3.877 (0.753)	4.166 (0.772)
Inert	0.008 (0.007)	0.010 (0.007)	0.032 (0.013)	0.075 (0.028)	0.205 (0.084)	0.330 (0.109)

Table 1. Average biomass (95% IC) according to coralline substrate (live and inert) and size groups ) Walu 1 0 2 1 -1 0 -~~ ~~ a ana atan dandir . 1 . . 100

## 3.2. Community Structure

Twenty-six taxonomic groups were found considering both coralline substrates, with 4 19 taxa occurring in live and 25 taxa in inert substrates. Relative densities per taxonomic 5 group revealed a single taxon dominance in live (Caridea: 49%) and in inert coralline sub-6 strates (Tanaidacea: 35%) (Figure 3A). The difference in macrofauna assemblage composition between substrates was statistically significant (p = 1e-04), accounting for  $\approx 78\%$ . 8 Caridea contributed the most to this dissimilarity (0.28, average dissimilarity contribu-9 tion), followed by Tanaidacea (0.11), and Polychaeta (0.09). Relative densities of these taxa 10 varied greatly between live and inert coralline substrates, Caridea (49% vs. 0.3%, respec-11 tively), Tanaidacea (5% vs. 35%), and Polychaeta (7% vs. 19%) (Figure 3A). 12



Figure 3. Taxonomic composition of macrofauna assemblages: (A) Bar graph comparing the relative taxonomic densities between live and inert macrofauna communities. \* Taxa grouped as Other com-15 prise Sipuncula, Arachnida, Mysida, Holothuroidea, Heterobranchia, and Platyhelminthes; (B) Rel-16 ative taxonomic densities among size groups in live coralline substrates; (C) Relative taxonomic 17 densities among size groups in inert coralline substrates. . 18

In live coralline substrates, dominance by a single taxon was observed in all but the 19 0.5–1 mm group (Figure 3B). This dominance was particularly pronounced in the 1–2 mm 20 and 2–4 mm groups (Caridea densities >70%) (Figure 3B). In the >8 mm group, inverte-21 brates belonging to the Caridea and Anomura-Brachyura taxa were mainly shrimps of the 22 genus Alpheus Fabricius, 1798 and crabs of the genus Trapezia Latreille, 1828, respectively. 23

In inert coralline substrates, a single taxon dominance pattern was observed in all 24 size groups, which was more conspicuous in the 4–8 mm and >8 mm groups, where den-25 sities were dominated by Ophiuroidea (Figure 3C). Dominance of Ophiuroidea decreased 26 in the smallest size fractions and it was progressively replaced by Polychaeta. The 0.5-1 27 mm size group was dominated by Tanaidacea (Figure 3C). 28

#### 4. Discussion

# 4.1. Biomass

Macroinvertebrate biomass showed the most conspicuous difference between coral-31 line substrates, with live coral having >12 times biomass than inert coralline substrates 32 (Figure 2A and Table 1). Higher biomass values in live than inert coralline substrates have 33 also been reported in the Western Micronesia [7] and the ETP [4]. 34

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Biomass distribution in live coralline substrates is remarkably outstanding because 1 total biomass is not only mainly contained in the >8 mm group, but also in very specific 2 taxa, the genera *Trapezia* and *Alpheus*. Several authors have described the intimate association between these two Decapoda and *Pocillopora* colonies in different regions [8,19,20], 4 including Isla Gorgona [21,22]. Therefore, total biomass in live coralline substrates is 5 mainly comprised of decapods that are obligate symbionts of *Pocillopora* colonies. 6

## 4.2. Community Structure

Macrofauna community composition is different between coralline substrates (Fig-8 ure 3A). When compared to other studies, the consistency of our result is not surprising, 9 considering that habitat has been widely identified as a key factor structuring the compo-10 sition of benthic communities [4,5,23,24], outweighing even latitude [6,15]. In our study, 11 this difference is due to taxa present in both substrates, but strongly correlated to only one 12 of them. Particularly, Caridea is strongly correlated to live coralline substrates and Tana-13 idacea to inert coralline substrates (Figure 3A). Hence, both macrofauna assemblages are 14 dominated by crustaceans. The importance of crustaceans in macrofauna communities 15 associated with different coralline microhabitats has been highlighted by several authors 16 [15,24,25]. 17

In live coralline substrate, crustaceans are mainly comprised by Decapoda (Figure 18 3B). The dominance of Decapoda in macrofauna assemblages associated with live coralline substrates, and particularly with branching corals, is consistent across the literature 20 due to their symbiotic associations [8,19,20,25]. Highest contribution of Decapoda to total 21 macrofauna density is maintained in all size groups, except the 0.5–1 mm group, which is 22 dominated by Peracarida (Figure 3B). The dominance of crustaceans according to size 23 groups in live coralline substrate has also been described by Fraser et al. (2021a)[15]. 24

However, when the distribution of crustaceans is studied among size groups in inert 25 coralline substrates, dominance of this taxonomic group is only observed in the 0.5–1 mm 26 group (Peracarida) (Figure 3C). The remaining size groups are dominated by Polychaeta 27 and by Ophiuroidea (Figure 3C). The occurrence of both taxa in macrofauna communities 28 associated with inert coralline substrates has been highlighted by different authors 29 [5,6,26], including in the ETP of Panama [27]. Furthermore, ophiuroids and polychaetes 30 have also been identified as key components in structuring these macrofauna assem-31 blages, due to their role as bioeroders [26]. 32

# 5. Conclusions

This study suggests that a coral degradation event in La Azufrada reef, induced for 34 example by thermal stress associated with an El Niño event could have a significant im-35 pact on this reef system. On the one hand, the composition of the macrofauna assemblage 36 could significantly change towards the dominance of smaller crustaceans (< 1 mm), likely 37 affecting the trophic relationships. On the other hand, and more outstanding, the amount 38 of energy and organic matter available for higher trophic levels could drastically decrease 39 following the death of obligate symbionts like the decapods associated with Pocillopora 40 colonies. These Decapoda are responsible for almost the total biomass contained in live 41 coralline substrates. Hence invertivore populations, such as coral-dwelling species, might 42 decline due to a decrease in their food resources. 43

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**Data Availability Statement:** The data presented in this study are available on request from the corresponding author.

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### References

- Reaka-Kudla, M.L. The global biodiversity of coral reefs: A comparison with rainforests. In *Biodiversity II: Understanding and Protecting Our Biological Resources*; Reaka-Kudla, M.L., Wilson, D.E., Wilson, E.O., Eds.; Joseph Henry Press: Washington, DC, USA, 1997; Volume 2, pp. 83–108.
- Taylor, R.B. Density, Biomass and Productivity of Animals in Four Subtidal Rocky Reef Habitats: The Importance of Small Mobile Invertebrates. *Mar. Ecol. Prog. Ser.* 1998, 172, 37–51. https://doi.org/10.3354/meps172037.
- 3. Kramer, M.J.; Bellwood, O.; Fulton, C.J.; Bellwood, D.R. Refining the Invertivore: Diversity and Specialisation in Fish Predation on Coral Reef Crustaceans. *Mar. Biol.* **2015**, *162*, 1779–1786. https://doi.org/10.1007/s00227-015-2710-0.
- Enochs, I.C.; Hockensmith, G. Effects of Coral Mortality on the Community Composition of Cryptic Metazoans Associated with Pocillopora Damicornis. In Proceedings of the 11th International Coral Reef Symposium, Fort Lauderdale, FL, United States, 7– 11 July 2008.
- 5. Ruiz-Abierno, A.; Armenteros, M. Coral Reef Habitats Strongly Influence the Diversity of Macro- and Meiobenthos in the Caribbean. *Mar. Biodivers.* **2017**, 47, 101–111. https://doi.org/10.1007/s12526-016-0553-7.
- 6. Fraser, K.M.; Stuart-Smith, R.D.; Ling, S.D.; Heather, F.J.; Edgar, G.J. Taxonomic Composition of Mobile Epifaunal Invertebrate Assemblages on Diverse Benthic Microhabitats from Temperate to Tropical Reefs. *Mar. Ecol. Prog. Ser.* **2020**, *640*, 31–43.
- 7. Wolfe, K.; Desbiens, A.; Stella, J.; Mumby, P.J. Length–Weight Relationships to Quantify Biomass for Motile Coral Reef Cryptofauna. *Coral Reefs* **2020**, *39*, 1649–1660. https://doi.org/10.1007/s00338-020-01993-9.
- 8. Alvarado, J.J.; Vargas-Castillo, R. Invertebrados Asociados Al Coral Constructor de Arrecifes *Pocillopora Damicornis* En Playa Blanca, Bahía, Culebra, Costa Rica. *Rev. Biol. Trop.* **2012**, *60*, 77–92.
- 9. Glynn, P.W. Coral Reef Bleaching: Ecological Perspectives. Coral Reefs 1993, 12, 1–17. https://doi.org/10.1007/BF00303779.
- 10. Zapata, F.A.; Vargas-Ángel, B. Corals and coral reefs of the Pacific coast of Colombia. In *Latin American Coral Reefs*; Cortés, J., Ed.; Elsevier Science B.V.: Heredia, Costa Rica, 2003; pp. 419–447.
- Cai, W.; Borlace, S.; Lengaigne, M.; Van Rensch, P.; Collins, M.; Vecchi, G.; Timmermann, A.; Santoso, A.; Mcphaden, M.J.; Wu, L.; et al. Increasing Frequency of Extreme El Niño Events Due to Greenhouse Warming. *Nat. Clim. Chang.* 2014, *4*, 111–116. https://doi.org/10.1038/nclimate2100.
- 12. Diaz-Pulido, G.; McCook, L.J.; Dove, S.; Berkelmans, R.; Roff, G.; Kline, D.I.; Weeks, S.; Evans, R.D.; Williamson, D.H.; Hoegh-Guldberg, O. Doom and Boom on a Resilient Reef: Climate Change, Algal Overgrowth and Coral Recovery. *PLoS ONE* **2009**, *4*, e5239. https://doi.org/10.1371/journal.pone.0005239.
- 13. Mazurkiewicz, M.; Górska, B.; Renaud, P.E.; Włodarska-Kowalczuk, M. Latitudinal Consistency of Biomass Size Spectra-Benthic Resilience despite Environmental, Taxonomic and Functional Trait Variability. *Sci. Rep.* **2020**, *10*, 4164.
- 14. Norkko, A.; Villnäs, A.; Norkko, J.; Valanko, S.; Pilditch, C. Size Matters: Implications of the Loss of Large Individuals for Ecosystem Function. *Sci. Rep.* **2013**, *3*, 2646. https://doi.org/10.1038/srep02646.
- 15. Fraser, K.M.; Stuart-Smith, R.D.; Ling, S.D.; Edgar, G.J. Small Invertebrate Consumers Produce Consistent Size Spectra across Reef Habitats and Climatic Zones. *Oikos* **2021**, *130*, 156–170. https://doi.org/10.1111/oik.07652.
- 16. Fraser, K.M.; Stuart-Smith, R.D.; Ling, S.D.; Edgar, G.J. High Biomass and Productivity of Epifaunal Invertebrates Living amongst Dead Coral. *Mar. Biol.* **2021**, *168*, 1–12. https://doi.org/10.1007/s00227-021-03911-1.
- 17. Díaz, J.M.; Pinzón, J.H.; Perdomo, A.M.; Barrios, L.M.; López-Victoria, M. Generalidades. In *Gorgona Marina: Contribución al Conocimiento de Una Isla Única*; Barrios, L.M., Ed.; INVEMAR: Santa Marta, Colombia, 2001; Volume 1, pp. 17–26.
- Glynn, P.W.; Alvarado, J.J.; Banks, S.; Cortés, J.; Feingold, J.S.; Jiménez, C.; Maragos, J.E.; Martínez, P.; Maté, J.L.; Moanga, D.A.; et al. Eastern Pacific Coral Reef Provinces, Coral Community Structure and Composition: An Overview. In *Coral Reefs of the Eastern Tropical Pacific*; Glynn, P.W., Manzello, D.P., Enochs, I.C., Eds.; Springer: Dordrecht, The Netherlands, 2017; Volume 8, pp. 107–176.
- Abele, L.G.; Patton, W.K. The Size of Coral Heads and the Community Biology of Associated Decapod Crustaceans. J. Biogeogr. 1976, 3, 35–47.
- 20. Stella, J.S.; Pratchett, M.S.; Hutchings, P.A.; Jones, G.P. Coral-associated invertebrates: Diversity, ecology importance and vulnerability to disturbance. *Oceanogr. Mar. Biol.* **2011**, *49*, 43–104.
- 21. Von Prahl, H.; Guhl, F.; Grógl, M. Crustáceos decápodos comensales del coral *Pocillopora damicornis* en la Isla de Gorgona, Colombia. *Bol. Investig. Mar. Costeras* **1978**, *10*.

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- Castro, P. Notes on symbiotic decapod crustaceans from Gorgona Island, Colombia, with a revision of the eastern Pacific species 22. 1 of Trapezia (Brachyura, Xanthidae), symbionts of scleractinian corals. Bol. Investig. Mar. Costeras 1982, 12. 2
- 23. Riddle, M. Patterns in the Distribution of Macrofaunal Communities in Coral Reef Sediments on the Central Great Barrier Reef. Mar. Ecol. Prog. Ser. 1988, 47, 281–292. https://doi.org/10.3354/meps047281.
- Kramer, M.J.; Bellwood, D.R.; Bellwood, O. Benthic Crustacea on Coral Reefs: A Quantitative Survey. Mar. Ecol. Prog. Ser. 2014, 24. 5 511, 105-116. https://doi.org/10.3354/meps10953.
- 25. Plaisance, L.; Knowlton, N.; Paulay, G.; Meyer, C. Reef-Associated Crustacean Fauna: Biodiversity Estimates Using Semi-Quan-7 titative Sampling and DNA Barcoding. Coral Reefs 2009, 28, 977–986. https://doi.org/10.1007/s00338-009-0543-3. 8
- 26. Glynn, P.W.; Enochs, I.C. Invertebrates and their roles in coral reef ecosystems. In Coral Reefs: An Ecosystem in Transition; Du-9 binsky, Z., Stambler, N., Eds.; Springer: Dordrecht, The Netherlands, 2011; pp. 273-325. 10
- 27. Clark, A.M. Echinoderms of coral reefs. In Biology and Geology of Coral Reefs; Jones, O.A., Endean, R., Eds.; Academic Press: New 11 York, NY, USA, 1976; Volume 3, pp. 95-123. 12

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