

# Sexual dimorphism of sharks from the amazonian equatorial coast

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## Abstract

Sexual dimorphism is a widespread feature in several groups of vertebrates. Chondrichthyans differ sexually due to the presence of the clasper, a structure for internal fertilization, and other sexual differences in secondary characteristics. Nevertheless, studies assessing these variations are fairly rare. The main goal of this study is to identify differences in sexual dimorphism in three species of sharks from the Carcharhinidae family (*Rhizoprionodon porosus*, *Carcharhinus porosus* and *Isogomphodon oxyrinchus*) using morphometric tools. A total of 213 specimens were captured in the Amazonian Equatorial Coast and analyzed using 65 morphometric characters. Discriminant analysis and The Student's t-test were used to demonstrate the morphological differences among sexes. Sexual dimorphism was reported at different levels for the three species. This study suggests that the most likely explanation for the presence of these variations is related to their reproductive characteristics and mating behavior.

**Keywords:** sexual dimorphism; sharks; ecological features; Carcharhinidae

**Edited by** Alberto Acosta & Juan Carlos Salcedo-Reyes✉

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**Received:** 16-09-2014 **Accepted:** 29-05-2015

**Published on line:** 18-08-2015

**Citation:** Barbosa Martins AP, da Silva Filho E, Manir Feitosa L, Nunes e Silva LP, da Silva de Almeida Z, Silva Nunes JL (2015) Sexual dimorphism of sharks from the amazonian equatorial coast. *Universitas Scientiarum* 20(3): 297-304 doi: 10.11144/Javeriana.SC20-3.sdos

**Funding:** REVIZEE, CNPq and BASA.

**Electronic supplementary material:** N/A

## Introduction

Sexual dimorphism is common among vertebrates (Kajiura et al. 2005). With regard to fishes, in mature specimens the internal differences between genders are usually clear, but externally they are not easily differentiated (Moyle & Cech 2004). Conversely, all cartilaginous fishes exhibit external sexual dimorphism brought about by the modification of males' pelvic fins. This results in a specialized structure for internal fertilization, called clasper. Beyond this evident characteristic, many other species also exhibit sexual dimorphism through other morphological features (Orlov et al. 2010, Orlov & Cotton 2011).

Males and females of the same species may respond differently to selective pressures (Filiz & Taşkavak 2006). In the case of elasmobranchs, the differences between males and females include variations in their external reproductive anatomy,



body size (Compagno 1984, Carlson & Parsons 1997), dental formula (Kajiura et al. 1996), number and distribution of Ampullae of Lorenzini (Crooks & Waring 2013a), and dermal thickness (Kajiura et al. 2000, Crooks & Waring 2013b). These morphological differences can impact ecological aspects such as foraging, competition, and habitat use (Ellis & Shackley 1995, Piorski & Nunes 2001). Also, males and females of the same elasmobranch species show different behaviors during mating (Gointein et al. 1998). Nonetheless, these variations in intra-specific secondary sexual characteristics have been poorly studied worldwide (Kajiura 2000, Filiz & Taşkavak 2006). The Amazonian Equatorial Coast, one of the elasmobranchs conservation hotspots of the world (Dulvy et al. 2014), is no exception.

It is known that *Rhizoprionodon porosus* and *Carcharhinus porosus* are the most abundant elasmobranch catches in Maranhão state (Almeida et al. 2011) and that *Isogomphodon oxyrinchus* is an endemic species of the coastal waters off northern South America. These species are listed as critically endangered by the Brazilian government (ICMBIO 2012) but are still captured in some of these areas (Lessa et al. 2006, Rodrigues-Filho et al. 2012). Despite their importance for the local ecosystem and the high fishing economic value in the state, information on the biology and ecology of the shark species of this region is not widely available.

Since updated data about these species is scarce and consistent knowledge is required for developing management strategies for local species, this study aims to evaluate the presence of sexual dimorphism within three Carcharhinidae sharks captured off the coast of Maranhão state, Amazonian Equatorial Coast, Brazil.

## Materials and methods

**Area of study:** The coastline of Maranhão is the second most extensive in Brazil, with a length of 640 km. The west side of the coastline belongs to the Amazonian Equatorial Coast and is characterized by the presence of recesses with excessively drained river valleys formed by the influence of fluvial estuarine bays (Muehe 2006). As a result, the region has high turbid waters, exuberant mangroves, and rich biotic communities (Souza-Filho 2005) (Figure 1).

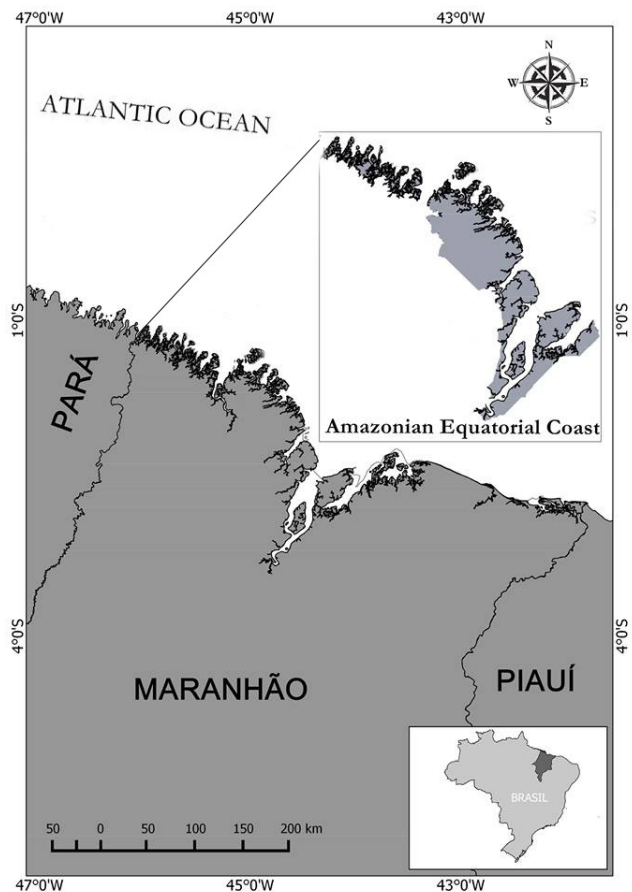


Fig. 1. Map of the coast of Maranhão State, Brazil.

**Sampling, morphometric measurements and data analysis:** The specimens were caught between October 1997 and March 1999 in shallow waters along the Amazonian Equatorial Coast of Maranhão using gillnets with a mesh size of 95-100mm. The sharks were stored in thermal coolers with ice and taken to the Ichthyology Laboratory of the *Universidade Federal do Maranhão* (Federal University of Maranhão) for morphometric analysis. The taxonomic identification was carried out using specialized literature (Compagno 1984).

Using a caliper and metric tape and applying the Piorski et al. (2010) protocol adapted from Compagno (1984), 65 morphometric characteristics were measured in various sized specimens of three shark species: *Carcharhinus porosus* (50 females with size range: 41.1 – 89.6 cm and 20 males with size range: 42.3 – 91.1 cm), *Rhizoprionodon porosus* (11 females with

size range: 47.6 – 83.5 cm and 58 males with size range: 52.3 – 85.7 cm), and *Isogomphodon oxyrinchus* (44 females with size range: 40.6 – 144.0 cm and 30 males with size range: 70.4– 115.0 cm) (**Table 1**).

In order to annul the effect of specimen size, the morphometric measurements were transformed using the Isometric Burnaby method, which projects

a series of measured distances within an orthogonal space. The data set was divided into three categories: head, body, and fins. The discriminant analysis (DA) was used to distinguish the differences between sexes in a dependent non-metric variable that characterized the several known classes. Therefore, the objective is to understand the differences between the sexes and predict the probability of an individual belonging

**Table 1.** Morphometric measures taken from shark species of the Carcharhinidae family caught in the State of Maranhão, Brazil. Ac = Acronyms.

| Head measures                   | Ac  | Fin measures                     | Ac   | Body measures                                   | Ac    |
|---------------------------------|-----|----------------------------------|------|---|-------|
| Upper labial fold               | ULF | D1 base                          | D1B  | Total length                                    | TL    |
| Lower labial fold               | LLF | Posterior margin D1              | PMD1 | Fork length                                     | FL    |
| Mouth width                     | MW  | Anterior margin D1               | AMD1 | Snout to nostrils                               | S_N   |
| Mouth length                    | ML  | Internal margin D1               | IMD1 | Snout to orbit                                  | S_O   |
| Inter-orbital                   | IO  | D1 height                        | D1H  | Snout to mouth                                  | S_M   |
| Orbit diameter                  | OD  | Free margin D1                   | FMD1 | Snout to 1 <sup>st</sup> gill slits             | S1G   |
| Inter-nostril internal distance | INI | D2 base                          | D2B  | Snout to 2 <sup>nd</sup> gill slits             | S2G   |
| Inter-nostril                   | IN  | Posterior margin D2              | PMD2 | Snout to 3 <sup>rd</sup> gill slits             | S3G   |
|                                 |     | Anterior margin D2               | AMD2 | Snout to 4 <sup>th</sup> gill slits             | S4G   |
|                                 |     | Internal margin D2               | IMD2 | Snout to 5 <sup>th</sup> gill slits             | S5G   |
|                                 |     | D2 height                        | D2H  | Snout to pectoral fin origin                    | SP1FO |
|                                 |     | Anal base                        | AB   | Snout to pectoral fin, end of base              | SP1FE |
|                                 |     | Posterior margin Anal            | PMA  | Snout to pelvic fin origin                      | SP2FO |
|                                 |     | Anterior margin Anal             | AMA  | Snout to 1 <sup>st</sup> dorsal fin origin      | SD1FO |
|                                 |     | Internal margin Anal             | IMA  | Snout to 2 <sup>nd</sup> dorsal fin origin      | SD2FO |
|                                 |     | Anal height                      | AH   | Snout to anal fin origin                        | SAFO  |
|                                 |     | Pectoral base                    | PB   | Snout to upper caudal fin                       | SUCF  |
|                                 |     | Posterior margin P1              | PMP1 | Snout to lower caudal fin                       | SLCF  |
|                                 |     | Anterior margin P1               | AMP1 | Inter-dorsal fins                               | ID    |
|                                 |     | Internal margin P1               | IMP1 | Pectoral to pelvic fin                          | PP2   |
|                                 |     | Pelvic base                      | P2B  | Pelvic to anal fin                              | P2A   |
|                                 |     | Posterior margin P2              | PMP2 | 2 <sup>nd</sup> dorsal to anal fin              | D2A   |
|                                 |     | Anterior margin P2               | AMP2 | Anal to caudal fin                              | AC    |
|                                 |     | Internal margin P2               | IMP2 | 1 <sup>st</sup> gill slits length               | 1GL   |
|                                 |     | Dorsal caudal fin lobe           | DCFL | 3 <sup>rd</sup> gill slits length               | 3GL   |
|                                 |     | Ventral caudal fin lobe          | VCFL | 5 <sup>th</sup> gill slits length               | 5GL   |
|                                 |     | Notch of the caudal fin's length | NCFL | Width of the trunk in the pectoral fin origins  | WTP1O |
|                                 |     | Notch of the caudal fin's depth  | NCFD | Height of the trunk in the pectoral fin origins | HTP1O |
|                                 |     |                                  |      | 1 <sup>st</sup> to 5 <sup>th</sup> gill slits   | 1-5G  |

to a particular sex by using the independent metric variables. This analysis was performed using the Paleontological Statistics Software (PAST) Package for education and data analysis, version 3 (Hammer et al. 2001). After normality of the data and homogeneity of the variances were verified, all variables were also tested dichotomously between sexes in order to detect possible sexual dimorphisms, with the Student's t-test considering a significance of 0.05. This analysis was performed using the program R (R Core Team 2013).

## Results and discussion

A total of 213 specimens were caught in shallow waters of Maranhão state, Amazonian Equatorial Coast. The discriminant analysis for *Isogomphodon oxyrinchus* showed no significant difference among the head measurements between sexes, while the measurements of body and fins resulted in discriminant effects (DA(HEAD) = 0.711,  $p = 0.7332$ , DA(BODY) = 7.57,  $p < .0001$ , DA(FINS) = 16.124,  $p < .0001$ ). This species showed

**Table 2.** Results of t-student test for *Rhizoprionodon porosus* and *Carcharhinus porosus*. F – Female, M – Male, SD – standard deviation, T – t-test result and p – p-value (considered significant if  $p < 0.05$ ).

| <i>R. porosus</i> | Variable | F      | SD     | M      | SD     | T     | p      | Variable | F      | SD     | M     | SD     | T     | p      |
|-------------------|----------|--------|--------|--------|--------|-------|--------|----------|--------|--------|-------|--------|-------|--------|
|                   | ULF      | 1.276  | ±0.183 | 1.610  | ±0.277 | -3.75 | 0.0005 | 3GL      | 1.467  | ±0.103 | 2.002 | ±0.278 | -4.62 | <.0001 |
|                   | LLF      | 0.817  | ±0.098 | 1.121  | ±0.176 | -4.12 | 0.002  | 5GL      | 1.067  | ±0.051 | 1.545 | ±0.308 | -3.76 | <.0001 |
|                   | MW       | 3.100  | ±0.253 | 4.200  | ±0.679 | -3.90 | 0.0003 | D1B      | 5.250  | ±0.207 | 6.493 | ±0.697 | -4.30 | <.0001 |
|                   | ML       | 4.017  | ±0.495 | 5.441  | ±0.741 | -4.54 | <.0001 | PMD1     | 5.450  | ±0.301 | 6.524 | ±1.028 | -2.52 | <.0001 |
|                   | IO       | 4.800  | ±0.456 | 6.619  | ±0.746 | -5.79 | <.0001 | AMD1     | 6.483  | ±0.479 | 8.000 | ±0.755 | -4.76 | <.0001 |
|                   | IN       | 2.850  | ±0.242 | 3.750  | ±0.375 | -5.67 | <.0001 | D1H      | 5.417  | ±0.360 | 6.112 | ±0.498 | -3.28 | 0.0020 |
|                   | S_N      | 2.817  | ±0.348 | 3.671  | ±0.390 | -5.07 | <.0001 | D2B      | 1.633  | ±0.280 | 2.088 | ±0.230 | -4.40 | <.0001 |
|                   | S_O      | 4.817  | ±0.213 | 5.912  | ±0.526 | -5.00 | <.0001 | PMD2     | 2.783  | ±0.040 | 3.626 | ±0.400 | -5.10 | <.0001 |
|                   | S_B      | 4.550  | ±0.197 | 5.536  | ±0.488 | -4.85 | <.0001 | AMD2     | 1.900  | ±0.400 | 2.205 | ±0.223 | -2.81 | 0.0072 |
|                   | S1G      | 10.450 | ±0.595 | 13.486 | ±1.480 | -4.93 | <.0001 | IMD2     | 2.433  | ±0.413 | 3.188 | ±0.339 | -4.97 | <.0001 |
|                   | S2G      | 10.983 | ±0.577 | 14.188 | ±1.544 | -4.99 | <.0001 | D2H      | 1.317  | ±0.183 | 1.710 | ±0.309 | -3.02 | 0.0041 |
|                   | S3G      | 11.567 | ±0.575 | 14.945 | ±1.644 | -4.95 | <.0001 | AB       | 2.717  | ±0.098 | 3.229 | ±0.407 | -3.04 | <.0001 |
|                   | S4G      | 12.117 | ±0.624 | 15.648 | ±1.687 | -5.04 | <.0001 | PMA      | 3.167  | ±0.121 | 3.843 | ±0.419 | -3.90 | <.0001 |
|                   | S5G      | 12.583 | ±0.624 | 15.017 | ±1.878 | -4.41 | <.0001 | AMA      | 2.400  | ±0.328 | 2.929 | ±0.269 | -4.37 | <.0001 |
|                   | SP1FE    | 14.283 | ±0.682 | 18.000 | ±1.969 | -4.55 | <.0001 | AH       | 1.783  | ±0.229 | 2.045 | ±0.288 | -2.13 | 0.0386 |
|                   | SP2FO    | 25.850 | ±1.893 | 32.712 | ±3.224 | -5.06 | <.0001 | PB       | 3.233  | ±0.287 | 3.881 | ±0.422 | -3.62 | <.0001 |
|                   | SD1FO    | 17.133 | ±0.880 | 22.252 | ±2.366 | -5.21 | <.0001 | PMP1     | 5.183  | ±0.895 | 6.636 | ±0.626 | -5.04 | <.0001 |
|                   | SD2FO    | 35.633 | ±1.894 | 45.890 | ±6.723 | -3.68 | <.0001 | AMP1     | 7.283  | ±0.457 | 9.248 | ±0.950 | -4.94 | <.0001 |
|                   | SAFO     | 34.050 | ±1.923 | 44.798 | ±4.512 | -5.72 | <.0001 | IMP1     | 2.567  | ±0.051 | 3.036 | ±0.304 | -3.73 | 0.0005 |
|                   | SUCF     | 41.817 | ±2.487 | 55.086 | ±6.369 | -5.01 | <.0001 | P2B      | 2.650  | ±0.207 | 3.293 | ±0.343 | -4.44 | <.0001 |
|                   | SLCF     | 41.067 | ±2.631 | 54.350 | ±6.174 | -5.16 | <.0001 | PMP2     | 2.450  | ±0.137 | 2.898 | ±0.322 | -3.34 | 0.0017 |
|                   | ID       | 13.733 | ±0.833 | 18.462 | ±2.048 | -5.55 | <.0001 | AMP2     | 2.633  | ±0.206 | 3.393 | ±0.389 | -4.66 | <.0001 |
|                   | PP2      | 12.533 | ±1.506 | 15.831 | ±1.560 | -4.86 | <.0001 | IMP2     | 3.133  | ±0.216 | 2.279 | ±0.325 | 6.22  | <.0001 |
|                   | P2A      | 6.450  | ±1.459 | 8.983  | ±1.156 | -4.87 | <.0001 | DCFL     | 15.500 | ±0.784 | 18.01 | ±2.350 | -2.57 | 0.0134 |
|                   | D2A      | 4.767  | ±0.680 | 6.210  | ±0.687 | -4.81 | <.0001 | VCFL     | 6.583  | ±0.778 | 8.157 | ±0.580 | -5.96 | <.0001 |
|                   | AC       | 4.967  | ±0.582 | 6.355  | ±0.713 | -4.54 | <.0001 | NCFD     | 1.750  | ±0.083 | 1.917 | ±0.139 | -2.84 | 0.0067 |
|                   | 1GL      | 1.083  | ±0.040 | 1.479  | ±0.190 | -5.02 | <.0001 |          |        |        |       |        |       |        |

only three significant variations (Paired t test,  $t_{P2A} = -3.47$ ,  $p = 0.0009$ ;  $t_{PMP2} = 2.29$ ,  $p = 0.0253$ ;  $t_{IMP2} = 9.07$ ,  $p < 0.0001$ ) in the t-test analysis conducted for each of the 65 morphometric measurements.

Distinctions between sexes for the *Rhizoprionodon porosus* were identified by all the discriminant analysis. Significant head, body, and fin measurements were obtained (DA(HEAD) = 3.641,  $p = 0.0013$ , DA(BODY) = 3.509,  $p = 0.0036$ , DA(FINS) = 68.176,

$p < .0001$ ). The t-test result identified 53 significantly distinct morphometric measurements among males and females (Table 2). Concerning *Carcharhinus porosus*, only the head measurements were statistically significant in the discriminant analysis, while the measures of body and fins were not (DA(HEAD) = 3.578,  $p = 0.0091$ , DA(BODY) = 1.319,  $p = 0.3726$ , DA(FINS) = 1.975,  $p = 0.5174$ ). The t-test showed 54 morphometric measurements significantly different between males and females (Table 2).

| <i>C. porosus</i> | Variable | F      | SD     | M      | SD     | T    | p      | Variable | F      | SD     | M     | SD     | T    | p      |
|-------------------|----------|--------|--------|--------|--------|------|--------|----------|--------|--------|-------|--------|------|--------|
|                   | SLS      | 0.300  | ±0.081 | 0.200  | ±0.077 | 3.29 | 0.0027 | 5GL      | 1.695  | ±0.297 | 1.391 | ±0.144 | 3.16 | 0.0037 |
|                   | MW       | 4.063  | ±0.501 | 3.118  | ±0.543 | 4.83 | <.0001 | D1B      | 6.900  | ±0.952 | 5.346 | ±0.621 | 4.83 | <.0001 |
|                   | ML       | 5.442  | ±0.683 | 4.182  | ±0.485 | 5.37 | <.0001 | PMD1     | 7.558  | ±1.186 | 5.546 | ±0.747 | 5.06 | <.0001 |
|                   | IO       | 6.353  | ±0.931 | 4.946  | ±0.965 | 3.93 | 0.0005 | AMD1     | 7.979  | ±1.208 | 6.182 | ±0.702 | 4.49 | 0.0001 |
|                   | IN       | 3.468  | ±0.400 | 2.682  | ±0.651 | 4.12 | 0.0003 | FMD1     | 3.537  | ±0.450 | 2.455 | ±0.317 | 7.01 | <.0001 |
|                   | S_N      | 3.821  | ±0.369 | 3.082  | ±0.312 | 5.58 | <.0001 | D1H      | 6.253  | ±1.053 | 4.573 | ±0.703 | 4.70 | <.0001 |
|                   | S_O      | 6.100  | ±0.597 | 4.882  | ±0.666 | 5.16 | <.0001 | D2B      | 2.258  | ±0.403 | 1.846 | ±0.206 | 3.15 | 0.0039 |
|                   | S_B      | 5.151  | ±0.539 | 4.718  | ±0.599 | 3.72 | 0.0009 | PMD2     | 3.074  | ±0.506 | 2.373 | ±0.431 | 3.85 | 0.0006 |
|                   | S1G      | 17.911 | ±2.213 | 14.645 | ±1.427 | 4.38 | 0.0002 | IMD2     | 2.737  | ±0.321 | 2.027 | ±0.195 | 6.61 | <.0001 |
|                   | S2G      | 13.611 | ±1.600 | 11.100 | ±1.097 | 4.58 | <.0001 | D2H      | 1.753  | ±0.412 | 1.273 | ±0.190 | 3.62 | 0.0012 |
|                   | S3G      | 14.363 | ±1.710 | 11.700 | ±1.192 | 4.55 | <.0001 | AB       | 2.879  | ±0.404 | 2.218 | ±0.240 | 4.91 | <.0001 |
|                   | S4G      | 15.047 | ±1.750 | 12.282 | ±1.226 | 4.61 | <.0001 | PMA      | 2.790  | ±0.546 | 2.146 | ±0.344 | 3.51 | 0.0015 |
|                   | S5G      | 15.721 | ±1.808 | 12.900 | ±1.253 | 4.56 | <.0001 | AMA      | 3.442  | ±0.567 | 2.773 | ±0.337 | 3.55 | 0.0014 |
|                   | SP1FO    | 14.389 | ±1.938 | 12.100 | ±1.141 | 3.56 | 0.0013 | IMA      | 2.642  | ±0.365 | 1.973 | ±0.214 | 5.52 | <.0001 |
|                   | SP1FE    | 17.911 | ±2.213 | 14.645 | ±1.427 | 4.38 | 0.0002 | AH       | 2.247  | ±0.453 | 1.573 | ±0.214 | 4.61 | <.0001 |
|                   | SP2FO    | 30.674 | ±4.610 | 24.209 | ±2.364 | 4.3  | 0.0002 | PB       | 4.190  | ±0.628 | 3.303 | ±0.234 | 5.27 | <.0001 |
|                   | SD1FO    | 19.711 | ±2.360 | 16.455 | ±1.496 | 4.11 | 0.0003 | PMP1     | 8.105  | ±1.425 | 5.755 | ±0.759 | 5.05 | <.0001 |
|                   | SD2FO    | 40.400 | ±6.231 | 32.300 | ±3.323 | 3.98 | 0.0004 | AMP1     | 9.221  | ±2.290 | 7.400 | ±0.883 | 2.52 | 0.0179 |
|                   | SAFO     | 39.000 | ±6.001 | 31.218 | ±3.212 | 3.96 | 0.0005 | IMP1     | 3.447  | ±0.545 | 2.709 | ±0.298 | 4.13 | 0.0003 |
|                   | SUCF     | 46.763 | ±6.045 | 38.073 | ±3.921 | 4.26 | 0.0002 | P2B      | 3.205  | ±0.458 | 2.436 | ±0.459 | 4.42 | 0.0001 |
|                   | SLCF     | 45.558 | ±5.952 | 37.145 | ±3.872 | 4.19 | 0.0003 | PMP2     | 3.253  | ±0.489 | 3.436 | ±0.528 | 3.85 | 0.0006 |
|                   | ID       | 14.458 | ±1.993 | 11.573 | ±1.358 | 4.25 | 0.0002 | AMP2     | 3.663  | ±0.409 | 2.736 | ±0.429 | 5.87 | <.0001 |
|                   | PP2      | 13.053 | ±1.961 | 10.673 | ±1.149 | 3.66 | 0.0010 | IMP2     | 3.984  | ±0.600 | 1.709 | ±0.661 | 9.64 | <.0001 |
|                   | D2A      | 4.726  | ±0.688 | 4.109  | ±0.568 | 2.51 | 0.018  | DCFL     | 15.800 | ±1.958 | 12.99 | ±1.497 | 4.10 | 0.0003 |
|                   | AC       | 4.942  | ±0.708 | 4.191  | ±0.678 | 2.84 | 0.083  | VCFL     | 7.726  | ±1.192 | 6.218 | ±0.625 | 3.88 | 0.0006 |
|                   | 1GL      | 1.503  | ±0.339 | 1.227  | ±0.161 | 2.54 | 0.0168 | ANCFL    | 4.500  | ±0.785 | 3.491 | ±0.493 | 3.83 | 0.0007 |
|                   | 3GL      | 2.068  | ±0.351 | 1.627  | ±0.190 | 3.83 | 0.0007 | NCFD     | 2.258  | ±0.285 | 1.864 | ±0.283 | 3.65 | 0.0011 |

The presence and absence of variations found in the three species could be related to different factors. According to Navarro et al. (2013) foraging specialization occurs between sexes within marine predators, including sharks. This could result in the development of distinctive features for each sex, which can arise in response to an increase in the competition for food resources (Wearmouth & Sims 2008). However, we discard this factor for two reasons: (1) *I. oxyrinchus* showed no differences in head measurements, the area most related to this process. This suggests that if differences in dietary habits existed, they were not sufficient to derive any morphological variations among sexes of this species; (2) The several studies that have evaluated feeding habits in Maranhão's coast have not found any sexually dimorphic diets for these three species (Lessa & Almeida 1997, Lessa et al. 1999, Silva & Almeida 2001, Almeida et al. 2011).

Another factor related to sexual dimorphism is body size. Sexual dimorphism expressed in body size is common among viviparous and ovoviviparous shark species (Sims 2003). According to Lessa et al. (2000), Machado et al. (2000), and Santos et al. (2000) females of the three species studied, as the vast majority of shark species, tend to be larger than males. This is the case for *I. oxyrinchus* and *R. porosus*, whose differences in body size can be related to the reproductive pattern of shark females. Females have larger energy expenditure than males due to the selection pressure for large offspring (Sims 2003). Additionally, males and females may present different growth rates as juveniles, which directly influences body size (Lessa et al. 2000). This hypothesis was also discarded given that the results were presented with the size effects standardized, preventing the size of the individuals from influencing the results.

In the present study, fin measurements for *R. porosus* and *I. oxyrinchus* showed significant variations. Likewise, head measurements for *R. porosus* and *C. porosus* also demonstrated significant values. Due to the lack of documented ecological data for these species it is hard to assess the more likely cause for these results. It is hypothesized that the distinction of head and fins may be related to reproductive behavior. According to Crooks & Waring (2013a)

sharks usually display a complex pattern of reproductive behavior in which the male has to bite one of the female's fins to enable copulation. The differences found may suggest adaptations of the male's head to bite and of the female's fins to receive these bites.

The t-test results for males and females of *R. porosus* and *C. porosus* showed the same morphological patterns. According to Lessa (1997), these species are very similar morphologically, which makes it difficult to distinguish them taxonomically, thus explaining these similar results. The t-test result for *I. oxyrinchus* was significant only for the pelvic area and may be related to the parturition and copulation processes. All three species analyzed were viviparous, and therefore had the same pelvic measurements with statistically significant values, as expected due to the known sexual dimorphism in primary characters of sharks.

## Conclusion

A possible cause for the variations in the external morphology among males and females of *I. oxyrinchus*, *R. porosus* and *C. porosus* is their reproductive patterns. It is clear that in order to further elucidate and test this hypothesis, further research on reproductive and behavioral features for each species must be carried out. This study revealed important details about the sexual dimorphism of *Isogomphodon oxyrinchus*, *Rhizoprionodon porosus* and *Carcharhinus porosus* through morphometric analysis, and will help facilitate the development of future studies in the area.

## Acknowledgements

We would like to thank REVIZEE, CNPq, and BASA for the financial support and Universidade Federal do Maranhão for the logistical support.

## Conflicts of interest

The authors certify that there are no conflicts of interest related to Individual Authors' Commitments, Project Support, Commitments of Editors, Journal Staff, or Reviewers. The authors also certify that they have NO affiliations with or involvement in any organization or entity with any financial interest or non-financial interest in the subject matter or materials discussed in this manuscript.

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### Dimorfismo sexual de tiburones de la costa ecuatorial amaz nica

**Resumen.** El dimorfismo sexual es una caracter stica presente en diversos grupos de vertebrados. Machos y hembras de Chondrichthyes se identifican f cilmente por la presencia de una estructura para la fertilizaci n interna, el clasper, y otras peque as diferencias en las caracter sticas secundarias. Sin embargo, los estudios que eval an estas variaciones en la morfolog a entre los sexos son poco frecuentes. El objetivo de este trabajo es identificar el dimorfismo sexual en tres especies de tiburones de la familia Carcharhinidae (*Rhizoprionodon porosus*, *Carcharhinus porosus* y *Isogomphodon oxyrinchus*) con el uso de herramientas morfom tricas. Se capturaron un total de 213 ejemplares en la costa Ecuatorial Amaz nica y fueron analizados utilizando 65 caracteres morfom tricos. Fueron aplicadas an lisis discriminantes y prueba t-student para mostrar las diferencias morfol gicas entre los sexos. Se observ  dimorfismo sexual en diferentes niveles para las tres especies. Sugerimos que la explicaci n m s probable para la presencia de estas variaciones est  relacionada con las caracter sticas reproductivas y el comportamiento de c pula.

**Palabras clave:** dimorfismo sexual, tiburones, aspectos ecol gicos; Carcharhinidae

### Dimorfismo sexual de tubar es da costa equatorial amaz nica

**Resumo.** O dimorfismo sexual   uma caracter stica presente em v rios grupos de vertebrados. Machos e f meas de Chondrichthyes s o facilmente identificados pela presen a de uma estrutura para a fertiliza o interna, o clasper, al m de outras diferen as em caracter sticas secund rias. No entanto, estudos que avaliam estas varia oes na morfologia externa entre os sexos s o raros. O principal objetivo deste estudo   identificar dimorfismo sexual em tr s esp cies de tubar o da fam lia Carcharhinidae (*Rhizoprionodon porosus*, *Carcharhinus porosus* e *Isogomphodon oxyrinchus*) atrav s do uso de ferramentas morfom tricas. Um total de 213 esp cimes foi capturado na costa Equatorial Amaz nica e analisado atrav s de 65 caracteres morfom tricos. An lises discriminantes e teste t-student foram aplicados para evidenciar as diferen as morfol gicas entre os sexos. N s observamos dimorfismo sexual, em diferentes n veis, para as tr s esp cies estudadas e sugerimos que a causa mais prov vel para a presen a de tais varia oes est  relacionada a caracter sticas reprodutivas e comportamentos populat rios.

**Palavras-chave:** dimorfismo sexual, tubar es, aspectos ecol gicos, Carcharhinidae