

## ***Mobula kuhlii* cleaning station identified at an inshore reef in southern Mozambique**

Calum JG Murie, Andrea D Marshall

Cleaning interactions between the short fin devil ray, *Mobula kuhlii*, and the blue streaked cleaner wrasse, *Labroides dimidiatus*, were observed at two sites on a single reef in southern Mozambique. Cleaning interactions were filmed and described, with the number and location of interactions recorded and subsequently binned into six distinct body patches. Cleaners preferentially foraged within certain ray body patches, and this was found to vary between the two sites, possibly signifying that variations in a habitats composition can influence cleaning. *Mobula kuhlii* were not found to clean sympatrically with their close relatives in the *Manta* genus, implying their cleaning requires a distinct habitat or that niche partitioning is required to stem competition for host cleaner fishes attention. In total, 15 individuals were observed interacting with cleaners, and they never arrived alone, suggesting they may travel to cleaning areas in an aggregative manner.

1 ***Mobula kuhlii* cleaning station identified at an inshore reef**  
2 **in southern Mozambique**

3  
4 Calum J.G. Murie<sup>1</sup> and Andrea D. Marshall<sup>1,2</sup>  
5

6 <sup>1</sup>Marine Megafauna Association, Manta Ray and Whale Shark Research Centre, Tofo Beach, Mozambique

7 <sup>2</sup>Marine Megafauna Foundation, Truckee, California, USA  
8

9 Corresponding Author:

10 Andrea Marshall

11 Marine Megafauna Foundation, Truckee, California, USA

12 Email: [andrea@marinemegafauna.org](mailto:andrea@marinemegafauna.org)  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23

24 **Abstract**

25 Cleaning interactions between the short fin devil ray, *Mobula kuhlii*, and the blue streaked  
26 cleaner wrasse, *Labroides dimidiatus*, were observed at two sites on a single reef in southern  
27 Mozambique. Cleaning interactions were filmed and described, with the number and location of  
28 interactions recorded and subsequently binned into six distinct body patches. Cleaners  
29 preferentially foraged within certain ray body patches, and this was found to vary between the  
30 two sites, possibly signifying that variations in a habitats composition can influence cleaning.  
31 *Mobula kuhlii* were not found to clean sympatrically with their close relatives in the *Manta*  
32 genus, implying their cleaning requires a distinct habitat or that niche partitioning is required to  
33 stem competition for host cleaner fishes attention. In total, 15 individuals were observed  
34 interacting with cleaners, and they never arrived alone, suggesting they may travel to cleaning  
35 areas in an aggregative manner.

36

37

38

39

40

41

42

43

44

45

46

47

48 **Introduction**

49 The *Mobula* genus contains nine dramatically understudied species. Currently three are classified  
50 by the I.U.C.N. as data deficient, four as near threatened, one as vulnerable to extinction, and one  
51 as endangered (IUCN, 2015). There has been an alarming lack of scientific publications focusing  
52 on the genus, and as a result particularly little is known about their biology, behaviour or daily  
53 movements (Couturier et al., 2013). Here we provide the first description of *Mobula* rays  
54 interacting with cleaner organisms.

55

56 The subfamily Mobulidae contains the *Mobula* as well as the two species of *Manta* (Ward-Page  
57 et al., 2013). Similar to the *Manta* species, *Mobula* are lowly fecund, birthing a single live young  
58 per pregnancy (Couturier et al., 2013). As a result, the targeted removal of individuals by  
59 directed fisheries has led to ubiquitous global population declines (Couturier et al., 2013). A  
60 frequently sighted species in southern Mozambique is the short fin devil ray, *Mobula kuhlii*, a  
61 species which is also commonly targeted by local artisanal fisherman (Couturier et al., 2013).  
62 Unfortunately, the extent with which they are targeted within the region has not been established  
63 as monitoring local artisanal fisheries is difficult along such a remote coastline and Mozambique  
64 is yet to record any landings with the F.A.O. (Ward-Paige, Davis & Worm, 2013).

65

66 While descriptions of *Mobula* rays' life history are incomplete, their diurnal behavioural  
67 patterns, particularly that of nocturnal foraging, significantly resembles the manta rays (Croll et  
68 al., 2012; Couturier et al., 2013). Both the bent tail devil ray, *Mobula thurstoni*, and the spine tail  
69 devil ray, *Mobula japonica*, have been shown to feed near exclusively on euphysiids which

70 nocturnally migrate to shallow waters (Gadig, Namora & Motta, 2003; Croll et al., 2012). During  
71 the hours of daylight, *Mobula* have been reported to remain in warm, shallow waters (Croll et al.,  
72 2012). Temperature can significantly influence poikilotherm physiological processes, and this is  
73 likely a key benefit to remaining in warm shallow waters, despite the decreased opportunities for  
74 foraging (Hochachka and Somero, 2002). Migrations inshore could also be attributed to social  
75 interactions, predator avoidance or their need to visit cleaning stations (Dewar, Mous &  
76 Domeier, 2008; Marshall, 2009; O'Shea, Kingsford & Seymour, 2010).

77

78 Cleaning ecology describes the interactions between client species and specialised cleaner  
79 organisms. Manta rays, the closest living relative to the *Mobula* species, are regularly observed  
80 interacting with cleaner fish hosts at aggregation sites worldwide (Dewar, Mous & Domeier,  
81 2008; Marshall, 2009; O'Shea, Kingsford & Seymour, 2010). They, in addition to a diverse set  
82 of client species, regularly visit reef systems to have mucus, algal build-up, necrotic tissue and  
83 ectoparasites removed by the cleaners (Grutter and Bshary, 2003). The most studied marine  
84 cleaner species is *Labroides dimidiatus*, the blue streaked cleaner wrasse. The wrasse primarily  
85 forages on ectoparasites, and preferentially targets gnathiid isopods (Grutter, 1997; Grutter and  
86 Poulin, 1998; Grutter and Bshary, 2003). The blue streaked wrasses presence on reefs has been  
87 shown to both increase species diversity and positively influence client health (Grutter, Murphy  
88 & Choat, 2003; Ros et al., 2010). Herewithin, we provide the first description of mobula rays  
89 being serviced by the blue streaked cleaner wrasse.

90

91

92 ***Materials and Methods***

93 Following numerous opportunistic encounters and reports from dive operators where mobula  
94 rays were observed soliciting cleaning interactions at a shallow reef in the Inhambane Province,  
95 5.17 hours of SCUBA dive transects (n = 7) were conducted to survey for established cleaning  
96 stations. The study reef, colloquially named “Two Mile”, is located within the Bazaruto  
97 Archipelago National Park, which is situated in the northern extent of the Inhambane province of  
98 southern Mozambique.

99

100 Subsequently, across two days and five separate S.C.U.B.A. dives, three and a half hours were  
101 spent recording cleaning interactions between *Mobula kuhlii* and *Labroides dimidiatus* (Fig. 1).  
102 Observations were split between two sites on ‘Two-Mile Reef’. Between low and mid tide, the  
103 two sites are separated by a rocky ridge breaking the water’s surface. One hour and five minutes  
104 were spent observing cleaning interactions at Fish Bowl, which sits at a depth of 8-14m on the  
105 leeward side of the reef. Two hours and twenty five minutes were spent observing cleaning  
106 interactions at Devil's Dance, which lies on the reefs seaward side at a depth of 16-20m. All  
107 research was conducted under approval of the National Administration of Conservation Areas  
108 (Department under the Environment Ministry of Mozambique) and was ethically approved by  
109 the Marine Megafauna Foundation.

110

111 During dives, two observers were present. One video-recorded the cleaning interactions, while  
112 the other positioned themselves on the edge of visibility to avoid influencing behaviours, but  
113 remaining accessible for safety. Videos were downloaded, replayed frame by frame, and  
114 interactions documented. A single interaction was characterised by a cleaner’s mouth making

115 contact with the ray's body. The location of each interaction was noted on a template of a *M.*  
116 *kuhlii* rays' body, and then binned within the six ray body patches for analysis (Fig. 2).

117

118 Statistical analysis was performed within R statistics (R version 3.1.3: "Smooth Sidewalk"). Chi  
119 squared testing was used to examine if interactions between the cleaner wrasse and rays differed  
120 from a uniform distribution between the six body patches and if interaction distributions varied  
121 between the two sampled sites.

122

123

## 124 **Results**

125 *Mobula kuhlii* are regularly encountered in southern Mozambique. Despite the authors, and other  
126 members of their team, completing 1,853 survey dives in the region (2003-2015) including the  
127 intensive surveillance of 10 inshore reefs known to host manta cleaning stations, no *Mobula*  
128 species have ever been seen interacting with cleaner fish other than on the single reef monitored  
129 in this study.

130

131 Preliminary surveying transects (n =7) were conducted on both the ocean, and lee, side of the  
132 study reef to identify established cleaning stations. Rays were only present on three of these  
133 initial transects but when present had a mean sighting of  $3.67 \pm 2.66$  (mean  $\pm$  SE). During this  
134 period of surveying, 11 *Mobula kuhlii* were observed cleaning in two specific areas of the reef  
135 known locally as 'Fish Bowl' and 'Devil's Dance'. At a later stage, during focal dives (n = 5) to  
136 observe cleaning behaviour, 15 individuals were observed interacting with cleaners. On two of

137 the dives rays were absent but when present the mean sighting was 5 individuals  $\pm$  0.58 (mean  $\pm$   
138 SE) per dive.

139

140 The recorded cleaning bouts at the monitored cleaning stations lasted for a mean time of 162s  $\pm$   
141 32s with the rays receiving direct cleaning interactions from the host fish for 67s  $\pm$  15s (mean  $\pm$   
142 SE). During cleaning bouts, the rays spent a large proportion of their time repositioning  
143 themselves in order to pass over spatially finite areas of the reef ( $t = 95s \pm 23s$ , (mean  $\pm$  SE)). A  
144 maximum of three rays were observed receiving simultaneous interactions, despite six being  
145 contemporaneously present.

146

147 The distribution of interactions varied significantly between the six body patches ( $\chi^2 = 23.419$ ,  $df$   
148  $= 5$ ,  $p < 0.001$ ). Overall, cleaners seemed to favour ventral patches, with the body patch  
149 receiving the highest absolute number of interactions (Fig. 5). Across the dorsal surface, the head  
150 patch received the most interactions from cleaner fish (Fig. 3).

151

152 Cleaner individuals also varied their patch specific foraging between the sampled sites ( $\chi^2 =$   
153  $42.895$ ,  $df = 5$ ,  $p < 0.001$ ). At Devil's Dance, all six patches received interactions. The majority  
154 were concentrated in the two head patches whilst the ventral body also received a relatively high  
155 number of interactions (Fig. 4). The interactions recorded at Fish Bowl were much less evenly  
156 distributed between the patches and were highly dominated by the two pelvic patches (Fig. 4).

157

158

159 ***Discussion***



160 As cleaning interactions provide individuals with tangible health benefits, it is widely regarded  
161 as an important aspect of a species life history (Ros et al., 2010). Cleaning behaviour has not  
162 previously been investigated for a *Mobula* species, however several reports describe both of their  
163 close relatives in the *Manta* genus cleaning habitually (Dewar, Mous & Domeier, 2008;  
164 Marshall, 2009; O'Shea, Kingsford & Seymour, 2010). During this study in southern  
165 Mozambique we observed *Mobula kuhlii* soliciting interactions from the blue streaked cleaner  
166 wrasse. These initial observations are both unique to the region and are the first time cleaning  
167 interactions similar to those commonly seen in manta rays have been reported for a *Mobula*  
168 species.

169

170 Despite almost 2,000 research dives being completed in the region across 13 years mobula rays  
171 have only ever been seen cleaning on this single reef. Intensive research has shown that manta  
172 ray species display distinct preferences for the reefs within the region that they visit to be  
173 serviced by cleaners (Marshall et al., in prep). However, despite regular surveys of over 10 major  
174 reef systems in the local area with established cleaning stations for manta rays, *Mobula* were  
175 never seen to solicit cleaning from host cleaner fish in these areas. It is likely that either mobula  
176 rays have their own, distinct, preferences for the reefs they visit to solicit cleaning services or  
177 they purposefully are partitioning their habitat to avoid competition with these larger rays  
178 species.

179

180 On the study reef in the Bazaruto Archipelago National Park *Mobula kuhlii* is commonly  
181 encountered and regularly observed being cleaned by host fish. Exhaustive survey work in the  
182 immediate area (40 km on either side of this particular reef) has failed to produce similar

183 observations. Studies have shown that clients have the ability to recognise a cleaning station, and  
184 then regularly return to it; a phenomenon often linked with the quality of the cleaning received  
185 (Tebbich, Bshary & Grutter, 2002). Variations in quality most likely arise from varying cleaner  
186 species abundance and composition, and propensity of cleaners to cheat (removal of non  
187 beneficial items by cleaners). Since *L. dimidiatus* is a true cleaner and is abundant throughout the  
188 surveyed sites, i.e. not a limiting factor in reefs shallower than 50 meters, reef preferences may  
189 rather be influenced by variations in topography and conditions, which may affect the range of  
190 movement of the cleaners and the quality of the service the cleaners can provide. Reef preference  
191 may also simply be a result of the reefs proximity to key habitats such as foraging grounds.

192

193 Group, or social, behaviour has been proposed as an important aspect of a manta ray's life history  
194 (Deakos, 2010a). Being part of a group is generally thought to increase a species awareness of  
195 predators and result in a greater probability of avoidance (Magurran, 1989; da Silva and Terhune,  
196 1998). As of yet, no investigations have focused on *Mobula* ray group behaviour. Mobulidae  
197 species have no form of parental care, and so any groupings are likely not driven by familial  
198 bonds, meaning individuals congregate opportunistically or as part of a roving clade (Deakos,  
199 2010b). During this preliminary study, no ray arrived at the designated cleaning stations alone,  
200 indicating they travel to cleaning areas as part of a structured group of some kind. Despite  
201 several rays being present around the cleaning station at once, individuals did not receive  
202 simultaneous interactions from the cleaners. So while social groups may navigate to cleaning  
203 stations together, it initially appears that at least the two monitored cleaning stations have a  
204 carrying capacity limiting the number of individuals that can be serviced at any one time. Certain  
205 individuals were noted to 'take turns' being cleaned, while other rays, that did not receive

206 cleaning services at all, appeared to wait in the vicinity for others to be finished before together  
207 moving away from the area.

208

209 Whilst on the cleaning station, individuals spent more of their time circling than actually  
210 cleaning in order to continually pass over specific areas of the reef where the cleaner hosts were  
211 aggregating. These observations suggest that at least the cleaning stations on this reef may be  
212 located in quite spatially finite regions, defined by the host fish themselves. The cleaners did not  
213 follow the rays into the water column, instead remaining within their territories until the ray  
214 returned, suggesting they themselves dictate the areas within which the rays can clean, a fact that  
215 has been proposed in previous cleaning studies (Nakashima et al., 2000; Tebbich, Bshary &  
216 Grutter, 2002). As ram ventilators (Correia, Graca & Hirofumi, 2008), mobula rays like manta  
217 rays are obligated to continually swim, meaning they are unable to hold position above a cleaners  
218 territory for long periods of time. This biological requirement, when coupled with the spatial  
219 limitations of cleaning areas, possibly limits the efficiency of the service and may dictate where  
220 this species is able to clean and under what conditions.

221

222 The distribution of cleaning interactions varied significantly between the two sites of the reef.  
223 The most obvious variations between the sites appeared to be their topography and substrate  
224 cover. Fish Bowl's benthos was covered with a bed of soft coral interspersed with loose rock,  
225 whilst Devil's Dance consisted of many large rocky outcrops encrusted with hard corals. Such  
226 differences likely influence the range and behaviour of the cleaner fish, and may alter both the  
227 wrasse's approach and their effectiveness as cleaners. Although no differences were noted during

228 our observations, varying environmental conditions, such as current strength, may also limit the  
229 cleaners movements, and may contribute to observed variations.

230

231 A fish's ectoparasite load positively correlates to body surface area and may be linked to the time  
232 a species is required to clean (Grutter, 1995; Sikkel, Fuller & Hunter, 2000). Descriptions of  
233 *Manta alfredi* cleaning have shown on average they clean for approximately twice the amount of  
234 time than the *M. kuhlii* observed within this study (Marshall et al., in prep). With surface areas  
235 orders of magnitude smaller than those of manta rays *M. kuhlii* may have much lower  
236 ectoparasite loads, which in turn may decrease the time and frequency of their cleaning  
237 requirement. Also unlike manta rays which are noted to be cleaned by a wide variety of cleaners  
238 (Marshall, 2009; O'Shea et al. 2010), *Mobula kuhlii* was only attended by a single cleaner  
239 species despite several known manta ray cleaners being present on the reef. If individual *Mobula*  
240 do indeed have low ectoparasite loads relative to an average manta ray, they may be perceived as  
241 less energetically rewarding clients for cleaner fish, perhaps explaining why interactions were  
242 only received from the small bodied, obligate cleaner, *L. dimidiatus*. With some studies  
243 indicating that cleaner fish species preferentially target certain parasite types (Marshall, 2009;  
244 Oliver et al., 2011), an alternate explanation may be that the parasites that these rays commonly  
245 possess may be favoured by this cleaner species explaining why they are attended solely by *L.*  
246 *dimidiatus*.

247

248 Cleaner fish should target body regions representing the most efficient foraging opportunities  
249 (Oliver et al., 2011). *L. dimidatus* has previously been shown to qualitatively assess a foraging  
250 opportunity and preferentially target it (Grutter, 1997). An uneven between-patch energy reward

251 may explain why the wrasse favoured certain body patches in *M. kuhlii*. The ventral body patch  
252 received the highest number of interactions suggesting the area may represent high quality  
253 foraging for the cleaners. The level of attention that this patch received may also have been  
254 exacerbated by the cleaner fish's approach from a benthic territory or by the patches relatively  
255 large surface area (Tetsuo, 1984; Green, 1994). However, as the vast majority of the recorded  
256 interactions were concentrated around the gill slits, the cleaners were more likely selectively  
257 targeting this region. Gnathiid isopods form the vast majority of *L. dimidiatus*'s diet, and manta  
258 rays are known to host dense aggregations of these parasites in their gill slits (Grutter and Poulin  
259 1998; Marshall, 2009). It is quite possible that the observed *M. kuhlii* may also have high  
260 gnathiid isopod loads in their gills, influencing the cleaners to forage in this area.

261

262

### 263 ***Conclusions***

264

265 Cleaning has never previously been investigated for a *Mobula* species. As their closest relatives,  
266 the giant and the reef manta ray (*Manta birostris*, *Manta alfredi*) extensively clean at inshore  
267 reefs, it is reasonable to assume that *Mobula* species seek out cleaning services as well. While  
268 this is the first reported account of cleaning in *Mobula*, this is likely due to lack of scientific  
269 effort. Alternatively, a variety of factors may result in these species being less heavily parasitised  
270 than manta rays requiring them to spend less time engaged in cleaning activities. It will be  
271 important to follow up on this preliminary study not just in this region or this species but across  
272 the rays in this genus. Understanding an animal's behaviour, habitat use, and movement patterns  
273 has significant implications on how it is most prudently managed. Further efforts must aim to

274 quantify the importance of this behaviour to mobula rays and better determine their use of  
275 inshore habitats during daytime hours. Being able to accurately describe the patterns of use of  
276 these critical inshore habitats may provide managers with the detail needed to more effectively  
277 safeguard these threatened species.

278

## 279 Acknowledgements

280

281 Many thanks go to Daniel Steuber for introducing us to the site and for Janneman Conradie's  
282 extensive logistical support. Thanks also to the Underwater Africa volunteers for aiding in data  
283 analysis.

284

## 285 References

286

287 Correia, JPS, Graca, JTC, Hirofumi, M. 2008. Long-term transportation, by road and air, of Devil-Ray (*Mobula*  
288 *mobular*), Meagre (*Argyrosomus regius*), and Ocean Sunfish (*Mola mola*). *Zoo Biology*. 27: 1-17.

289

290 Couturier, LIE, Marshall, AD, Jaine, FRA, Kashiwagi, T, Pierce, SJ, Townsend, KA, Weeks, SJ, Bennett, MB,  
291 Richardson, AJ. 2012. Biology, ecology and conservation of the Mobulidae. *Journal of Fish Biology* 80: 1075-1119.

292

293 Croll, DA, Newton, KM, Weng, K, Galvan-Magana, F, O'Sullivan, J, Dewar, H. 2012. Movement and habitat use  
294 by the spine-tail devil ray in the eastern Pacific Ocean. *Marine Ecology Progress Series* 465: 193-200.

295

296 da Silva, J, Terhune, JM. 1988. Harbour seals grouping as an anti-predator strategy. *Animal Behaviour* 36: 1309-  
297 1316.

298

299 Deakos, MH. 2010a. Ecology and social behaviour of a resident manta ray (*Manta alfredi*) population off Maui,  
300 Hawaii. D. Phil. Thesis, University of Hawaii. Available online:  
301 [http://www.hamerinhawaii.org/resources/deakos\\_pdf/DeakosDissertation2010.pdf](http://www.hamerinhawaii.org/resources/deakos_pdf/DeakosDissertation2010.pdf).

302

303 Deakos, MH. 2010b. Paired-laser photogrammetry as a simple and accurate system for measuring the body size of  
304 free-ranging manta rays *Manta alfredi*. *Aquatic Biology* 10:1-10.

305

306 Dewar, H, Mous, P, Domeier, M. 2008. Movements and site fidelity of the giant manta ray, *Manta birostris*, in the  
307 Komodo Marine Park, Indonesia. *Marine Biology* 155:121-133.

308

309 Gadig, OBF, Namora, RC, Motta, FDS. 2003. Occurrence of the bentfin devil ray, *Mobula thurstoni*  
310 (Chondrichthyes: Mobulidae), in the western Atlantic. *Journal of the Marine Biological Association of the U.K.* 83:  
311 869-870.

- 312  
313 Green, AL. 1994. The early life history of *Labroid* fishes at Lizard Island, Northern Great Barrier Reef. D. Phil.  
314 Thesis, James Cook University. Available online: <http://eprints.jcu.edu.au/27393/>.
- 315  
316 Grutter, AS. 1995. Relationship between cleaning rates and ectoparasite loads in coral reef fishes. Marine Ecology  
317 Progress Series 118: 51-58.
- 318  
319 Grutter, AS. 1997. Size-selective predation by the cleaner fish *Labroides dimidiatus*. Journal of Fish Biology 50:  
320 1303-1308.
- 321  
322 Grutter, AS; Poulin, R. 1998. Intraspecific and interspecific relationships between host size and the abundance of  
323 parasitic larval gnathiid isopods on coral reef fishes. Marine Ecology Progress Series 4: 263-271.
- 324  
325 Grutter, AS, Bshary, R. 2003. Cleaner wrasse prefer client mucus: support for partner control mechanisms in  
326 cleaning interactions. Biology Letters 270: 242-244.
- 327  
328 Grutter, AS, Murphy, JM, Choat, JH. 2003. Cleaner fish drives local fish diversity on coral reefs. Current Biology  
329 13: 64-67.
- 330  
331 Hochachka, PW, Somero, GN. 2002. Biochemical adaptation: mechanism and process in physiological evolution.  
332 Oxford University Press, New York, NY.
- 333  
334 International Union for the Conservation of Nature, IUCN. 2015. The IUCN Red List of Threatened Species.  
335 Version 2015-3. <http://www.iucnredlist.org>. Downloaded on 10 October 2015.
- 336  
337 Magurran, AE. 1990. The adaptive significance of schooling as an anti-predator defence in fish. Behaviour of Fish  
338 2: 51-66.
- 339  
340 Marshall, AD. 2009. Biology and population ecology of Manta birostris in southern Mozambique. D. Phil. Thesis,  
341 University of Queensland, Australia.
- 342  
343 Nakashima, Y, Sakai, Y, Karino, K, Kuwamura, T. 2000. Female-female spawning and sex change in a harem  
344 coral-reef fish, *Labroides dimidiatus*. Zoological Science 17: 967-970.
- 345  
346 Oliver, SP, Hussey, NE, Turner, JR, Beckett, AJ. 2011. Oceanic sharks clean at coastal seamount. PLOS ONE,  
347 e14755.
- 348  
349 O'Shea, OR, Kingsford, MJ, Seymour, J. 2010. Tide-related periodicity of manta rays and sharks to cleaning  
350 stations on a coral reef. Marine and Freshwater Research 61: 65-73.
- 351  
352 Ros, AFH, Lusa, J, Meyer, M, Soares, M, Oliveira, RF, Brossard, M, Bshary, R. 2010. Does access to the blue  
353 streaked cleaner wrasse *Labroides dimidiatus* affect indicators of stress and health in resident reef fishes of the red  
354 sea. Hormones and Behaviour. Published Online: doi:10.1016/j.yhbeh.2010.11.006.
- 355  
356 Sikkell, PC, Fuller, CA, Hunter, W. 2000. Habitat/sex differences in time at cleaning stations and ectoparasite loads  
357 in a Caribbean reef fish. Marine Ecology Progress Series 193: 191-199.
- 358  
359 Tebbich, S, Bshary, R, Grutter, AS. 2002. Cleaner fish, *Labroides dimidiatus*, recognised familiar clients. Animal  
360 Cognition 5: 139-145.
- 361  
362 Tetsuo, K. 1984. Social structure of the protogynous fish *Labroides dimidiatus*. Publications of the Seto Marine  
363 Biological Laboratory 293: 117-177.

364

365 Ward-Paige, CA, Davis, B, Worm, B. 2013. Global Population Trends and Human Use Patterns of Manta and  
366 Mobula Rays. PLoS ONE 8(9):e74835 [doi:10.1371/journal.pone.0074835].

367

368

369



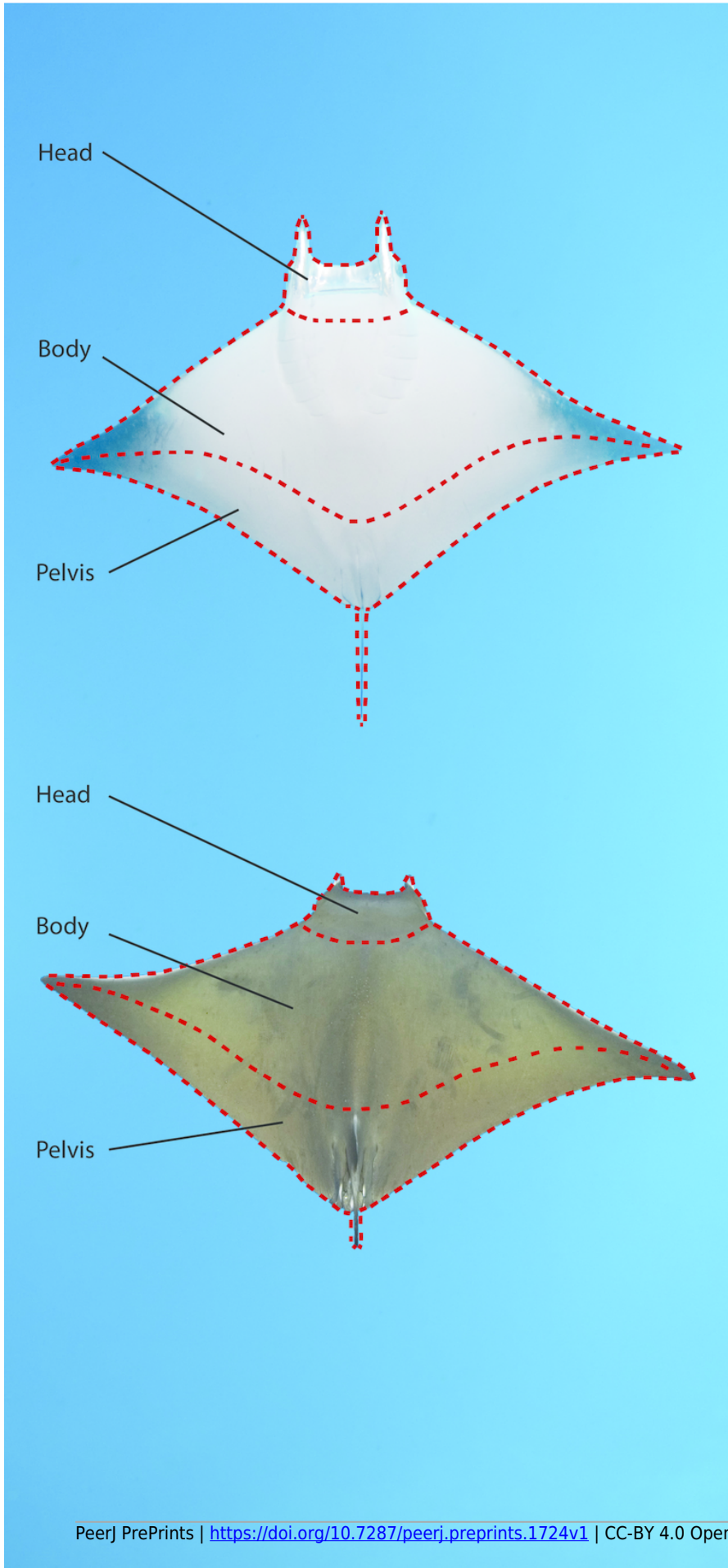
## 1

A *Mobula kuhlii* interacting with *Labroides dimidiatus* individuals during a pass over a cleaning station on the site “Devils Dance”.



## 2

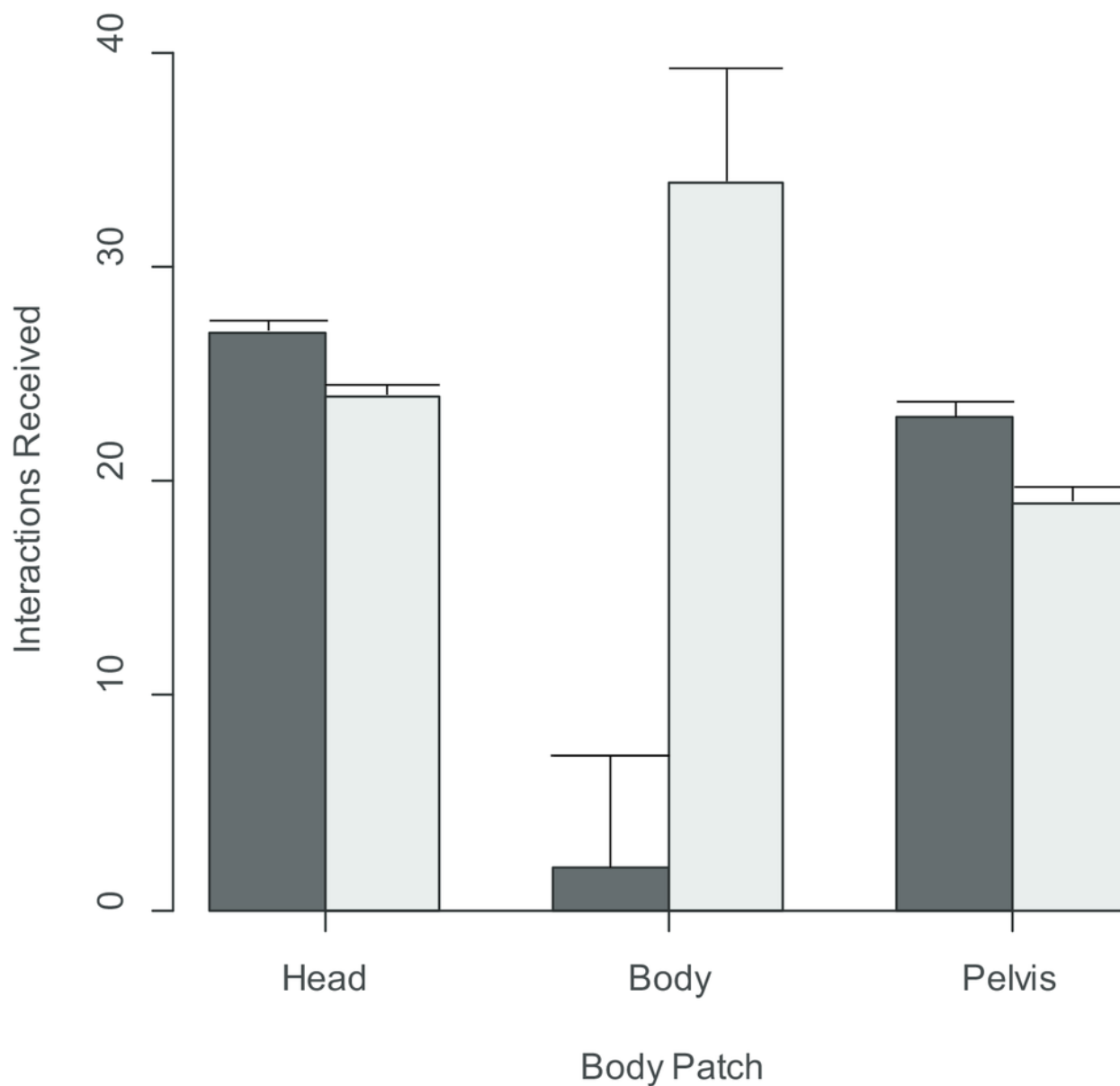
The body patches that recorded cleaning interactions between *Mobula kuhlii* and *Labroides dimidiatus* were binned within.



## 3

The mean number ( $\pm$  SE) of interactions that a ray's body patch received from cleaners.

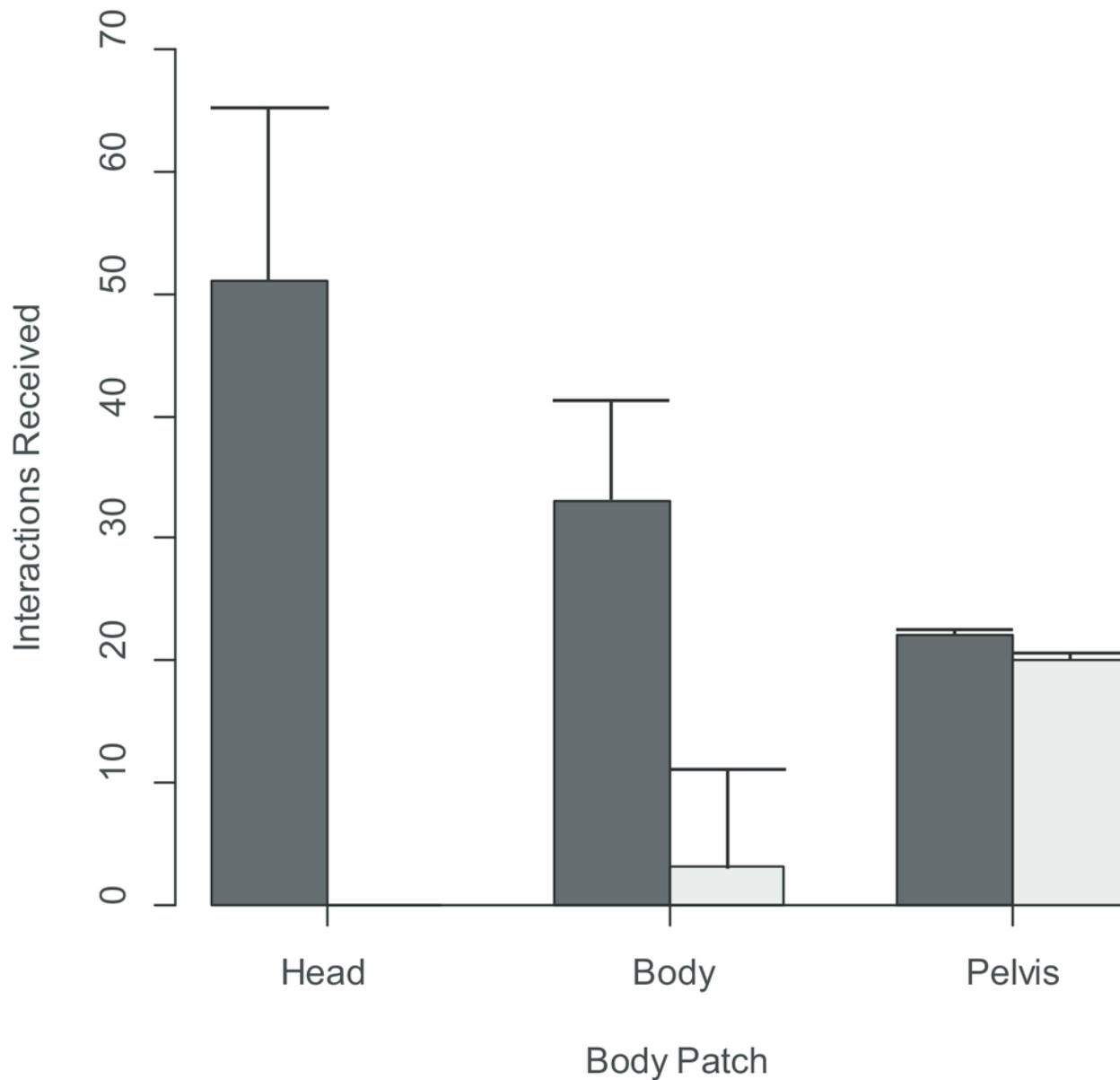
Light bars denote the ventral surface, and dark the dorsal. [b]



## 4

The mean number ( $\pm$  SE) of interactions a ray's body patch received from the cleaner fish *Labroides dimidiatus*.

Light bars symbolise the reef "fish bowl", and dark "Devil's dance".



# 5

Recorded cleaning interactions between the fish *Labroides dimidiatus* and *Mobula kuhlii*.

The origin of a line represents the point at which contact occurred. Red lines represent the site “Fish Bowl” and black lines the site “Devil's Dance”.

