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1 **Subconscious Biases in Coral Reef Fish Studies**

2

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11 **Abstract**

12 In complex, diverse ecosystems, one is faced with an exceptionally challenging decision:
13 which species to examine first, and why? This raises the question: is there evidence of
14 subconscious biases in study species selection? Likewise, in selecting methods, locations and
15 times? We addressed these questions by surveying the literature on the most diverse group of
16 vertebrates (fishes) in an iconic high-diversity ecosystem (coral reefs). The evidence suggests
17 that we select study species that are predominantly yellow. Reef fish studies also selectively
18 examine fishes that are behaviourally bold, and in warm, calm, attractive locations. Our
19 findings call for a re-evaluation of study species selection, and methodological approaches,
20 recognising the potential for subconscious biases to drive selection for species that are
21 attractive rather than important, and methods that give only a partial view of ecosystems.
22 Given the challenges faced by high diversity ecosystems, we may need to question our
23 decision-making processes.

24

25 **Keywords**

26 Biodiversity; Coral reef; Fish; Function; Subconscious bias

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37 Overview of the issue

38 Coral reefs are one of the world's most iconic high-diversity ecosystems. Such
39 diversity offers a plethora of potential study species, leaving one faced with an exceptionally
40 challenging decision: which species do we examine first? And where, when and how should
41 we conduct our study? Logically, it is in such high-diversity systems that the potential for
42 subconscious biases are likely to be greatest, given the range of study options available
43 (Bonnet et al. 2002; Clark and May 2002). This issue is particularly pressing as many high-
44 diversity ecosystems are rapidly reconfiguring in response to climate-induced environmental
45 disruption (Barlow et al. 2018; França et al. 2020). As a result, there have been urgent calls to
46 understand and maintain the ecosystem functions that sustain high-diversity ecosystems, such
47 as coral reefs, and the services they provide to humanity (Hughes et al. 2017; Brandl et al.
48 2019b). However, our understanding of ecosystem functions depends on the species we
49 examine and how we study them (Bellwood et al. 2019). Such selection processes may
50 involve subconscious biases.

51 There is a burgeoning literature on the extent, nature and impacts of subconscious
52 biases (also termed unconscious and, perhaps most accurately, implicit biases) (e.g. McNutt
53 2016; Knezek 2017; Asplund and Welle 2018; Baum and Martin 2018). These biases have
54 been repeatedly shown to influence the nature of academia, especially in terms of selecting
55 researchers for funding, promotion and publication (Wenneras and Wold 1997; Bornmann et
56 al. 2007; Moss-Racusin et al. 2012). However, there have been few critical evaluations of our
57 decisions when undertaking scientific research, especially in terms of what species we study
58 (but see Bonnet et al. 2002; Clark and May 2002), or how we conduct our research. To
59 address this knowledge gap we asked two key questions: 1) is there evidence of biases in our
60 selection of study species and, 2) are there biases in our approaches when undertaking
61 research in high-diversity systems? Ultimately, this raises the question: to what extent may
62 these biases shape our understanding of ecosystem processes?

63 To address these questions, we surveyed the published literature on the most diverse
64 group of vertebrates (fishes) in an iconic high-diversity ecosystem (coral reefs) (Fig. 1). Coral
65 reef fishes represented a particularly amenable study group because of their ease of
66 identification to a species level, taxonomic stability, pantropical distribution, and exceptional
67 taxonomic and morphological diversity (Fig. 1). Our focal literature source was the
68 international journal *Coral Reefs*; the world's primary journal for coral reef studies. This
69 journal was specifically selected for its research breadth, while offering the highest
70 concentration of papers on coral reefs. The sole restriction for papers in this journal, apart
71 from scientific merit, is that they pertain to coral reefs; there is no restriction on geographic
72 location or approach (i.e. field or experimental). Limiting our study to this one broad journal
73 therefore minimizes the potential for other biases, associated with journal selection, to
74 confound our results. For each article related to reef fishes published between 1982 and 2018
75 that involved a field-based component (e.g. fish collection, observation or quantification) (n =
76 377 articles) we recorded details pertaining to: a) selected study species, b) the month/s when
77 the field-component of the study occurred, c) fish abundance quantification methods and d)
78 the habitat where and fish quantification methods were performed (studies may involve only
79 some or all of the above; see the supporting information for a full overview of the literature
80 survey and associated methods). It is important to note that our interest is in the selection of
81 study species, locations, methods and times by researchers, i.e. decisions that will
82 fundamentally shape our understanding of these systems. The focus is solely on the decisions
83 made by the scientist(s), not the subsequent popularity or perceived importance of the study
84 based on metrics such as citations or journal impact factor.

85

86 **Species biases**

87 Although biases were recorded in all four study criteria, it was in the selection of focal
88 species that the most striking patterns were revealed. Of an estimated 6000+ reef fish species,
89 less than 7% (396 species) were selected for study, with just 0.1% (6 species) examined 10%
90 of the time (Fig. S2). Most research is restricted to a small range of quintessential coral reef
91 butterflyfishes and damselfishes. The selection of these specific species may be influenced by
92 a range of factors, however, it is particularly interesting to consider the colouration of these
93 species; almost all focal study species had bright colours, especially yellow (Fig. 2a).

94 At the broadest scale, looking at all species examined, our results strongly suggest
95 that selected study species are not random with regards to colour (Fig. 2a). Naturally, there is
96 a range of other potential explanations. Yellow fishes may be more territorial and thus easier
97 to observe, see or catch. Most importantly, yellow fish species may be more common or
98 abundant than fishes with other colours. To directly test whether species selection does favor
99 predominantly yellow species, rather than other traits (e.g. territoriality), we looked in detail
100 at the family Pomacentridae. This family was selected as it is, by far, the most frequently
101 studied, making up more than a third of all records. It also contains numerous species, with a
102 wide variety of colour patterns, thus permitting robust analyses. Focusing on the GBR, almost
103 all species had appropriate photographs available. Most importantly, the family is composed
104 of species with very similar traits (other than colour); all have relatively small body sizes, are
105 strongly site attached, show minimal diver avoidance, live in relatively shallow waters, are
106 easily collected and are omnivorous, herbivorous, or planktivorous.

107 We can therefore ask: compared to the Pomacentridae on the GBR, are study species
108 selected randomly with regards to their colour patterns? The answer is no (Fig. 2b) (see ESM
109 for statistical values). We cannot say species are selected because they are yellow, inferring
110 causation, but we can say with confidence that yellow species are studied most often (and
111 equally, that dark fishes appear to be strongly avoided). Furthermore, if we focus specifically
112 on the abundance of species (around one of the peak research locations in the GBR - Lizard
113 Island); asking if damselfish species are selected randomly with regards to their estimated
114 local abundance, we get the same pattern. Yellow species were preferred, dark ones avoided
115 (Fig. 2c). Taken together, these patterns raise questions about the possibility of a colour-
116 based bias. Interestingly, a recent study using simulated reefs provides strong
117 supporting evidence, documenting human preferences for attractive reefs organisms based on
118 colour; with a clear indication that yellow fishes are by far the most attractive (Tribot et al.
119 2019). Thus, the possibility of yellow fish being both overrepresented and positively selected
120 for based on their colour, represents a distinct possibility.

121

122 **Methodological biases**

123 We also found evidence of preference or bias in the three methodological approaches.
124 The relationship between sampling period and time of year was strongly selective, with
125 almost twice the research effort in the summer months (Fig. 3a) (offset in northern and
126 southern hemisphere locations Fig. S1). Likewise, habitats were unevenly studied, with 50%
127 of all fish censuses undertaken on the reef slope or crest where fish densities and coral cover
128 are often highest (Russ 1984; Wismer et al. 2009); they are therefore, arguably, the most
129 attractive habitats. When examined as a proportion of the available reef area, across four
130 standard reef habitats, this selectivity is striking (Fig. 3b), with 64% of the censuses looking

131 at just 35% of the reef area. Fish census methods were also revealing (Fig. 3c), with most
132 studies using methods which are likely to have strong diver-effects (potentially missing up to
133 70% of fishes) (Dickens et al. 2011; Emslie et al. 2018). Counts from these approaches,
134 therefore, focus on bold, non-diver-averse fishes.

135 Overall, it appears reef fish studies focus on fishes with bright predominantly yellow
136 hues, using field approaches that focus on relatively warm-water seasons, in attractive
137 locations (that do not represent the majority of reef area), using methods that favor bold
138 fishes. Furthermore, it should be noted that although we looked at four separate aspects, there
139 is a possibility that these factors may be operating synergistically. For example, we may
140 choose locations because they support more bold, yellow fishes or select survey methods that
141 are particularly good at censusing bold (territorial) brightly coloured fishes (cf. Emslie et al.
142 2018).

143

144 **Biases and our approach to research**

145 Biases, be they conscious or sub-conscious may be logical: in summer, reefs are
146 usually warm and calm with high fish recruitment (Meekan et al. 1993; Booth and Beretta
147 1994). The crest and slope often support the highest densities of fish and corals (Russ 1984;
148 Wismer et al. 2009), and standard fish censuses are quick and easy to conduct. Colour-based
149 biases, however, appear to have a stronger influence from subconscious biases; yellow fishes
150 are undeniably attractive to humans (as reflected by their frequent occurrence on marketing
151 images in magazines) (cf. Tribot et al. 2019), and in reef environments they stand out clearly
152 (Marshall 2000). Unfortunately, whether conscious or subconscious, such selectivity may lead
153 to a biased or partial understanding of coral reef fish ecology.

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155

156 **Why does this matter?**

157 The problem with subconscious or implicit biases are many fold, with the potential for
158 missed opportunities, partial understanding and misleading interpretations. Indeed, they may
159 lead to an over- or under-estimation of the impacts of climate change or a redirection of
160 research resources to functionally irrelevant species. We provide four examples where biases
161 may change our understanding of reef ecosystems.

162 Firstly, for example, many of the highly-studied, yellow-hued species have extremely
163 tight associations with live branching coral and may be severely impacted if corals are lost.
164 Indeed, yellow damselfishes (e.g. *Pomacentrus moluccensis*) and brightly-coloured yellow
165 and white butterflyfishes show some of the strongest declines following coral loss (Pratchett
166 et al. 2006, 2008; Wismer et al. 2019). However, losses in these species may not be
167 representative of other species or families. For example, many dark-coloured damselfishes
168 often show significant increases following bleaching events (Pratchett et al. 2008; Wismer et
169 al. 2019). Our favourite fishes may just be exceptionally sensitive.

170 Secondly, and by contrast, is the case of the gobies and their allies, collectively
171 termed cryptobenthic reef fishes; one of the most overlooked fish groups on coral reefs
172 (Brandl et al. 2018). Frequently overlooked because of their small size and cryptic behavior,
173 recent work has identified cryptobenthics as a major driver of trophodynamics on coral reefs
174 (Brandl et al. 2019c), supplying up to 70% of consumed fish flesh (Brandl et al. 2019a). Yet,

175 these fishes are strongly selected against in surveys, remaining largely invisible in visual
176 censuses (Ackerman and Bellwood 2000) due to their predominantly drab or cryptic colours
177 (Fig. 1c), and their low densities on the upper reef slope and crest (Depczynski and Bellwood
178 2005). A similar situation is seen in some off-reef plankton-feeding species, e.g. the fusiliers
179 (Caesionidae), which are often overlooked on visual censuses because they occur high off the
180 reef (Hamner et al. 1988; Russ et al. 2017). Despite being missed, they represent one of the
181 most important conduits for supplying energy to coral reefs via pelagic subsidies (Morais and
182 Bellwood 2019).

183 Third, is the example of herbivorous reef fishes, a group widely regarded as critically
184 important on coral reefs (Hughes et al. 2010; Bellwood et al. 2019; Brandl et al. 2019b). Even
185 in this group, biases have potentially shaped and/or hindered progress in this field. For
186 example, it was not until detailed video-based assessments of herbivory were conducted that
187 we were able to identify the potential importance of three drab reef fishes *Platax pinnatus*
188 (Bellwood et al. 2006), *Melichthys niger* (Tebbett et al. 2020) and *Siganus canaliculatus* (Fox
189 and Bellwood 2008) in macroalgae removal on coral reefs. Until these video-based studies
190 were performed, the former two species were not recognized as significant reef herbivores in
191 the study areas, while the later had not been recorded from the study location despite the
192 widespread use of traditional census techniques. Thus, highlighting the potential for ‘how’ we
193 conduct our studies to provide only a partial understanding of specific processes.

194 Finally, the impacts of biases associated with selective seasonal and location sampling
195 might have a particularly pronounced effect on our understanding of the process of herbivory.
196 This process can be strongly related to seasonally-variable factors such as temperature
197 (Longo et al. 2019). Indeed, evidence suggests that macroalgae removal on GBR reefs can
198 decrease by over 60% in the winter relative to summer months (Lefèvre and Bellwood 2011),
199 while algal turf consumption by herbivorous fishes in the Caribbean can decrease by over
200 20% in the winter (Van Rooij et al. 1998). In addition, while herbivorous fish densities are
201 higher per unit area on the crest and slope (Russ 1984; Wismer et al. 2009), where studies
202 generally count fishes (Fig. 3b), the reef flat is the most substantive reef habitat by total area
203 (Bellwood et al. 2018). Consequently, reef flat habitats support nearly 80% of the
204 herbivorous fish populations on reefs and account for approximately 75% of the herbivorous
205 fish biomass production on reefs (Bellwood et al. 2018). Despite this, to-date, we have
206 focused our research on a small subset of the available reef area, providing only a partial
207 view of reef-wide processes. As such, there is a clear potential to underestimate or
208 overestimate the rates of specific functions depending on ‘when’ and ‘where’ we perform our
209 studies.

210 Many of the examples above stand in marked contrast to the overwhelming attention
211 paid to damselfishes and butterflyfishes, with >35% and >11% of all studies that selected
212 species involving these families, respectively (compared to <7% and <2% of studies
213 involving the cryptobenthic gobies or blennies, respectively, and <1% involving the off-reef
214 fusiliers). Arguably, damselfishes and butterflyfishes are among the most intensely studied
215 reef fish families. However, their sensitivity to coral loss (Pratchett et al. 2006, 2008) offers a
216 stark contrast to the patterns seen in the examples where herbivores, cryptobenthics and off-
217 reef planktivores show an unexpected degree of resilience, especially in supporting
218 ecosystem processes, even in the face of coral loss (Morais and Bellwood 2019; Robinson et
219 al. 2019; Taylor et al. 2019). Unfortunately, it is fishes such as damselfishes and
220 butterflyfishes that are often used to examine the impacts of future climate change scenarios,
221 habitat degradation and predator-prey interactions; potentially biasing our view towards one
222 of high-sensitivity (cf. Clark et al. 2020). Much of the research to-date appears to be looking

223 at the ‘passengers’ rather than the ‘drivers’ of ecosystems (sensu Walker 1992); many of
224 which are evolutionary baubles on the tree of life (sensu Bellwood et al. 2017). This results in
225 a partial understanding of reef ecosystems. Previous results, therefore, are not wrong, just
226 incomplete. The most valuable step is in recognizing the potential for such oversights. Thus,
227 the selection of study species, as well as when, where and how we conduct our research, has
228 the potential to profoundly change our perception of coral reef ecosystems and associated
229 critical processes.

230 **Conclusions**

231 As coral reefs reconfigure in response to anthropogenic stressors, it is becoming
232 increasingly clear that we need to understand what keeps reefs functioning if we are to steer
233 them through the challenges they will face in the near future (Hughes et al. 2017; Bellwood et
234 al. 2019). Yet, it appears that other factors, not necessarily functional importance, may have
235 largely influenced our selection of study species, and when, where and how we have studied
236 them. These factors, potentially shaped by human preferences or biases, may have limited our
237 ability to fully understand reef functions.

238 If we are to understand high-diversity ecosystems, be they coral reefs, alpine grasslands
239 or rainforests, it is imperative to understand ourselves first. There is undoubtedly a place for
240 interest-based science looking at morphologically unique or colourful species. But in a
241 rapidly changing world where the functionality of high-diversity ecosystems is under threat, a
242 new focus on function rather than convenience or appearance may be warranted (Bellwood et
243 al. 2019). Brightly coloured fishes may be interesting, but the future of coral reefs may
244 depend on their drab counterparts that do not make it into advertisements but do keep coral
245 reefs alive.

246

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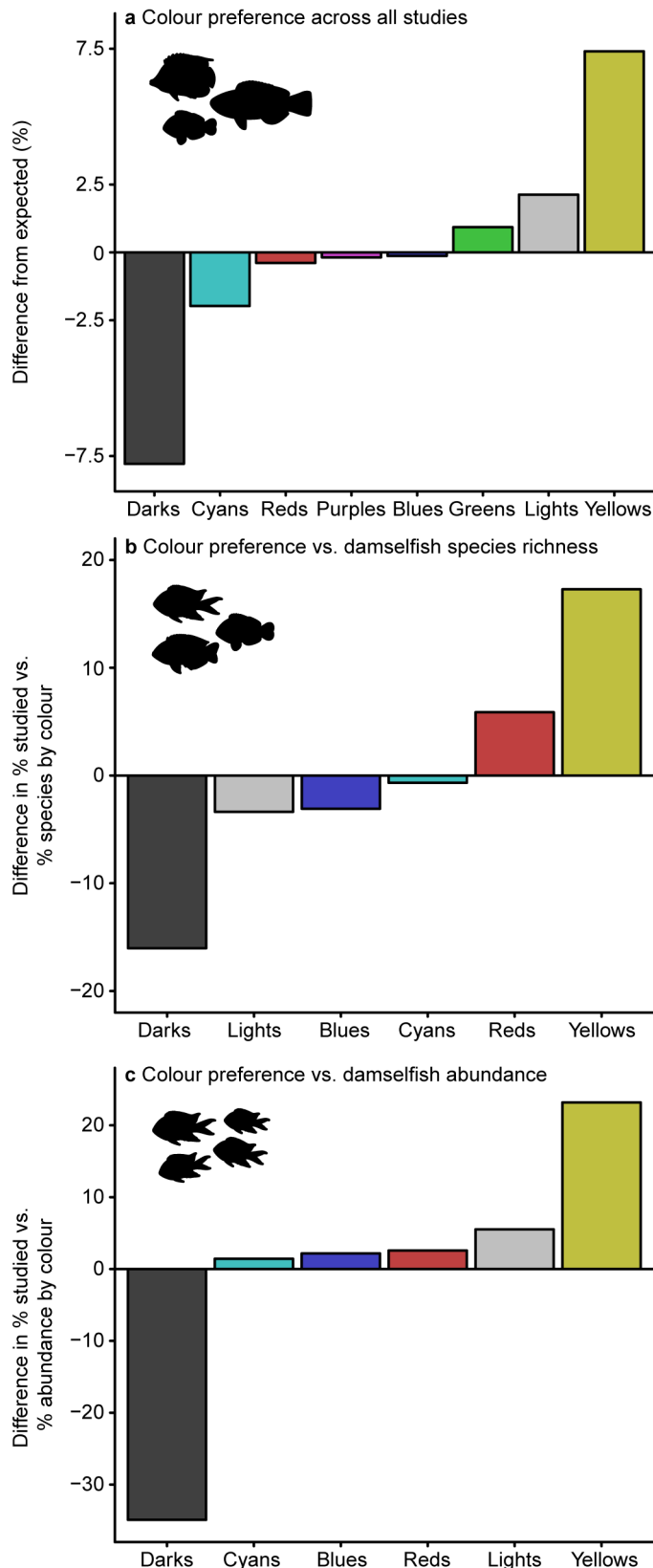
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- 365



366

367 **Figure 1** Four coral reef fish species; a) *Pomacentrus brachialis*, b) *Pomacentrus*
368 *moluccensis*, c) *Crossosalarias macrospilus* and d) *Chaetodon rafflesii*. Of these species, two
369 have characteristics that would lead to strong positive selection as study species (b, d), while
370 two (a, c) are rarely studied. (Photos. a, c Victor Huertas, b. Christopher Hemingson, d.
371 Renato Morais).

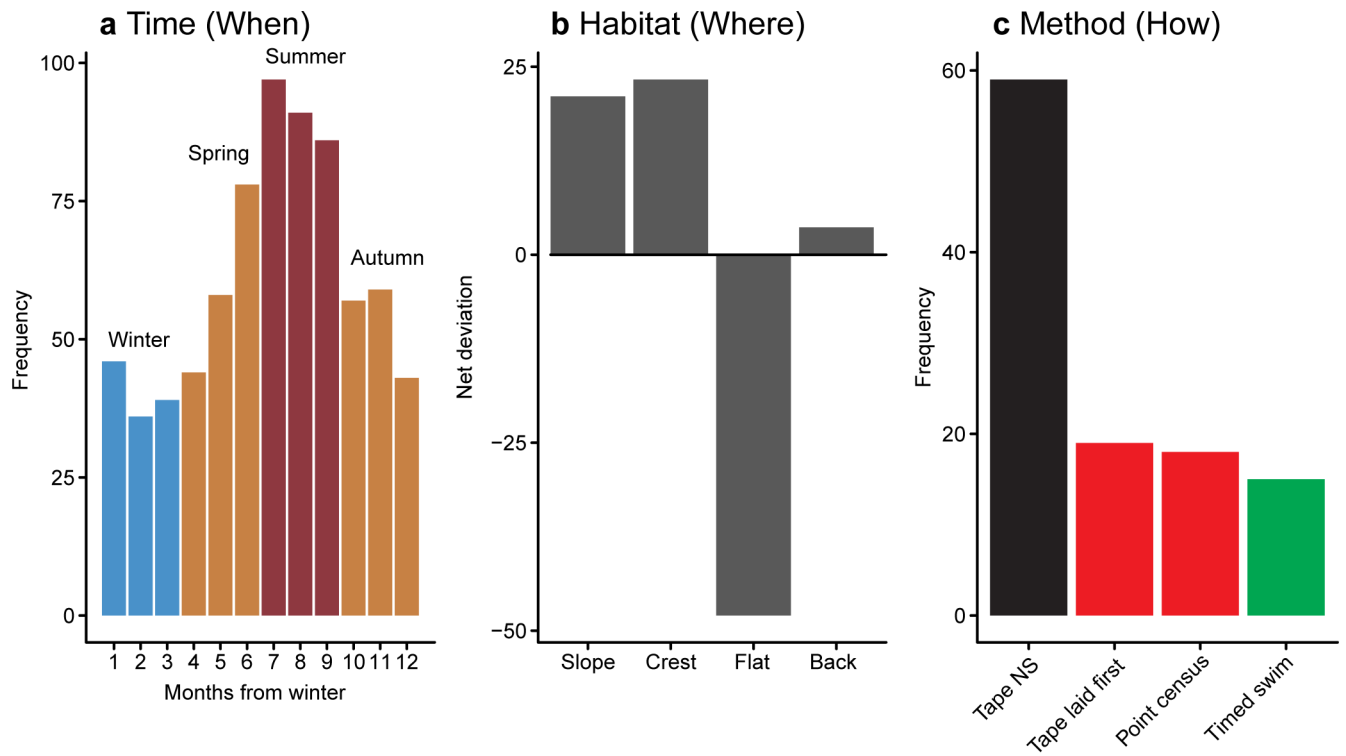


372

373 **Figure 2** Species selection in coral reef fish research (based on 37 years of research in the
 374 journal *Coral Reefs*). Whether examining species colouration relative to a) all species studied,
 375 b) all damselfish species on the GBR, or c) damselfish abundances on GBR mid-shelf reefs,
 376 the pattern was the same: research was overwhelmingly focused on yellow fishes with a
 377 strong negative selection for dark-coloured fishes.

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380

381 **Figure 3** The when, where and how of coral reef fish research. a) Frequency distribution of
382 sampling months during field-based studies (standardized for the northern and southern
383 hemisphere as months since first winter month). b) The selectivity of four major reef habitats
384 as fish census locations (deviation from expected if habitats were selected based on their area
385 covered). c) The frequency distribution of the four most common fish census methods (NS =
386 not specified). For more details please see the supporting information.

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