

14 **Highlights**

- 15 • Microplastics (MPs) have been determined in tissues of fish and crustaceans from the Musa
16 estuary and Persian Gulf
- 17 • 828 MPs of mainly a fibrous nature were detected in all tissues and species examined
- 18 • Mean abundance ranged from 7.8 in tiger prawn to 21.8 in bartail flathead
- 19 • The means by which MPs enter non-digestive tissues is unclear
- 20 • The occurrence of MPs in seafood for human consumption is cause for concern

21

22 **Abstract**

23 Commercially-important species of fish and a crustacean from four sites in the Musa estuary and
24 a site in the Persian Gulf have been analysed for the presence and location of microplastics (MPs).
25 A total of 828 MPs were detected in the guts (gastrointestinal tracts), skin, muscle, gills and liver
26 of demersal and pelagic fish (*Platycephalus indicus*, *Saurida tumbil*, *Sillago sihama*, *Cynoglossus*
27 *abbreviatus*) from all five sites and in the exoskeleton and muscle of the tiger prawn, *Penaeus*
28 *semisulcatus*, from three sites. On an individual basis, MPs were most abundant in *P. indicus*
29 (mean = 21.8) and least frequently encountered in *P. semisulcatus* (mean = 7.8), but when
30 normalized on a mass basis, MPs ranged from 0.16 g⁻¹ for *C. abbreviatus* to 1.5 g⁻¹ for *P.*
31 *semisulcatus*. Microscopic analyses (polarized light, fluorescence, SEM/EDS) revealed that MPs
32 were mainly fibrous fragments (with a few angular fragments) of various colour and size (< 100
33 µm to > 1000 µm) and with strong C and O signatures. Additional particles detected that were
34 distinctly different in colour, morphology, brittleness and elemental composition (part-metallic,
35 and containing Cu) were suspected of being fragments of antifouling paint. The means of entry of

36 MPs into tissues not involved in digestion are unclear but could be related to translocation or
37 adherence. Regardless of the mode of accumulation, the presence of MPs in heavily fished species
38 of fish and crustacean raises concerns about the potential transfer of synthetic materials into
39 humans.

40

41 **Keywords:** Microplastics; fish; prawns; accumulation; microscopy; Persian Gulf

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43

44 **1. Introduction**

45 While the effects of chemical pollutants on marine ecosystems have been studied for many
46 decades, the pervasiveness and impacts of litter on marine life have been recognized more recently
47 (Auta et al., 2017; do Sul and Costa, 2014). Plastics, as identifiable primary objects or secondary
48 fragmented pieces, comprise the largest pool of litter on both a mass and number basis and enter
49 the oceans via rivers, sewage discharge, land run-off, and spillages and discharges from ships at
50 sea (Moore, 2008; Andrady, 2011; Barnes et al., 2009; Gregory and Andrady, 2003). Given the
51 expected future demand and discharges of plastic, coupled with the resistance of synthetic
52 polymers to environmental degradation, it is clear that the marine plastic inventory will continue
53 to increase beyond at least the next decade (Jambeck et al., 2015).

54

55 Of particular concern with respect to both the direct impacts on marine life and transfer through
56 the foodchain are readily ingestible microplastics (MPs), or synthetic particles ranging from a few
57 micrometers to five millimeters in any dimension (Alomar et al., 2016; Turner, 2017; Abbasi et

58 al., 2017). Primary MPs include abrasive micro-beads in face scrubber cosmetics and toothpaste,
59 synthetic fibres and pre-production resin pellets, while secondary MPs are generated in situ by the
60 mechanical and oxidative breakdown of larger plastics (Hidago-Ruz et al., 2012). As well as the
61 inherent composition of the polymer and the presence of any additives, the chemistry of MPs may
62 be modified by the adsorption of toxic substances from ambient sea water to the hydrophobic
63 plastic surface (e.g. organic pollutants; Teuten et al. 2007; Ziccardi et al. 2016) or to more
64 hydrophilic, hydrogenous or biogenic phases coating the surface (e.g. heavy metals; Ashton et al.,
65 2010; Holmes et al., 2014).

66

67 A wide range of marine organisms, including bivalves, zooplankton, fish, invertebrates, birds and
68 cetaceans, incidentally take up MPs from sediment or the water column because they mistake them
69 for food (Cole et al. 2013; Lusher et al. 2015; Ferreira et al. 2016). Ingesting MPs of no nutritional
70 value may induce physical and chemical toxicity, block or damage the digestive tract, or decrease
71 individual fitness, ultimately resulting in death (Wright et al., 2013; Luís et al. 2015; De Sá and
72 Guilhermino 2015). Moreover, MPs of tens of micrometers in dimension have the propensity to
73 translocate from the gut to the circulatory system in many organisms where they may reside for
74 relatively long periods of time (Browne et al., 2008; van Cauwenberghe et al., 2015; Collard et al.,
75 2017). While the effects of translocated MPs on chronic animal health are unknown, their presence
76 is of particular concern because consumption of contaminated food, including fish and shellfish,
77 may act as a vehicle for the ingestion and translocation of MPs in humans (Li et al., 2015; Rist et
78 al., 2018).

79

80 The Musa is one of the biggest estuaries in the northern Persian Gulf and is the most important
81 fishery resource for people in the cities of Mashahr (population 150,000), Sarbandar (75,000) and
82 Hendijan (50,000). While the coast of the estuary is flanked by agricultural land, there are also
83 various industrial plants and extensive docks that support the petrochemical and shipping
84 industries and municipal and industrial sewage from the catchment is poorly treated (Hosseini et
85 al., 2013; Rastegari Mehr et al., 2016). Given these conflicting uses of the estuary, the aim of the
86 present study was to determine whether MPs are accumulating in different organs of five abundant
87 and commercially valuable species of fish and crustacean that are heavily consumed by local
88 people. Specifically, we target the skin, gastrointestinal tract, liver, muscle and gills of two
89 demersal fish, the bartail flathead (*Platycephalus indicus*) and greater lizardfish (*Saurida tumbil*),
90 one pelagic fish, the northern whiting (*Sillago sihama*), and one mesopelagic species fish, the
91 tongue sole (*Cynoglossus abbreviatus*), and the skin and muscle of the tiger prawn, *Penaeus*
92 *semisulcatus*.

93

94 **2. Materials and methods**

95 **2.1. Sampling and sample preparation**

96 Fish and prawn samples were caught from along the coastal waters of the Persian Gulf during June
97 2015 by a trawl net from five locations (see Figure 1), one of which served as a control site (S5;
98 the fishery port of Hendijan located outside the estuary and 70 km from any petrochemical
99 facilities). At each station, up to five samples of each species were collected, with a total catch of
100 56 specimens among all species. Samples were transported in a cooler to the laboratory where they
101 were stored at -20 °C pending processing and analysis.

102

103 Given the ubiquity of MPs in the indoor environment (Gasperi et al., 2018), suitable measures
104 were undertaken to prevent plastic and fibre contamination in the laboratory. Thus, all chemical
105 reagents were filtered (8- μ m, Whatman No. 540) before use and white cotton laboratory coats,
106 single-use latex gloves and face masks were used throughout sample manipulation and processing.
107 Working surfaces were thoroughly cleaned with ethanol and all glassware, tools and fish and
108 prawn skin surfaces were washed successively with a commercial dishwashing liquid, HPLC-
109 grade distilled water and ethanol before being dried in an oven at 105 °C (glassware and tools) or
110 at room temperature in a metal cabinet (skin surfaces). Analysis of two procedural blanks (without
111 tissues) and distilled water contained in two wide dishes that had been left exposed during the
112 duration of sample processing revealed no contamination from MPs under the working conditions
113 in the laboratory.

114

115 **2.2. Extraction of MPs**

116 As required, specimens were thawed and the fork length from the mouth to the central point of the
117 caudal fin and body weight were recorded. Each fish was gutted and dissected in a metal tray using
118 a scalpel, forceps and scissors and the muscle, skin, gills, liver and gut (gastrointestinal tract)
119 retrieved. The pooled livers, guts and gills from each species and at each site were transferred
120 directly to covered petri dishes while pooled muscles and skin, after separation, were homogenized
121 using an Electric Meat Grinder (KENWOOD MG510, UK) before about 15 g of each was retained
122 and stored in a petri dish. For the (smaller) prawns, tissue retained for analysis was restricted to
123 the muscle and skin (exoskeleton) that was pooled from individuals and homogenized as above.

124

125 Tissues of fish and prawn were subject to digestion to remove organic matter and leave behind
126 silica/aluminosilicates and any plastic (Karami et al., 2017). Thus, tissues were emptied into a
127 series of 500 mL glass beakers to which approximately 30 mL of 35% H₂O₂ (Arman Sina) and 30
128 mL of 4% KOH (Merck) were added. The contents were digested for 72 h at 60 °C in an oven to
129 dissolve the soft organic components of the tissues, before a 10:40 ml mixture of 68% HClO₄ and
130 65% HNO₃ (both Merck) was added to completely digest more resistant material like the gills and
131 skin-exoskeleton. After a few minutes of acid extraction, digests were diluted with warm distilled
132 water to preserve the integrity of MPs. Plastics were separated from all tissues with the exception
133 of the gut by shaking digests at 350 rpm for 5 min and subsequently centrifuging triplicate aliquots
134 for 5 min at 4000 rpm. Supernatants were directly filtered under vacuum through S & S grade
135 589/3 filters which were subsequently stored and dried (at room temperature) in individual petri
136 dishes pending analysis.

137

138 For MPs embedded in the gastrointestinal tract of fish, remaining digests were agitated at 350 rpm
139 for 5 min in a solution of concentrated sodium iodide (NaI, Merck; density = 1.6-1.8 g cm⁻³) to
140 separate plastics from additional material that had been ingested with subsequent filtration and
141 storage undertaken as above.

142

143 **2.3. Observation and validation of MPs**

144 A visual assessment of material retained on the filters, and including any arising from the
145 procedural control, was made according to colour, size and morphology (elongated fibre versus
146 angular fragment) and at up to 200 x magnification using a Carl-Zeiss binocular microscope. The
147 presence of plastic was verified by the colours returned by polarized light microscopy using an

148 Olympus BX41TF microscope and by fluorescence microscopy using an Olympus CX31
149 microscope. Images from all microscopic techniques were captured using an Olympus Pen EPL 1
150 digital camera.

151
152 Based on the optical microscopy results, the topography and elemental composition of selected
153 MPs were determined through high vacuum SEM/EDS. We used a Tescan VEGA 3 electron
154 microscope (with a resolution of 2 nm at 20 kV) and an Oxford Instruments X-Max 50 silicon drift
155 detector with AZtec and INCA software after samples that had been carefully brushed from the
156 filters were mounted on double-sided adhesive carbon tabs on aluminium SEM stubs.

157

158 **3. Results**

159 **3.1. Size and weight of fish and prawns**

160 Table 1 summarises the catch from each sampling site (note that the number of species caught at
161 each site varied and that some species were absent from sites 1, 2 and 3). Also shown are the mean,
162 minimum and maximum lengths and weights of each of the five species, serving to illustrate
163 differences in size among species and between sites and, for a given species, differences in age
164 and, therefore, propensity to have accumulated MPs.

165

166 **3.2. MPs in fish and prawns**

167 Table 2 shows the number of MPs in the tissues of the five species at each site, with data pooled
168 for the number of individuals indicated in Table 1. Note that MPs were detected visually (Figure
169 2), with the synthetic nature of samples confirmed by fluorescence and polarized light

170 microscopies for characteristic response to visible and ultraviolet light (Woodall et al., 2015; Wang
171 et al., 2016; Figure 3) and, for selected samples, by SEM/EDS for surface morphology and
172 elemental composition (mainly carbon). By comparison, no particles of this nature were observed
173 on the two filters arising from the procedural controls.

174
175 Among the catch, 828 pieces of MP were detected, being encountered across all tissues from each
176 species. In only isolated cases (e.g. the liver of *P. indicus* from sites 1, 2 and 5 and the gut of *P.*
177 *indicus* at site 5) were MPs absent, with numbers exceeding 25 in the skin of *S. sihama* at site 2,
178 the gills of *P. indicus* at site 4 and the skin of *P. indicus* at site 5. On this basis, there were no clear
179 differences in the total number of MPs accumulated by each species or between sites (and
180 including the control site), but numbers tended to be higher in the skin, muscle and gills than the
181 gut and liver of *S. sihama* and *P. indicus* and were always greater in the skin than in muscle from
182 *P. semisulcatus*. When considered on an individual basis, or after total numbers for each species
183 had been normalized for the number of samples analysed, MPs are most abundant in *P. indicus*
184 (mean = 21.8) and least frequently encountered in *P. semisulcatus* (mean = 7.8); when normalized
185 on a mass basis, however, the mean abundance of MPs ranged from 0.16 g⁻¹ for *C. abbreviatus* to
186 1.5 g⁻¹ for *P. semisulcatus*. By comparison, a recent study by Akhbarizadeh et al. (2018) in the
187 northeast of the Persian Gulf reports an average abundance of MPs in muscle of the fish, *P. indicus*,
188 *Sphyraena jello* and *Epinephelus voioides*, and the shrimp, *Alepes djedaba*, of 1.85 ± 0.46, 0.57 ±
189 0.17, 0.78 ± 0.22 and 0.80 ± 0.12 g⁻¹, respectively.

190
191 Nearly all MPs encountered were filamentous fragments (consisting of single fibres) of different
192 size and colour and as illustrated in Figure 2. In only five cases were non-fibrous plastics found

193 among the different species of fish: specifically, two white fragments in the muscle of *C.*
194 *abbreviatus* from site 5, one yellow fragment in the gills of *S. sihama* from site 4 and one blue
195 fragment in the gastrointestinal tract of both *S. sihama* at site 4 and *S. tumbil* at site 1.

196
197 The size distributions of MPs are shown in Table 3 for individual tissues and in Figure 4 for whole
198 organisms. Thus, there is a wide range of lengths of (mainly) filamentous material across all
199 species, with the most abundant sizes between either 100 and 250 μm (*S. sihama*, *P. indicus*, *P.*
200 *semisulcatus*) or 250 to 500 μm (*C. abbreviatus*, *S. tumbil*). With respect to the different tissue
201 types, the digestive organs appear to contain a high proportion of relatively large MPs, while
202 particles above 250 μm are absent from the liver.

203
204 The colour distribution of the MPs that had visibly accumulated is shown in Figure 5. Thus,
205 overall, 71% of MPs consisted of black or grey filamentous fragments, with blue and green
206 fragments comprising about 12% of the MP pool. White-transparent and red-pink fragments
207 contributed about 7 and 8%, respectively, with yellow-orange material lowest in overall abundance
208 at about 1.3%. There were no clear differences in colours accumulated by different species or in
209 different organs. However, there were notable differences in the distribution of certain colours
210 between the different sites; for instance, only one white-transparent fragment and no yellow-
211 orange fragments were recorded at site 1 while six yellow-orange and 20 white-transparent
212 fragments were observed at sites 4 and 5, respectively.

213
214 In addition to the MPs described above and quantified in Tables 2 and 3, a number of fragmented
215 particles of between a few tens of nm to a few hundred μm in diameter were observed in the guts

216 and gills of (mainly) pelagic fish that were distinctly different. Thus, EDS revealed the presence
217 of metals, and mainly Cu, in addition to C and O, while manipulation during analysis and SEM
218 imagery showed that the material was highly brittle (Figure 6). It is possible that these particles
219 were of metal construction, at least in part. However, given the detection of both organic material
220 and Cu, we suspect that these particles are small flakes of paint impregnated with Cu. Most
221 contemporary antifouling paint formulations employ Cu as a biocide and are generated abundantly
222 at boat maintenance and repair facilities and are also shed from boat hulls and other painted
223 maritime structures while in use (Turner, 2010).

224

225 **4. Discussion**

226 This study is one of an emerging number demonstrating the accumulation of MPs by marine
227 organisms. Of the MPs detected, and consistent with previous environmental studies, they are
228 mainly fibrous (Lusher et al., 2013; Rochman et al., 2015; Pazos et al., 2017), with sizes ranging
229 from $< 100 \mu\text{m}$ to $> 1000 \mu\text{m}$. MPs are generally larger in the gills and gastrointestinal tract than
230 in other organs because larger material can readily enter the digestive environment with relatively
231 little obstruction; the abundance of MPs in the digestive environment is also rather variable,
232 reflecting variations in the amount and type of consumed food both between individuals of the
233 same species and among different species.

234

235 Despite some planktivorous fish seeming to select MPs that are visually similar to their diet (i.e.
236 blue fragments) (Ory et al., 2017), without information on the colour distribution of MPs in the
237 water column or sediments of the Musa estuary and Persian Gulf there is no evidence in the present
238 study for the preferential ingestion or accumulation of MPs according to appearance. We also do

239 not have specific information on the type of plastics found in the organisms sampled, although
240 MPs retrieved from littoral sediments of the Persian Gulf indicate a predominance of polyethylene,
241 nylon and polyethylene terephthalate (Naji et al., 2017).

242
243 On an individual basis, MP abundance ranges from about 8 for the prawn, *P. semisulcatus*, to over
244 20 for the demersal fish, *P. indicus*, that forages in the sediment and where most of the denser MPs
245 reside. These values are higher than those reported for fish in previous studies; for example, up to
246 7.2 items per individual were observed in coastal and freshwater fish from China (Jabeen et al.,
247 2017), up to about 4 per individual were detected in the semi-pelagic Mediterranean fish, *Boops*
248 *boops* (L.). (Nadal et al., 2016), and an average of 1.6 items per fish were recorded in various
249 demersal fish in Spanish coastal waters (Bellas et al., 2016). However, it is important to appreciate
250 that these studies focused on the retrieval of MPs from the digestive tract only. When our data are
251 restricted to the gut, the average number of MPs per individual ranges from about 1.5 in *S. sihama*
252 to 3 in *C. abbreviatus* (see Table 2).

253
254 The discrepancies referred to above arise from the general assumption that accumulation of plastics
255 by fish and other organisms proceeds mainly through ingestion and is, therefore, dependent on
256 factors like feeding strategy and gut structure as well as the extent of local plastic pollution (Romeo
257 et al., 2015; Jabeen et al., 2017). Thus, MPs may be accumulated directly and incidentally or
258 deliberately while feeding from the water column or sifting through contaminated sediment, or
259 indirectly through the consumption of contaminated prey (Cannon et al., 2016; Jovanović, 2017).
260 The detection of MPs in the present study in organs not directly involved with ingestion-digestion

261 suggests that other factors may be significant for the accumulation and, potentially, translocation
262 of MPs in fish.

263
264 Results of laboratory experiments have reported the occurrence of MPs in the circulatory system
265 or non-digestive organs of marine invertebrates (Browne et al., 2008; von Moos et al., 2012) and
266 in the liver of zebrafish (Lu et al., 2016). However, particles employed in these studies were on
267 the order of tens of micrometers in diameter or less, thereby facilitating passage across the gill or
268 gut epithelium through cell internalization and subsequent translocation. Collard et al. (2017)
269 suggest that detection of larger MPs (and of dimensions comparable to those observed here) in the
270 livers of European anchovies (*Engraulis encrasicolus*) may result from two processes: the
271 agglomeration of smaller particles and/or passage through the gut barrier by some form of
272 intracellular or paracellular endocytosis. The former mechanism is unlikely in the present study
273 because SEM images revealed distinct and relatively smooth fibrous fragments, and without
274 knowledge of the locations of MPs in (homogenized) tissue the latter mechanism cannot be fully
275 explained.

276
277 Alternatively, it has recently been suggested that adherence affords an additional means by which
278 fibrous MPs may associate with organs independent of the digestive system, in a manner by which
279 seaweeds accumulate plastics (Gutow et al., 2016). Thus, under laboratory conditions, about 50%
280 of microfibrils exceeding 100 µm in marine mussels could be accounted for through adherence,
281 with surface area and “stickiness” two important controls in this respect (Kolandhasamy et al.,
282 2018). Regardless of the mechanisms by which MPs enter or associate with non-digestive tissues,
283 their occurrence has a number of implications for evaluating the inventory, location and toxicity

284 of MPs in marine animals, as well as for human health through seafood consumption. Specifically,
285 if the gut is considered as the sole receptacle, where MPs may either be in transit or entrapped, the
286 total number of MPs accumulated by an individual may be considerably underestimated. With
287 respect to toxicity, accumulation outside the digestive tract may induce histological changes and
288 oxidative stress (Lu et al., 2016) or release contaminants associated with or adsorbed to MPs
289 (Ashton et al., 2010). The potential for MPs to be transferred to humans should not be
290 underestimated given that the soft tissue of the species considered are important to the regional
291 fishing industry. According to the Institute of Standards and Industrial Research of Iran in 2010,
292 daily average fish muscle consumption is about 7 g/person/day, meaning that about 5 MPs could
293 be consumed on a daily basis. While there is currently no regulatory framework concerning the
294 presence of MPs in sea food (European Food Safety Authority 2016), this does not exclude the
295 possibility that MPs are able to interact with human cells and tissues and facilitate the delivery of
296 harmful contaminants to the bloodstream (Santillo et al., 2017).

297

298 **5. Conclusions**

299 This study has demonstrated the presence of MPs of mainly a fibrous nature and of length < 100
300 μm to > 1000 μm in various commercially important species of fish and a crustacean collected
301 from the Musa estuary and the Persian Gulf. Average quantities of MPs ranged from 0.16 g^{-1} for
302 the mesopelagic fish, *C. abbreviatus*, to 1.5 g^{-1} for the prawn, *P. semisulcatus*, with particles
303 encountered in various tissues from both digestive and non-digestive organs across all species. The
304 occurrence of MPs outside the digestive system suggests that material can be translocated
305 following ingestion or that additional, non-ingestive mechanisms (e.g. adherence) are significant.

306 The presence of MPs in non-digestive organs has the potential to induce toxic effects on
307 individuals and affords an exposure route to humans who consume contaminated fish.

308

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312

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457 **Figure captions:**

458 **Figure 1:** Locations of the five sampling sites along the coast of the Musa estuary.

459 **Figure 2:** Examples of MPs encountered in fish and prawn tissues and as captured by binocular
460 microscope. Note that fibres in panels (b) and (g) are extremely thin and, therefore, have a
461 relatively high propensity to penetrate tissue, and that fibres in panels (c), (d), (f), (h), (i) and (j)
462 exhibit partial entrapment in half-digested tissues.

463 **Figure 3:** An image and the composition of a fibre obtained by SEM/EDS (W% = weight percent
464 and A% = atomic percent) (a); fibre images obtained using upper-light fluorescence microscopy
465 (b,c); fibre images obtained by polarized downward projecting light microscopy (e,g) and
466 corresponding images obtained without polarized light (d,f).

467 **Figure 4:** The net distribution of MPs among different size categories (in μm) in the five species.

468 **Figure 5:** Overall colour distribution of the MPs observed in the samples.

469 **Figure 6:** SEM/EDS image and composition of a particle of a relatively brittle and non-fibrous
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472 **Table 1:** Number of species caught from each site (*n*) together with the mean (and minimum and
 473 maximum) lengths (cm) and weights (g).

		<i>S. sihama</i>	<i>P. indicus</i>	<i>C. abbreviatus</i>	<i>S. tumbil</i>	<i>p. semisulcatus</i>
S1	<i>n</i>	4	3	4		
	length	20.1 (17.2-20.1)	17.3(16.5-18.5)	17.7 (14.2-20.5)		
	weight	67.8 (49.3-95.1)	23.8 (18.3-32.7)	33.1 (14.2-56.6)		
S2	<i>n</i>	4	1	4		
	length	16.6 (13.0-20.0)	16.0	17.7 (14.2-20.5)		
	weight	39.4 (14.2-62.4)	16.8	33.1 (14.2-56.6)		
S3	<i>n</i>					5
	length					7.8 (5.5-10.0)
	weight					5.4 (2.3-10.6)
S4	<i>n</i>	4	4	3	4	3
	length	16.6 (15.5-18.0)	19.5 (18.5-21.5)	23.7 (23.0-24.0)	15.7 (13.0-18.0)	7.3 (6.0-8.5)
	weight	45.6 (36.4-53.3)	46.7 (35.7-63.6)	115.9 (109.3-123.0)	36.1 (18.6-50.4)	5.2 (2.5-8.0)
S5	<i>n</i>	5	4	4		4
	length	20.5 (18.5-24.5)	20.5 (20.0-22.0)	23.8 (22.5-26.0)		7.6 (4.5-10.5)
	weight	72.2 (51.8-119.9)	41.7 (39.1-46.5)	88.4 (75.2-115.4)		4.9 (1.4-8.7)
total	<i>n</i>	17	12	15	4	12
	length	18.6 (13.0-24.5)	19.0 (16.0-22.0)	24.6 (14.2-21.7)	15.7 (13.0-18.0)	7.6 (4.5-10.5)
	weight	57.2 (14.2-119.9)	36.8 (16.8-63.6)	75.8 (14.2-123.0)	36.1 (18.6-50.4)	5.2 (1.4-10.6)

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480 **Table 2:** Number of MPs detected in the five species pooled from each site (with the number of
 481 species given in Table 1). Also shown is the total number of MPs in each species, the mean number
 482 when normalized for the number of individuals analysed and the average mass of individuals, and
 483 the mean number per individual when only the gut was considered.

		<i>S. sihama</i>	<i>P. indicus</i>	<i>C. abbreviatus</i>	<i>S. tumbil</i>	<i>P. semisulcatus</i>
S1	skin	7	27			
	muscle	14	7			
	gut	1	11			
	gills	15	22			
	liver	6	0			
S2	skin	29	14	8		
	muscle	20	21	10		
	gut	9	4	11		
	gills	12	12	12		
	liver	4	0	5		
S3	skin					23
	muscle					12
	gut					
	gills					
	liver					
S4	skin	14	14	8	6	21
	muscle	19	14	12	12	14
	gut	12	12	18	11	
	gills	20	27	13	8	
	liver	11	13	24	17	
S5	skin	11	27	13		14
	muscle	11	13	12		10
	gut	4	0	15		
	gills	8	23	8		
	liver	12	0	11		
total		239	261	180	54	94
mean/individual		14.1	21.8	12.0	13.5	7.8
mean/g		0.25	0.59	0.16	0.37	1.51
mean/gut		1.5	2.3	2.9	2.8	

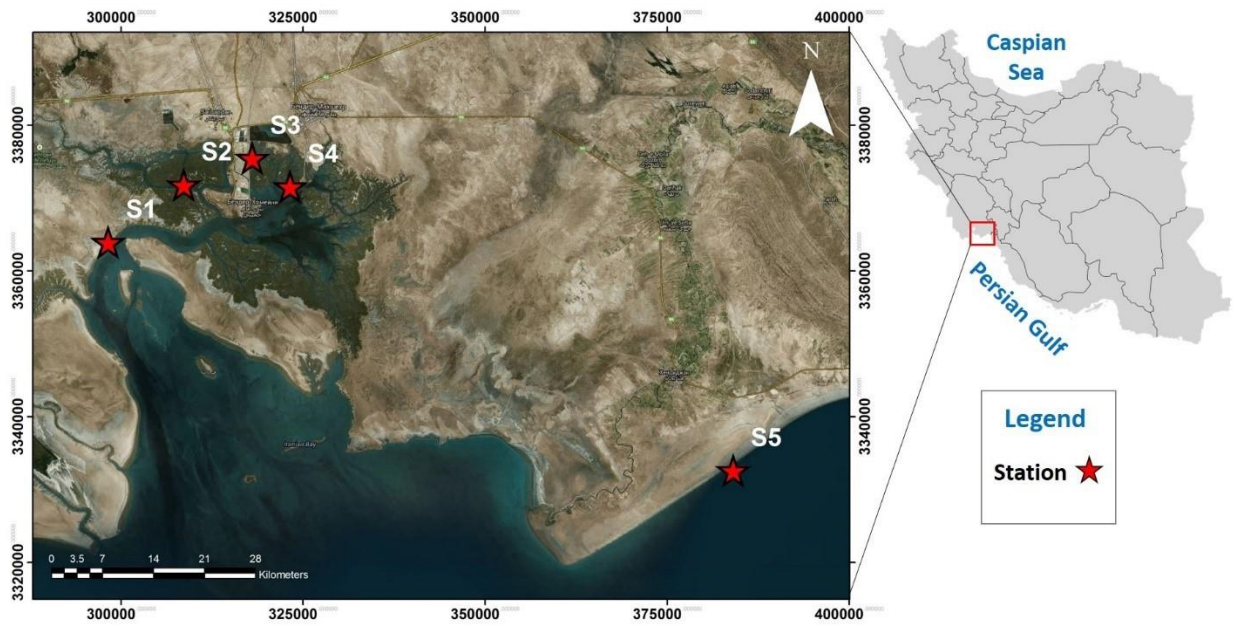
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487 Fig 1.

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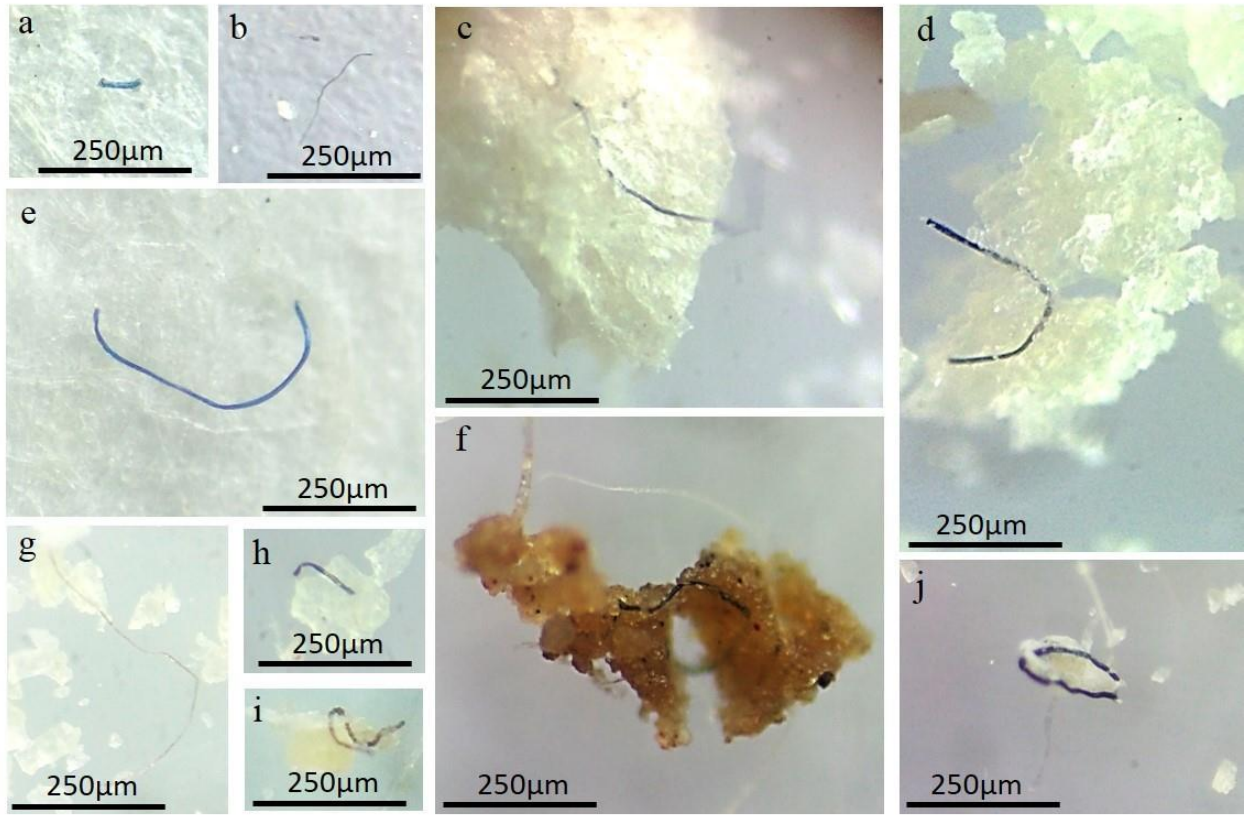
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495 Fig 2

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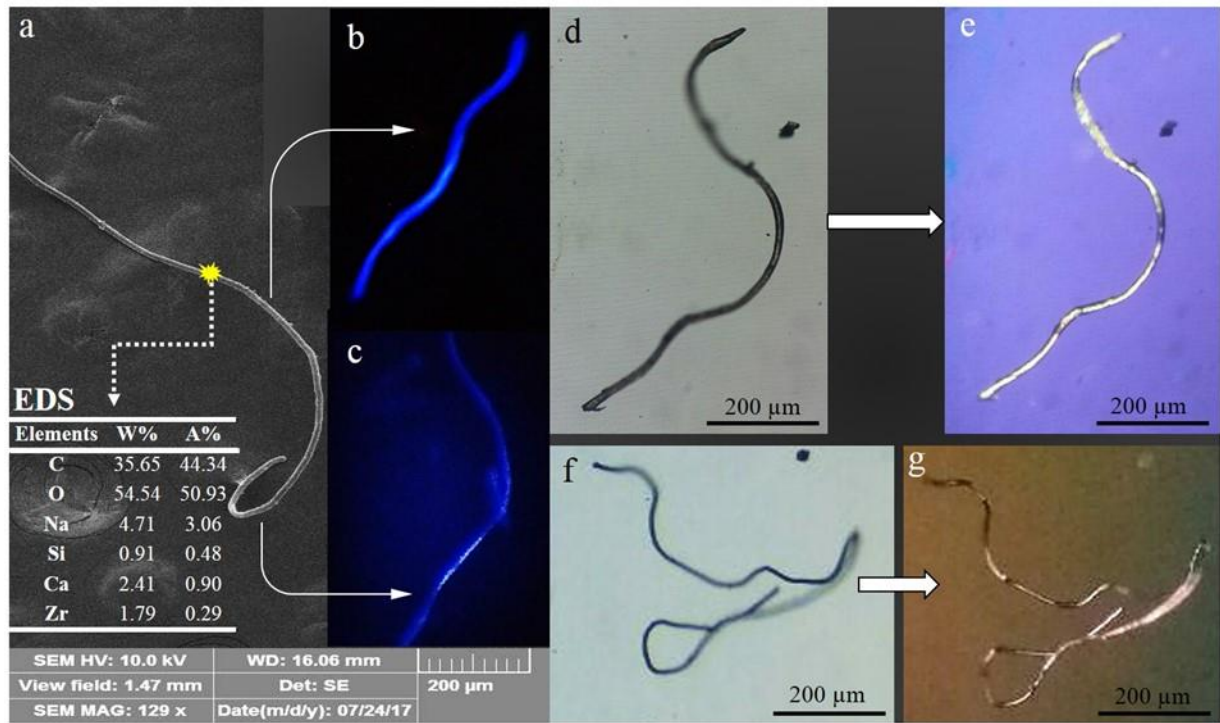


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499 Fig 3

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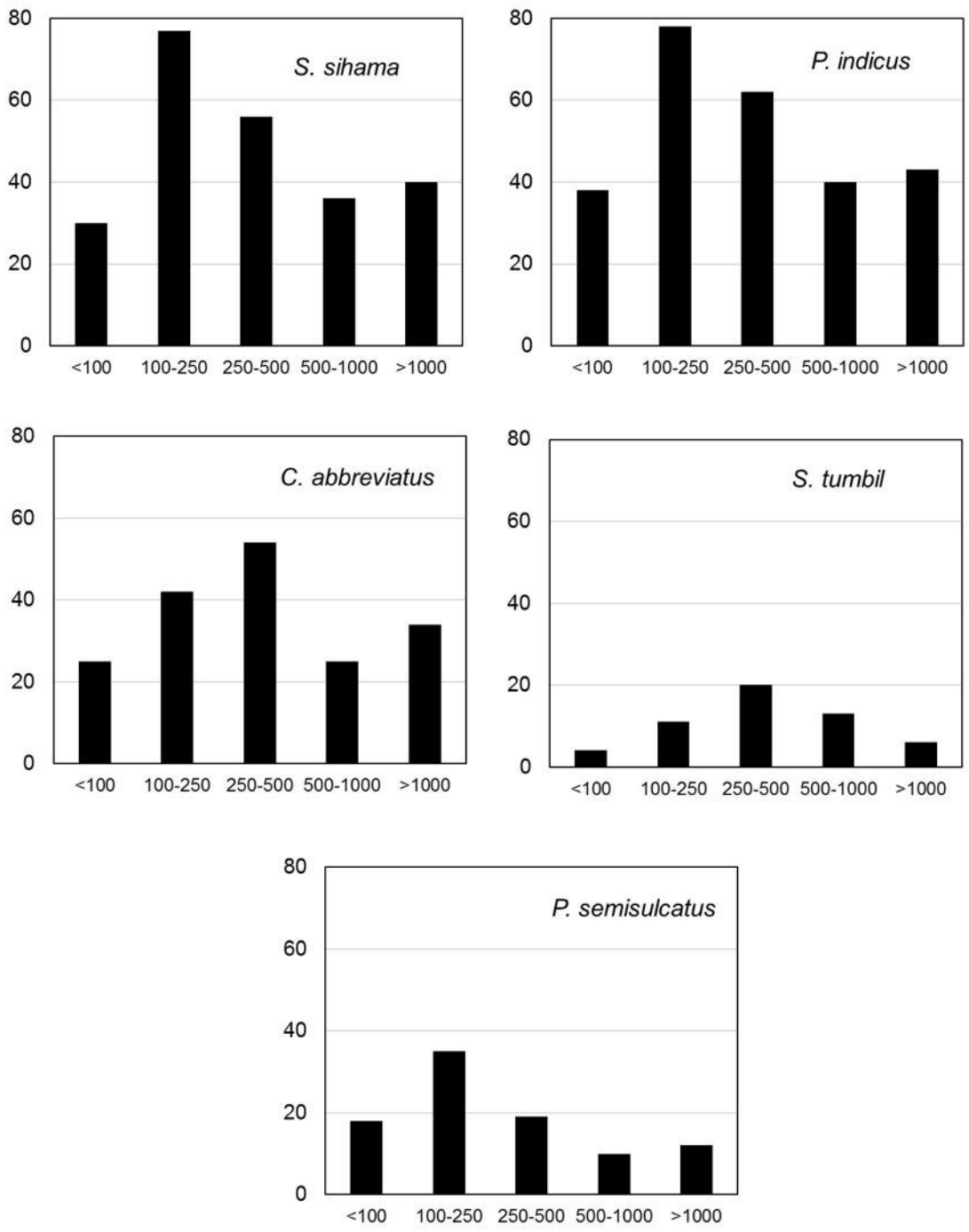
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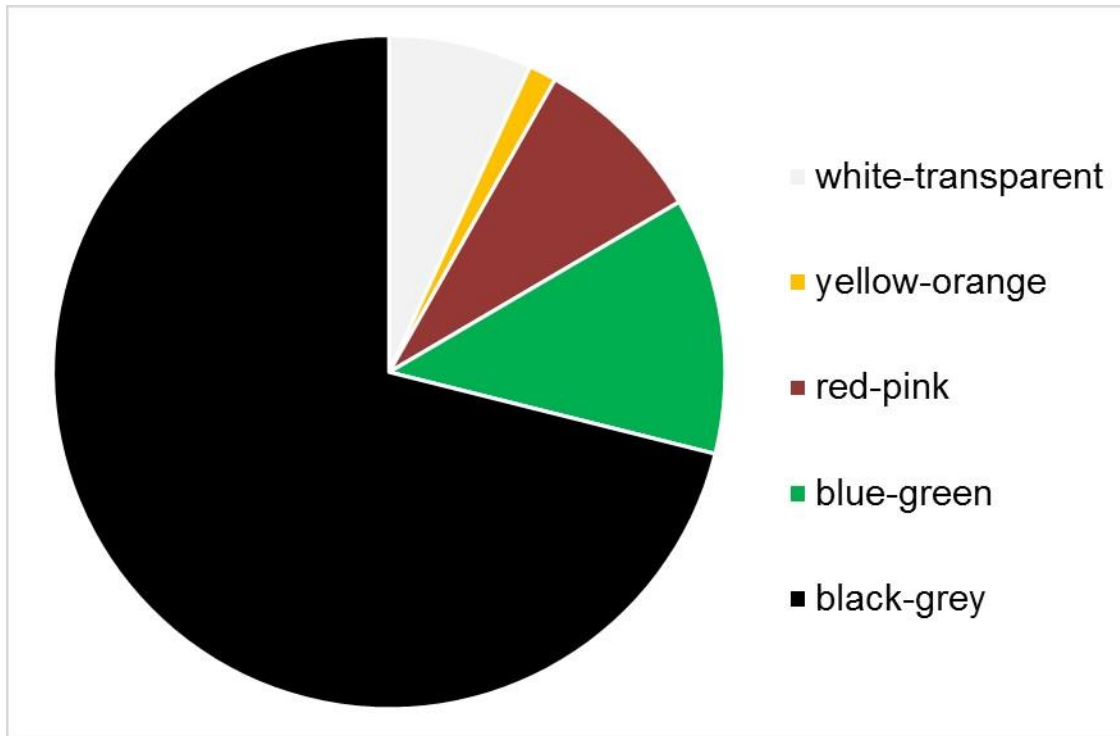


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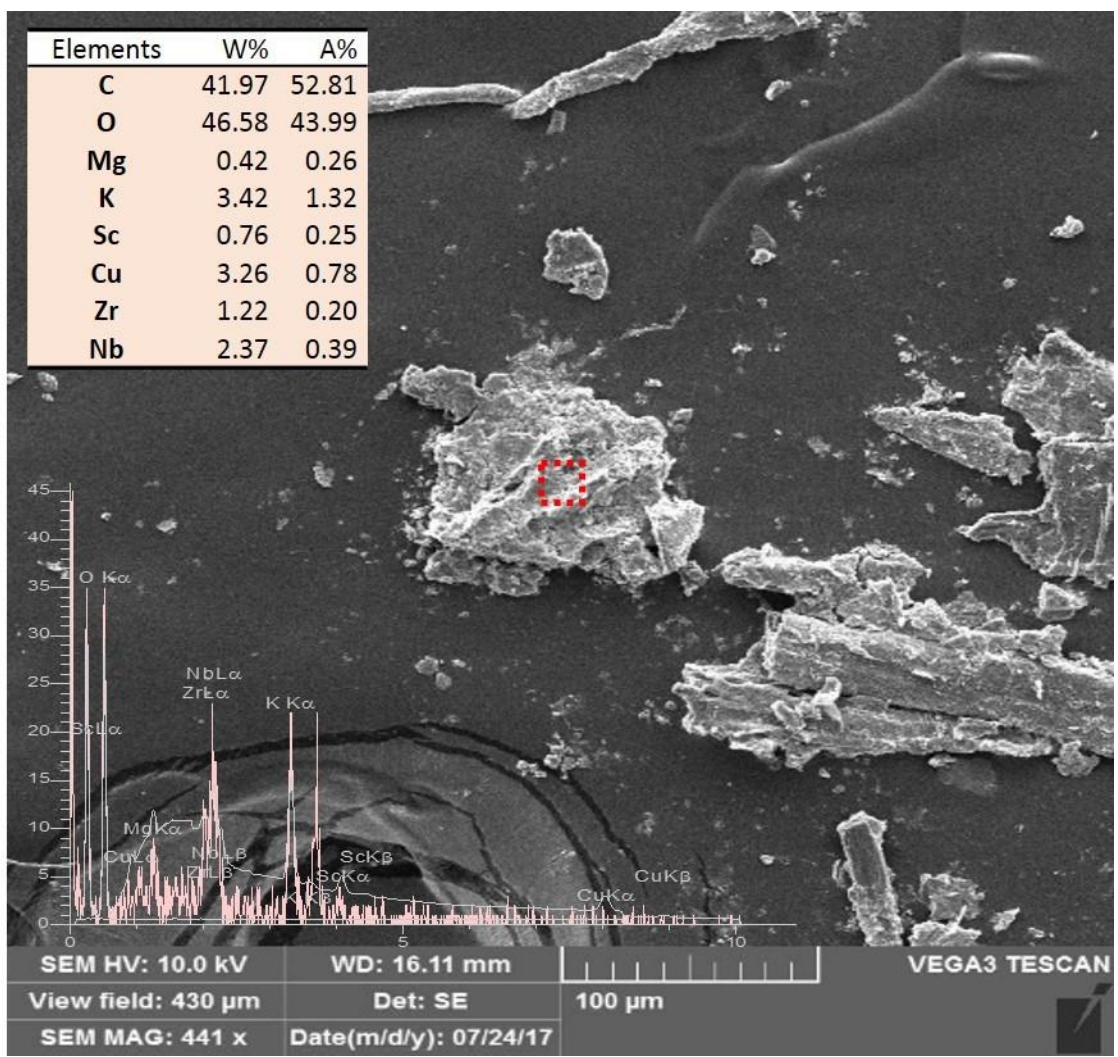
512 Fig 5



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515 Fig 6



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519 re (W% = weight percent and A% = atomic percent).