

1 **Long-term sublethal exposure to polyethylene and tire wear**
2 **particles: Effects on risk-taking behaviour in invasive and native fish**

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16 **Abstract**

17 Anthropogenic polymer particulates pollute even the most remote ecosystems and may
18 compromise the behaviour and movement skills of organisms. It is expected that invasive species
19 cope better with pollution than native species (i.e., pollution resistance hypothesis). In this study,
20 invasive gibel carp (*Carassius gibelio*) and native crucian carp (*Carassius carassius*) were used as
21 model organisms. Specimens were fed daily with food pellets (1% body weight) added with 0.1%
22 polyethylene (PE), TWPs and control. Their behavioural parameters were compared before and
23 after 14 and 60 days of exposure. Additionally, we evaluated burst swimming capacity after 60
24 days of exposure to the treatments. The fishes exposed to the PE and TWPs treatments showed
25 significant trends toward increased boldness scores and, in the PE treatment, higher utilization of
26 the open field, and both behavioural changes are associated with higher risk-taking. Invasive gibel
27 carp had substantially better swimming performance than crucian carp, but the expected trend in
28 relation to the treatments was not found. Fish exposed to sublethal doses of PE and TWPs showed
29 signs of behavioural changes after two months of exposure that may affect risk-taking behaviour,
30 which might impact species interactions with predators.

31 **Keywords**

32 habitat degradation; pollution; invasive species; behavioural changes; invasion ecology

33 **1. Introduction**

34 Plastic production worldwide has resulted in the generation of over 8 billion tons of waste; over
35 80% of it is currently dispersed across all environmental matrices (Geyer et al., 2017). Plastic
36 waste ends up in water bodies such as lakes and oceans, with rivers playing a major role in its
37 transport and transfer across different habitats (Cau et al., 2022; McNeish et al., 2018; Palmas et

38 al., 2022). The scientific community has paid particular attention to plastics that deteriorate and
39 fragment into smaller particles, generating microplastics (MPs): plastic particles with dimensions
40 comprised between 1 μm and 5 mm (Frias and Nash, 2019). Generally, MPs in rivers originate
41 from anthropogenic land-based sources due to industrial activities and population densities (Birch
42 et al., 2020). Microplastics can be found in soil, atmosphere, marine and freshwater sediments,
43 surface or ground water and biota (Boyle and Örmeci, 2020; Hidalgo-Ruz et al., 2012; Lee et al.,
44 2013; Liu et al., 2021; Robin et al., 2020). Microplastics are ingested by plethora of aquatic
45 organisms and may cause mechanical damage to tissues but also physiological or behavioural
46 responses such as immune system disturbances, oxidative stress, developmental defects, growth
47 suppression and abnormal feeding selectivity (Au et al., 2015; Caccamo et al., 2016; Cau et al.,
48 2023, 2020; de Sá et al., 2015; Espinosa et al., 2019; Rist et al., 2016; Zhang et al., 2022).

49 Among conventional plastics that dominate the world market, polyethylene (PE) has a
50 major role. PE has experienced widespread use since the 1950s in packaging products with usually
51 short shelf lives, such as plastic bags, bottles, cups and containers. Thus, it is not surprising that it
52 is one of the most recurrent types of MPs retrieved in aquatic environments, regardless of the
53 matrix of dominium explored (Kumar et al., 2021).

54 Tire wear particles (TWPs) are generated from the mechanical abrasion of tire material
55 during use on roads and have gained considerable attention as part of MPs that contaminate aquatic
56 environments (Wagner et al., 2018). Tire wear particles are dispersed in aquatic environments
57 through various pathways, such as wastewater, road water runoff, and atmospheric deposition
58 (Kukutschová et al., 2011; Sugiura et al., 2021; Ziajahromi et al., 2020). It is estimated that
59 approximately 500,000 tonnes of TWPs are generated annually in the European Union (EU) alone,
60 and 50–140,000 tonnes are released annually in EU surface waters (Hann et al., 2018; Page et al.,

61 2022). TWPs are made of a complex mixture of rubber, e.g., styrene butadiene, embedded asphalt,
62 pavement minerals and other minerals (copper and zinc) (Eisentraut et al., 2018; Panko et al.,
63 2013). Till date, specific chemical components of TWP have been the main focus for ecotoxicity
64 studies, while studies that incorporate the physical components of the contaminant are quite rare
65 and should be prioritized (Wagner et al., 2018). TWPs can also include metals (e.g., copper) that
66 can affect both biological (i.e., reduced growth) and ecological (i.e., prey–predator interactions,
67 behavioural patterns, swimming performance) traits (Gosavi et al., 2020; Siddiqui et al., 2022).

68 While great effort has been made over the last years to document the contamination by
69 MPs in aquatic environments, knowledge on the effects that exposure to specific polymers might
70 have on aquatic organisms is still scarce (Bucci et al., 2020), even in freshwater environments,
71 where PE and TWPs are more prone to be collected and transferred (Cunningham et al., 2022;
72 Wagner et al., 2018). Behavioural changes in response to exposure to pollutants are becoming an
73 increasingly important topic in ecotoxicology (Oulton et al., 2014). As a general pattern,
74 behavioural changes may occur even with small quantities of pollutants (Bae and Park, 2014), as
75 already documented in invertebrates (Van Colen et al., 2020), amphibians (Araújo and Malafaia,
76 2020) and fish (Critchell and Hoogenboom, 2018). Despite the fact that changes in animal health
77 may not be noticeable, behavioural changes may have severe consequences on animal interactions
78 within the community (Scott and Sloman, 2004), thus allowing us to identify and infer sub-
79 individual effects that eventually can alter population or even community dynamics due to their
80 adaptive and maladaptive responses to the pollution in the wild and their resistance to the pollutant
81 due to multiple stressors along the biological hierarchy (Ashauer and Jager, 2018; Sih et al., 2011).

82 As a potential cumulative effect of MP contamination in aquatic environments, the
83 "pollution resistance hypothesis" postulates that invasive species may have a competitive

84 advantage in environments that are polluted or contaminated with toxic substances (Crooks et al.,
85 2011). The hypothesis proposes that invasive species have evolved traits that enable them to
86 tolerate or detoxify pollutants, giving them a competitive edge over native species that may be
87 more susceptible to pollution (Crooks et al., 2011; El Haj et al., 2019; Varó et al., 2015). This may
88 be due to the nature of invasive species establishment, where many more species are translocated
89 and only a handful become successful and invasive (Blackburn et al., 2011; Sakai et al., 2001). As
90 invasions often start near urban areas with high levels of pollution and environmental change (Qiao
91 et al., 2022), this can act as a selective mechanism for pollution-resistant invasive species
92 (Camacho-Cervantes and Wong, 2023). Hence, to address this hypothesis with declining native
93 species and corresponding invasive species, we used the native crucian carp (*Carassius carassius*
94 L., 1758), a sharply declining species in European waters due to invasion by gibel carp (*Carassius*
95 *gibelio* K., 1782) (Jeffries et al., 2016; Kottelat and Freyhof, 2007). The invasive gibel carp was
96 imported into Romania in the mid-20th century (Szalay, 1954; Tóth, 1976) and has since spread
97 across the European continent (Kottelat and Freyhof, 2007; Perdikaris et al., 2012; Ribeiro et al.,
98 2015; Wouters et al., 2012). During gibel carp invasion, most crucian carp populations were
99 extirpated due to the higher competitive ability of the invasive carp (Tapkir et al., 2022).

100 With these premises, the present study compares the behavioural and performance responses
101 of two *Carassius* species exposed to 0.1% mass of PE and TWPs. We conducted manipulative
102 experiments to test as to whether exposure could i) alter fish behaviour either towards more
103 cautious or bolder attitude; ii) reduce fish swimming capacity, thus potentially affecting predator–
104 prey relationships and iii) differ in these effects among the native and the invasive species.

105

106 **2. Materials and Methods**

107

108 *2.1. Preparation of microplastics*

109 To realistically simulate environmental scenarios, manipulative experiments described in the
110 present work were conducted using particles produced from discarded commercial tires and plastic
111 bottles. For both polymers, toxicity levels are known to vary across different stages of weathering
112 (Arp et al., 2021; Halle et al., 2021); thus, in the present study we used end-life tires, that do show
113 a different toxicity compared to new tires, and bottles retrieved from the environment. Both items
114 were retrieved from aquatic environments, during river clean-up for macrolitter. This was done to
115 ensure that tires and plastics underwent not only typical usage but also environmental weathering
116 (i.e., photo-, thermo- and mechanical deterioration). In detail, PE-based MPs were produced from
117 a single PE item collected during the abovementioned activities and analytically characterized via
118 micro Fourier transform infrared spectroscopy (μ FT-IR) (spectra available in Supplementary
119 Figure S1). Scanning electron microscopy (SEM) was used to produce detailed images and
120 evaluate morphological features of TWPs (Supplementary Figure.S2), for their shape to be as
121 similar as possible similar to those produced by the friction of tires on asphalt Knight et al.(2020);
122 Kreider et al.(2010)The elemental analysis of the particles was carried out by X-ray Energy
123 Dispersive Spectrometer of SEM (Supplementary Figure S3). The elemental composition TWPs
124 observed on the surface of the particles (Supplementary table T2) are used in tire manufacture
125 Yang et al.(2022). The organic constituents of the TWPs were analysed with a micro-furnace
126 pyrolyzer frontier coupled to gas chromatography combustion isotope ratio mass spectrometry
127 (GC/IRMS), the selected markers used and the compounds detected are provided in
128 (Supplementary Table T1).

129 Effects on biota can be caused by physical interactions between particles and organisms, according
130 to particle size and shape, and by associated compounds released from the particles (Skjolding et
131 al., 2016). Thus, it is crucial to create experimental designs that allow scientists to properly
132 discriminate either effects caused by physical factors (size, volume and shape of particles) or from
133 chemical factors such (e.g., different polymers, production additives, etc.), within the same level
134 of the biological hierarchy (Bucci et al., 2020). To meet this need, particles of both polymers were
135 manually grated to create irregularly shaped particles (especially for TWPs, that derive from tires
136 abrasion with asphalt) and then sieved to an expected size range comprised between 70 and 210
137 μm (estimated relative abundance within this size range is reported in Supplementary Table X).
138 This size range was considered relevant for the purpose of the experiment since it simultaneously
139 encompasses the most common shape and size range of runoff and/or shredded tires (Charters et
140 al., 2015).

141

142

143 *2.2. Diet*

144 The feeds were manufactured in the laboratory at the Institute of Aquaculture and Protection of
145 Waters, University of South Bohemia. A hammer mill grinder (Mistral 50 L, River System SRL,
146 Italy) with an 18-mm mesh was used to grind commercial feed (C-3 Carpe F, Skretting, Stavanger,
147 Norway). A portion of the ground feed (1000 g) and PE (1 g) or TWPs (1 g) were thoroughly
148 mixed in a container to achieve homogeneity and the experimental groups 0.1% TWPs and 0.1%
149 PE. Subsequently, 0.2 L of warm water (40 °C) was gradually added and thoroughly mixed into
150 the feed. Water was added until the mixture become adhesive. Then, the mixture was passed
151 through the mixer, mini pelleting machine (2-mm diameter plate, Bottene, Bottene Fratelli Snc,

152 Italy) and cutter (Cutter LT3, FAC srl, Italy) to form pellets. The pellets were then placed in paper-
153 lined trays and dried in an air oven for 24 h at 55 °C. The oven was shut off after the feeds were
154 dried and they were left to cool to room temperature (20 °C). Feeds were stored at -20 °C in plastic
155 bags until fed to the fish. The control diet contained 0% plastic, and it was prepared using the same
156 procedure to achieve similar surface textures and to avoid any unforeseen interferences.

157

158 2.3. Fish collection and maintenance

159 Live crucian carps (n=60) with an average size of 93.5 ± 6.5 mm (range 81–108 mm) were
160 collected from Pavlov u Herálce (49.5039411N, 15.4285000E), and live invasive gibel carp (n=60)
161 with an average size of 91 ± 5.7 mm (range 81–105 mm) were collected from the same site. To
162 prevent mortality, the fish were transported to the laboratory in plastic barrels filled with
163 dechlorinated oxygenated water. Fish were kept in glass aquariums (64 x 60 x 45 cm) filled with
164 130 L of clean, oxygenated 20 °C water in the laboratory in groups of ten fish. The photoperiod
165 was 12 h light (9:00 a.m. light on) and 12 h dark (9:00 p.m. light off). The aquariums were fitted
166 with aerators for continuous oxygenation, and water from all the aquariums was changed every 14
167 days. Fish were allowed to acclimatize for 30 days prior to the experiment.

168

169 2.4. Experimental design A total of 120 fish; 60 (*Carassius Carassius*), 60 (*Carassius gibelio*) with
170 similar body weights were distributed into 12 experimental tanks (n=10 fish per tank). The native
171 and invasive species were kept separately in three treatments: i) control treatment, ii) PE treatment,
172 and iii) TWPs treatment, using 20 fish per treatment for each species (Supplementary Fig S.4).
173 The fish were anaesthetised with MS-222 before tagging, and their standard length (in mm) and
174 weight (in g) were recorded. During tagging, 3-4 scales were removed, and a 2-3 mm vertical

175 incision was made 3 cm posterior to the pelvic fin (Šmejkal et al., 2019). Then, a passive integrated
176 transponder tag (PIT tag, Oregon RFID, Oregon, USA; half-duplex; length: 12 mm; diameter: 2.12
177 mm; weight: 0.1 g; ISO 11784/11785 compatible) was inserted into the body cavity. The fish were
178 given a healing time of 30 days before starting the experiment. The fish were exposed for 14 and
179 60 days to the three different treatment diets (control, PE, and TWPs) and were fed daily at 1% of
180 body weight supplied once a day. Fish were monitored throughout the experiment for any potential
181 indications of poor health status (i.e., feeding behaviour, swimming activity and condition of fins).
182 Physical and chemical water parameters (pH and dissolved oxygen mg L⁻¹) were measured on a
183 weekly basis using a water quality sensor (Digisens-OPTOD, Ponsel, France) and a computer
184 running CalSens 1.4 software (Aqualabo, France).

185

186 *2.5. Fish behaviour assays*

187 *2.5.1 Boldness – emergence time*

188 The scoring behaviour was performed in a rectangular arena (57 cm long × 57 cm wide) filled with
189 dechlorinated tap water. A start box was placed at one end of the arena, and an air stone was placed
190 for aeration. The start box had a sliding door on one side operated by a pulley (Supplementary
191 Figure S5). The body weight, standard length and total length of each fish were measured one day
192 prior to the experiment. The fish were fed 24 h before the experiment. The test fish were removed
193 from the fish tanks using a hand net and placed into a start box. Using a pulley, the adjustable
194 doors were maintained closed to keep the fish out of the test room. The fish were allowed to
195 acclimatize for 10 min, and later, with the help of a pulley, the arena door was gently opened,
196 allowing the fish to emerge in the test arena. The videos were recorded for 20 min after opening
197 the arena door. To minimize the effects of human observers on fish behaviour, a webcam (Logitech

198 C270, Laussane, Switzerland) was positioned directly above the apparatus and utilized to record
199 videos. Videos were recorded for fish emergence into the test arena, and the emergence time was
200 noted when the complete fish (i.e., fish fully visible in the arena) exited the start box (Tosetto et
201 al., 2017). The emergence time and boldness have an inverse relationship, i.e., a decrease in the
202 emergence time leads to an increase in the boldness value. The highest scores, i.e., 1200 seconds,
203 were given to the fish that did not appear in the test arena.

204

205 *2.5.2. Open field test*

206 The open field was represented by a rectangular arena (57 cm long × 57 cm wide) and was filled
207 with oxygenated water. The bottom of the apparatus was divided into the centre, which made up
208 60% of the whole arena, and the periphery, which made up 40% (Supplementary Figure S6)
209 (Araújo and Malafaia, 2020). Body measurements (weight, standard length and total length) of
210 each fish were measured one day prior to the experiment. Fish were fed 24 h before the experiment.
211 The fish were individually introduced in the centre of the arena and allowed to acclimatize for 10
212 min and later recorded for 20 min using a webcam (Logitech C270, Laussane, Switzerland)
213 connected to a computer running iSPY software (www.ispyconnect.com). The distance travelled
214 by the fish (measured in cm) was calculated as well as the ratio of locomotion in the arena centre
215 to total locomotion during the test period. Video recordings were analysed using LoliTrack 5
216 (Loligo Systems, Tjele, Denmark).

217

218 *2.5.3. Swimming performance*

219 Evaluation of swimming performance and oxygen consumption was performed in a 5-L (testing
220 section 28 x 7.5 x 7.5 cm) Steffensen-type swimming tunnel respirometer (Loligo systems, Tjele,

221 Denmark) submerged in a buffer tank that was connected to an aerated temperature-controlled 50-
222 L reservoir tank allowing continuous water exchange (Supplementary Figure S7). The swimming
223 chamber was connected to a buffer tank through a flush pump (5 L/min, Eheim GmbH, Deizisau,
224 Germany) to ensure an adequate dissolved oxygen concentration for swimming performance
225 testing. Flow calibrations were performed using a handheld flowmeter (Flowtherm, Höntzsch
226 GmbH, Waiblingen, Germany). The level of dissolved oxygen in the test chamber was kept above
227 70% throughout the experiment (Tran et al., 2021). The temperature and dissolved oxygen in the
228 swimming chamber were continuously monitored using a fibreoptic oxygen probe and a
229 temperature probe coupled to a Witrox 1 sensor meter (Loligo Systems, Tjele, Denmark). The
230 water temperature was maintained at 20 ± 0.1 °C. The tank was covered with paper to prevent
231 disturbance from the outside. The water flow and dissolved oxygen in the swimming chamber
232 were monitored and controlled by the system via AutoResp software (Loligo Systems, Tjele,
233 Denmark).

234

235 *2.6. Experimental protocol*

236 A total of 120 fish (20 fish/diet group) were used in swimming tests following 24 h without
237 feeding. The standard length (in mm) and weight (in g) were recorded prior to the experiments.
238 Fish were randomly and rotationally selected from the species and treatment groups (a single fish
239 was measured at a time) and identified by the PIT tag number. Individual fish were transferred to
240 the swimming tunnel and allowed to acclimatize to the test conditions for 10 min with a water flow
241 velocity of 10 cm s^{-1} . The tunnel was closed to avoid water exchange with the surrounding tank.
242 The initial velocity was set at 10 cm s^{-1} and increased by 5 cm s^{-1} increments every two min,
243 terminating at a maximum of 150 cm s^{-1} or when the fish was exhausted (stopped swimming). The

244 small increments of velocity and time in our protocol were set to minimize stress on the tested fish,
245 and the protocol follows Tran et al. (2021). The swimming test was terminated when the test fish
246 remained at the rear grid for more than 10 s without any activity. The critical swimming speed
247 (U_{crit} , cm s^{-1}) was calculated using the previously described equation (Brett, 1964):

$$248 \quad U_{crit} = U_{max} + (T_{max}/T_{interval} * U_{interval})$$

249 where U_{max} is the highest velocity recorded at fatigue (cm s^{-1}); $U_{interval}$ is the velocity interval (5
250 cm s^{-1}); T_{max} is the time spent at fatigued velocity; and $T_{interval}$ is the time interval (2 min).

251

252 *2.7. Data analysis*

253 Possible effects of species, treatment, initial standard length (SL) and time of exposure on the
254 behavioural parameters (emergence time, swimming activity and open field utilization) were
255 analysed using linear mixed effects models (LMMs). Individual fish identity was used as a random
256 intercept in the models to account for the among-individual differences in behaviour. Model
257 selection based on the corrected Akaike information criterion (AICc) was used to identify the most
258 parsimonious model with the lowest AICc and other plausible models ($\Delta\text{AICc} \leq 2$) for the data
259 (Burnham and Anderson, 2002). Four models per behavioural variable were fit, including the full
260 model with species, treatment, initial SL and the statistical interaction between treatment and time,
261 and possible submodels (behaviour parameter \sim species; species + treatment; species + treatment
262 + SL) were among the candidate models of individual fish behaviour in the experiment. The
263 analyses were conducted in R (R Core Team, 2022) using the packages nlme and jtools (Long,
264 2022; Pinheiro et al., 2021). To gain better insight into changing behaviour across different
265 treatments and times of exposure, forest plots were obtained using log-transformed ratios of mean
266 response variables (i.e., boldness, open field utilization and swimming activity) in fish fed

267 contaminated pellets (PE and TWPs). Forest plots were generated using R Studio (R Core Team
268 2016) through the meta-analysis packages “metafor” (Quintana, 2015; Viechtbauer, 2010) and
269 “robumeta”; the script was created by Quintana (2015) and modified using the log-transformed
270 ratios of means as the effect size.

271 To test whether exposure to TWPs or PE reduced the fish swimming capacity using cm s^{-1}
272 and body lengths s^{-1} , a general linear model (GLM) was constructed with swimming performance
273 as the response variable and species, treatment and fish weight as explanatory variables.

274

275 **3. Results and discussion**

276 No fish mortality under experimental conditions was observed. Comparison of LMMs of fish
277 emergence time showed that the most parsimonious model was the most complex and included the
278 explanatory variables species, initial SL, treatment, interaction of treatment and time and fish ID
279 as a random intercept, with an R^2 of 0.27. The effect of species and SL was not significant in the
280 model, nor was the effect of treatment, but the emergence time significantly decreased as a result
281 of the interaction between treatment and time at 60 days, being lowest in both the PE and TWPs
282 treatments (Table 1, Figure 1). Fish behaviour in the open field test was best explained by a simple
283 model including only species and treatment (AIC=413); however, fish behaviour in the open field
284 test was best explained by the most complex model that also involved SL and time effects, with
285 an R^2 of 0.36 (AIC=420). Species identity was significant in the model, and a further significant
286 change was observed as a result of the interaction between time at 60 days and PE, with higher
287 utilization of the open field in crucian carp than in the other groups and experiment time (Table 2,
288 Figure 2). In the case of fish activity, the most parsimonious model was a simple model with only

289 the explanatory variable species; however, the AIC was very similar for all models. The most
290 complex model showed a significant effect of time at 14 and 60 for TWPs (Table 3, Figure 3).

291 The most complex GLM with fish swimming speed was chosen based on the AIC, and it
292 showed high significance of species on maximum swimming speed ($t = 9.34$, $p < 0.001$; Figure 4)
293 and a positive effect of the TWPs treatment on swimming speed in gibel carp ($t = 2.22$, $p = 0.03$;
294 Figure 4). The best model with the response variable swimming speed expressed as body lengths
295 s^{-1} was a GLM with species as the explanatory variable ($t = 9.63$, $p < 0.001$).

296 Forest plot analysis showed that, compared to controls, PE contamination led to a
297 significantly negative effect (i.e., decrease in exploratory behaviour) on crucian carp in the open
298 field test, after 14 days of exposure ($\text{LnR} = -0.49 \pm 0.28$; 95% CI) and decrease in the emergence
299 time of the crucian carp after 60 days ($\text{LnR} = -0.58 \pm 0.28$; 95% CI). In contrast, PE led to a
300 positive (i.e., decrease in the emergence time) effect on bolder behaviour for gibel carp after 60
301 days ($\text{LnR} = 0.43 \pm 0.46$; 95% CI; Figure 5). TWPs exerted a significantly positive effect on gibel
302 carp in the open field test after 14 days of exposure ($\text{LnR} = 0.20 \pm 0.19$; 95% CI), and a significant
303 negative effect on emergence time after 60 days of exposure ($\text{LnR} = -0.47 \pm 0.46$; 95% CI; Figure
304 5).

305
306 Our results suggest that within laboratory settings, sublethal doses of PE and TWPs may
307 cause fish to change their behaviour in the direction of more active and bold behaviour, especially
308 in native crucian carp. In the present study, we also obtained highly significant results in the case
309 of species for maximum swimming speed, indicating a positive shift caused by TWPs exposure.
310 The swimming speed of the invasive species increased significantly, which indicates that there is
311 a possible threat to species interactions, since locomotion is being affected by the experimental

312 treatments, especially TWPs. Such induced behavioural changes may lead to more risk-taking
313 behaviour in the wild, with potential repercussions for predator-induced mortality rates (Houston
314 et al., 1993; Hulthén et al., 2017; McCormick et al., 2018). The ecotoxic effect of PE MPs on
315 different fish species was addressed in a laboratory study, and trophic transfer of PE from *Poecilia*
316 *reticulata* (primary consumer) to *Danio rerio* adults (secondary consumer) resulted in behavioural
317 changes such as deficits in the anti-predatory defensive response in the organisms in the upper
318 trophic level (da Costa Araújo et al., 2020). The study of Araújo and Malafaia (2020) suggested
319 that PE MP accumulation in the tadpole *Physalaemus cuvieri* affects the locomotion ability,
320 anxiogenic behaviour, and antipredator response deficit in anurans exposed to potential predators.

321 **3.1 Evidence of the pollution resistance hypothesis**

322 Our results also provided support for the pollution resistance hypothesis, showing that
323 invasive species reacted to the toxicity of PE and TWPs with less pronounced changes in behaviour
324 triggered by PE and TWPs compared to native species. Bold individuals of zebrafish were found
325 to have a higher exposure burden to MPs than shy individuals, which is contrary to the common
326 perception of better survival chances among bolder individuals (Chen et al., 2022). The pollution
327 resistance hypothesis is likely valid only in selected scenarios. The opposite example is that native
328 *Artemia* species are extremely resistant to Hg, which prevents the invasion of non-native *Artemia*
329 *franciscana* (Pais-Costa et al., 2020).

330 Most of the available knowledge highlights the low sensitivity of gibel carp to
331 environmental contaminants and pollution (Gkelis et al., 2006; Kagalou et al., 2008; Perdikaris et
332 al., 2012), while comparative information on the two study species in relation to pollutants is still
333 missing. The results presented here further corroborate the higher resistance of invasive species to
334 MPs and provide insights into the potential effects that, synergically with other disturbances, could

335 strengthen biological invasion and replacement in European freshwater ecosystems. While the
336 replacement of native fish by invasive species likely occurs for many reasons, including the
337 competitive abilities of the two species (Tapkir et al., 2022), the most pronounced decline in
338 crucian carp was observed in degraded and suboptimal habitats, such as the oxbows of large river
339 systems (Buj et al., 2023; Kottelat and Freyhof, 2007; Lusk et al., 2010), which supports the
340 pollution resistance hypothesis.

341 **3.3 Toxicity of PE and TWPs: current knowledge**

342 The compounds leached from tires into water include a multitude of chemicals (Chibwe et
343 al., 2022; Halle et al., 2021), which have potential toxic effects on fish behaviour and performance
344 (Chang et al., 2023; Chibwe et al., 2022). Toxicity studies documented differential effects of
345 leachate originating from worn and new tires, with the latter showing long-term acute toxic effects
346 in freshwater crustacean *Hyalella azteca* (Halle et al., 2021). The toxicity of tire leachate has also
347 been addressed in early life stages of zebrafish (*Danio rerio*) embryos, which displayed a dose-
348 dependent reduced swimming performance (i.e., velocity, locomotive behaviour, total distance
349 travelled) (Varshney et al., 2022). Studies also indicated the toxic effect of TWP leachate on
350 marine phytoplankton, the base of marine food webs, with a reduced growth rate observed in three
351 phytoplankton species, namely, the cryptophyte *Rhodomonas salina*, the diatom *Thalassiosira*
352 *weissflogii* and the dinoflagellate *Heterocapsa steinii* (Page et al., 2022).

353 In adult zebrafish, the exposure to 10–600 μm PE particles at a concentration of 2 mg L^{-1}
354 resulted in abnormal behaviours such as erratic movement, seizures and downwards tail bending
355 (Mak et al., 2019). The short-term trophic transfer of MPs from beach hoppers to fish did not affect
356 fish behaviour; however, there was a shift in boldness with fish becoming shyer due to changes in
357 diet (Tosetto et al., 2017).

358 The research conducted thus far indicates that the effect of TWPs, PE and other pollutants
359 on behaviour may vary depending on the fish species tested. These studies are of ecological
360 importance; longer exposure to pollutants can disturb the fitness, cognitive, physiological and
361 behavioural patterns of fish from an early stage of life (Jacquin et al., 2020). Accumulation of these
362 pollutants can result in interpopulation divergence and affect ecosystems disproportionately by
363 selectively removing sensitive species.

364 **3.4 Role of behaviour in predator–prey interactions**

365 In the comparison of invasive gibel and native crucian carp, the study showed that although
366 gibel carp had an overall trend toward higher scores in activity, maximum swimming performance
367 and open field utilization (i.e., more active and exploratory behaviour) in the experiments, gibel
368 carp also took much longer to emerge from the box, suggesting more timid behaviour. During the
369 experiment, there was a likelihood for habituation in fish, e.g., reduced time of emergence and
370 increased activity, which could also diminish the perception of the impacts of PE and TWPs on
371 behaviour. To avoid habituation effects, more natural conditions, such as outdoor mesocosms and
372 telemetry testing, can be used to address behavioural changes (Lennox et al., 2021; Šmejkal et al.,
373 2022). In particular, crucian carp were found to be susceptible to habituation, and their behaviour
374 also changed in the expected direction in the control treatments, with crucian carp showing an
375 overall trend of being substantially more relaxed in the laboratory than gibel carp.

376 Native crucian carps are usually not strong in avoiding predation (Holopainen et al., 1997),
377 and their main adaptation involves inducible morphological changes in body depth (Brönmark and
378 Miner, 1992; Domenici et al., 2008; Hulthén et al., 2014) or an eventual switch from nocturnal to
379 aperiodic activity (Pettersson et al., 2001). In comparison with crucian carp, invasive gibel carp
380 thrive even in predator-rich environments, such as the main channel and oxbows of the Elbe River,

381 Czech Republic (Daněk et al., 2012; Lusk et al., 2010). The observed change in the behavioural
382 trends in relation to predation risks in crucian carp needs to be tested before drawing any
383 conclusions, but it may have the potential to further weaken its anti-predation skill.

384 **3.5 Relevance for species and environment**

385 Despite the extreme resistance of crucian carp to abiotic factors (Blažka, 1958), native
386 crucian carp is facing a decline due to the introduction and competitive abilities of invasive gibel
387 (Jeffries et al., 2016; Kottelat and Freyhof, 2007; Tapkir et al., 2022). Both native crucian carp and
388 invasive gibel carp are well adapted to the non-favourable conditions of ageing pools of
389 floodplains; e.g., they are extremely hypoxia- and heat-tolerant species (Antonova, 2010;
390 Bundgaard et al., 2020; Jackson, 2000; Karvonen et al., 2005; Piironen and Holopainen, 1986).
391 Furthermore, gibel carp were found to survive ammonia concentrations (12.5 mg L^{-1} ; pH 8.6)
392 (Nathanailides et al., 2003) and toxic cyanobacterial blooms (Perdikaris et al., 2012) by storing
393 toxins in the liver and other tissues (i.e., ovaries, brain, intestine, muscle and kidneys) (Gkelis et
394 al., 2006; Kagalou et al., 2008). Such high environmental tolerance to abiotic environmental
395 factors is hypothesized to make gibel carp strong enough to outcompete native crucian carp in its
396 suboptimal and degraded habitat (Kottelat and Freyhof, 2007), rendering gibel carp one of
397 Europe's most effective invaders (Perdikaris et al., 2012). While crucian carp is known to prefer
398 floodplain channels with rooted floating aquatic vegetation (Sayer et al., 2011; Tonn et al., 1992;
399 Wheeler, 2000), gibel carp can withstand man-made as well as natural waterways such as streams,
400 rivers, canals, dams, reservoirs, estuaries and ponds (Tarkan et al., 2012; Vetemaa et al., 2005),
401 which could be exposed to high levels of contamination, and where crucian carp populations
402 dropped in recent decades (Lusk et al., 2010; Lusková et al., 2010).

403 In the present work, we used a concentration of 0.1% of MPs, incorporated in food pellets.
404 We acknowledge that comparison with environmental concentration of MPs across different
405 matrixes is not feasible, since it is impossible to determine the actual exposure to biota starting
406 from the environmental concentration. However, the very few reported weight-based
407 concentrations of MPs, regardless of the matrix, were higher or similar to those used in the present
408 study, and we thus believe that our exposure concentrations can be considered relevant and useful,
409 particularly for foundational studies such as the present one. In order to more accurately replicate
410 the experimental conditions such as concentration and modes of ingestion in the wild, a more
411 natural scenario and typical ingestion of filter feeders like *Daphnia* can be developed for future
412 work based on these behavioural experiments. With respect to the studied species, it may be
413 expected that native crucian carp receives a slightly higher dose of contaminants than gibel carp
414 since the trophic position of crucian carp is higher (feeding solely on zooplankton and benthic
415 invertebrate prey) (Batel et al., 2016; Farrell and Nelson, 2013; Setälä et al., 2014), while part of
416 the gibel carp diet consists of plant material (de Meo et al., 2022; Özdilek and Jones, 2014; Tapkir
417 et al., 2023).

418 **4. Conclusions and future directions**

419 Available scientific literature provides evidence of acute poisoning from road run-off in the case
420 of TWPs and negative effects associated with the accidental ingestion of PE particles. Our study,
421 on the contrary, documents the long-term effects of sublethal concentrations of PE and TWPs,
422 comparing native and invasive species of carps, within freshwater ecosystems. We observed a
423 tendency of crucian carp to reduce its scores related to cautious behaviour (reduced emergence
424 time and increased use of open space) in the PE treatment. It remains to be tested whether higher
425 doses or longer exposure to pollutants would lead to more pronounced changes and whether such

426 changes would be relevant for interactions with other organisms, especially with predators. Our
427 results, based on relevant sublethal concentrations of TWPs, emphasize the need to further identify
428 the potential effects of the polymers included in MPs.

429 Plastic production, use and management will likely undergo a deep remodulation that will
430 hopefully result in reduced plastic input into the environment (e.g., global plastic treaty; Bergmann
431 et al., 2022). Ideally, this should apply also to TWPs, which should experience the same
432 remodulation forecasted for plastic. Unfortunately, this cannot be fully true for TWPs because, at
433 present, there are no other options that could replace tires for large-scale road transportation, even
434 though a shift in materials used for their production can be foreseen. However, focusing on the
435 near future, there will likely be a consistent input of TWPs through its numerous pathways that
436 span from atmospheric deposition to runoff waters (Wagner et al., 2018). Therefore, the important
437 goal is to evaluate potential negative impacts on the environment and important changes in species
438 interactions that, in specific cases, can eventually foster biological invasions in some ecosystems.

439

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447 **Ethical statement**

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453

454 **Conflict of interest**

455 The authors declare no competing financial interests.

456

457 **Ethical approval**

458 The field sampling and experimental protocols used in this study were performed in accordance
459 with the guidelines and permission from the Experimental Animal Welfare Commission under the
460 Ministry of Environment of the Czech Republic (ref. no. CZ 01679). The methods and ethics of
461 the study were approved by the Experimental Animal Welfare Commission of Biology Centre of
462 the Czech Academy of Sciences, and a certificate of approval is available upon request.

463

464 **Informed consent**

465 Not applicable.

466

467 **Authors' contributions**

468 Pankaj A. Gorule: data curation; investigation; writing—original draft; visualization; formal
469 analysis; methodology; writing—review and editing

470 Marek Šmejkal: conceptualization; methodology; data curation; investigation; validation; formal
471 analysis; supervision; visualization; funding acquisition; writing—original draft; writing—review
472 and editing; resources
473 Sandip Tapkir: formal analysis, writing review and editing
474 Yevdokiia Stepanyshyna: methodology; writing—review and editing
475 Vlastimil Stejskal: methodology; data curation; conceptualization; writing original draft;
476 writing—review and editing
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479 funding acquisition; writing—original draft; writing—review and editing

480

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830 **Tables**

831 **Table 1.** Linear mixed effects model for the boldness-emergence time of gibel carp and crucian
 832 carp from the observations recorded for a group of 120 fishes, with 40 fishes each in the control
 833 and 14-day and 60-day exposure periods. The estimated parametric coefficients and their
 834 significance (coefficients for Species, SL, Treatment and the interaction between Treatment and
 835 Exposure Time) in the model are shown. The adjusted R² of the model with the random intercept
 836 fish ID and dependent variable emergence time was 0.27, and the deviance explained by the ICC
 837 values was up to 21%. Significant p values for the explanatory variables were calculated using
 838 Satterthwaite d.f.
 839

	Est	S.E.	t val.	d.f.	p
Intercept	18.96	203.77	0.09	174.77	0.93
Species (gibel carp)	47.69	28.24	1.69	116.88	0.09
SL	1.90	2.15	0.88	171.48	0.38
Treatment PE	-23.12	49.36	-0.47	319.08	0.64
Treatment TWPs	18.30	49.36	0.37	319.07	0.71
Treatment control Time 14	-33.41	43.84	-0.76	234.08	0.45
Treatment PE Time 14	-35.23	43.91	0.80	235.26	0.42
Treatment TWPs Time 14	-73.85	43.92	-1.68	235.37	0.09
Treatment control Time 60	-133.43	44.21	-3.02	240.41	0.00
Treatment PE Time 60	-106.37	44.39	-2.40	243.44	0.02
Treatment TWPs Time 60	-182.14	44.23	-4.12	240.78	0.00

841 **Table 2.** Linear mixed effects model for the open field utilization of gibel carp and crucian carp
842 from the observations recorded for a group of 120 fishes, with 40 fishes each in the control and
843 14-day and 60-day exposure periods. The estimated parametric coefficients and their significance
844 (coefficients for Species, SL, Treatment and interaction between Treatment and Exposure Time)
845 in the model are shown. The adjusted R^2 of the model with the random intercept fish ID and
846 dependent variable RatioN was 0.36, and deviance explained by the ICC values was up to 28%.
847 Significant p values for the explanatory variables such as fish identity were calculated using
848 Satterthwaite d.f. The most appropriate parsimonious model was the most complex model, selected
849 based on the AIC.

850

	Est	S.E.	t val.	d.f.	p
Intercept	-0.12	0.13	-0.89	184.93	0.37
Species (gibel carp)	0.08	0.02	4.45	117.10	0.00
SL	0.00	0.00	1.96	182.51	0.05
Treatment PE	-0.03	0.03	-0.85	300.65	0.40
Treatment TWPs	-0.00	0.03	-0.12	300.65	0.91
Treatment control Time 14	0.06	0.03	2.12	234.07	0.04
Treatment PE Time 14	0.05	0.03	1.82	235.39	0.07
Treatment TWPs Time 14	0.06	0.03	2.15	235.51	0.03
Treatment control Time 60	0.03	0.03	1.08	241.09	0.28
Treatment PE Time 60	0.06	0.03	2.28	244.44	0.02
Treatment TWPs Time 60	0.01	0.03	0.27	241.50	0.79

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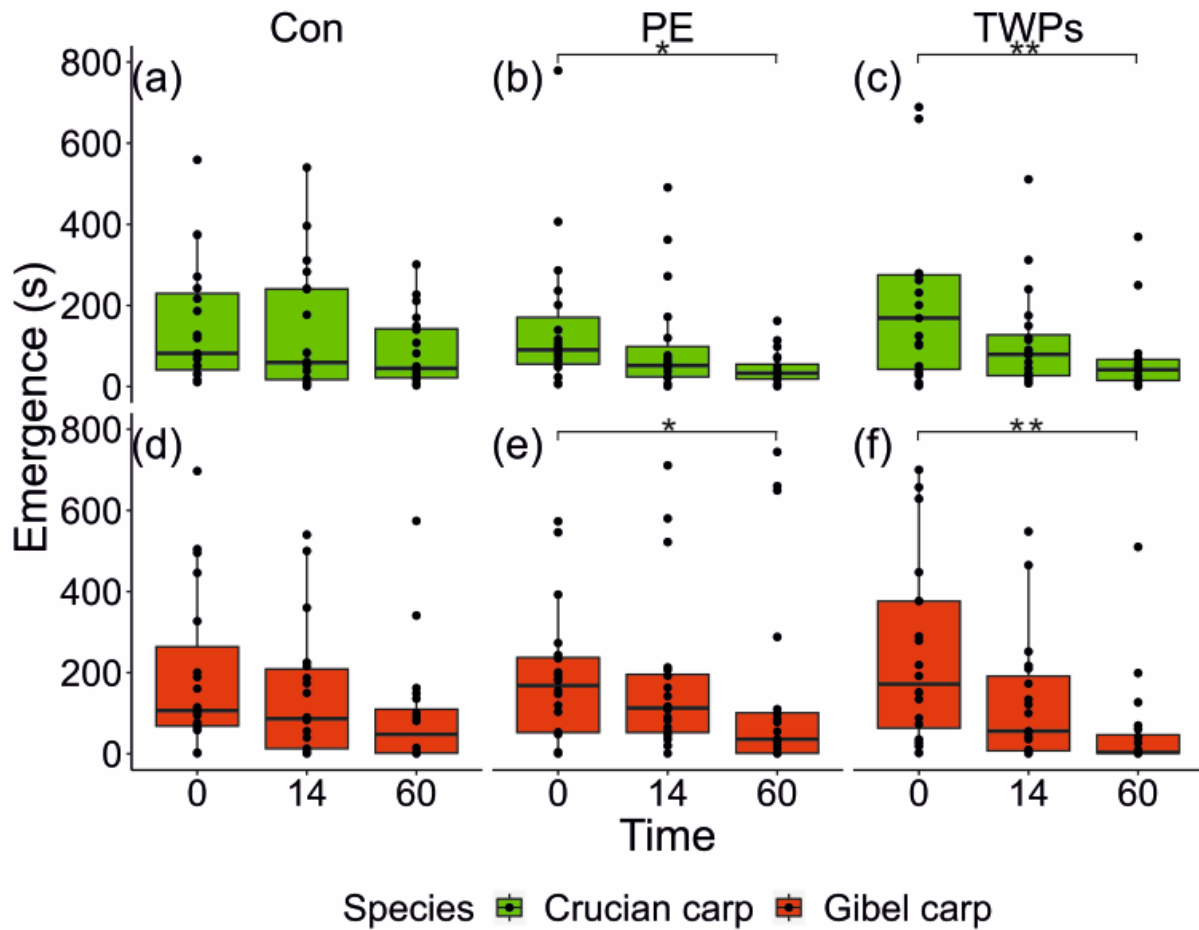
853 **Table 3.** Linear mixed effects model for the total activity in the open field test of gibel carp and
854 crucian carp from the observations recorded for a group of 120 fishes, with 40 fishes each in the
855 control and 14-day and 60-day exposure periods. The estimated parametric coefficients and their
856 significance (coefficients for Species, SL, Treatment and interaction between Treatment and
857 Exposure Time) in the model are shown. The adjusted R² of the model with the random intercept
858 fish ID dependent variable TotalN was 0.18, and the deviance explained by the ICC values was up
859 to 14%. Significant p values for the explanatory variables such as fish identity with respect to time
860 were calculated using Satterthwaite d.f.

	Est	S.E.	t val.	d.f.	p
Intercept	6087.21	2783.74	2.19	165.27	0.03
Species (gibel carp)	588.99	378.98	1.55	116.40	0.12
SL	-1.27	29.40	-0.04	161.02	0.97
Treatment PE	-737.33	698.99	-1.05	335.18	0.29
Treatment TWPs	-760.97	699.00	-1.09	335.17	0.28
Treatment control Time 14	95.57	648.98	0.15	233.83	0.88
Treatment PE Time 14	-35.10	649.85	-0.05	234.87	0.96
Treatment TWPs Time 14	1684.65	649.93	2.59	234.97	0.01
Treatment control Time 60	253.28	653.64	0.39	239.42	0.70
Treatment PE Time 60	1043.68	655.89	1.59	242.10	0.11
Treatment TWPs Time 60	1546.91	653.91	2.37	239.74	0.02

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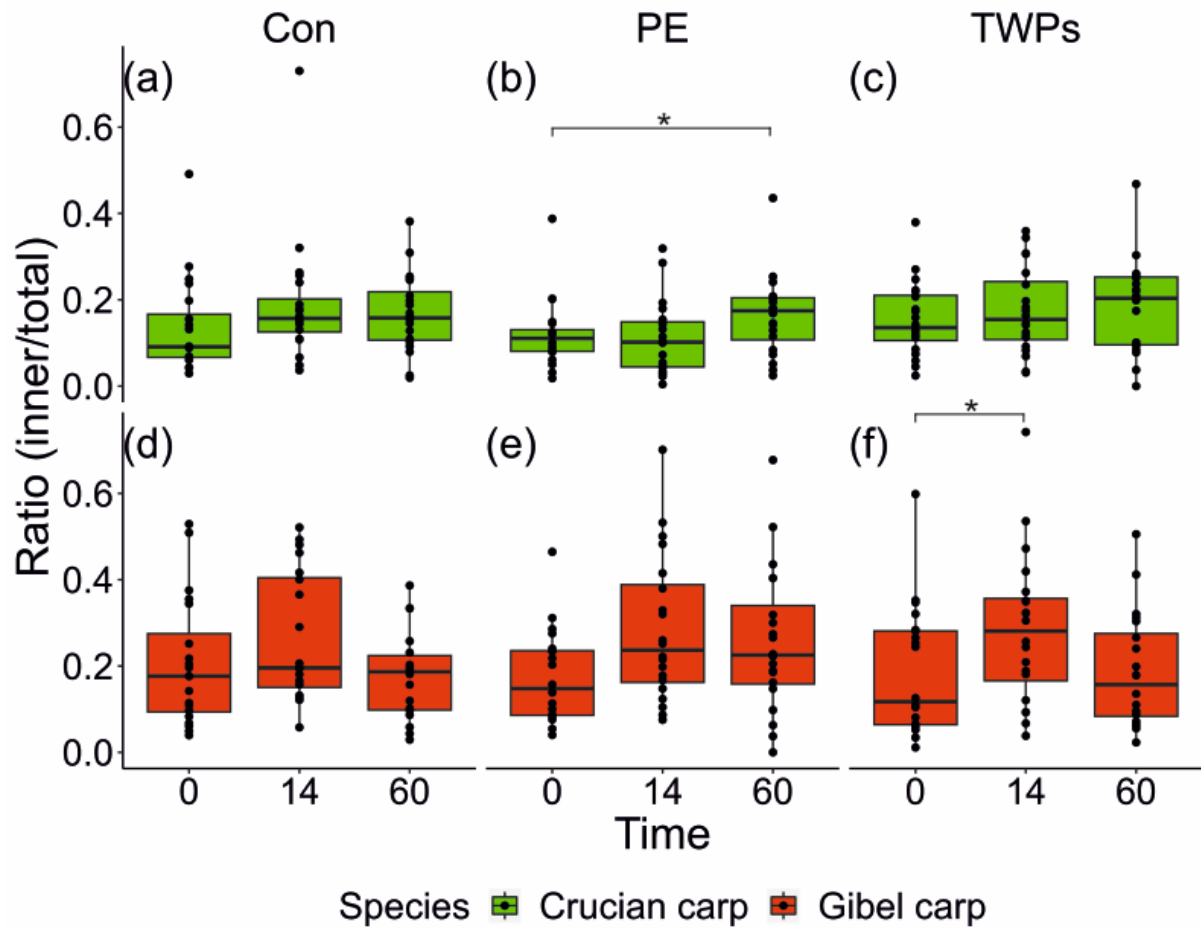
863 **Figures**



864

865 **Figure 1.** Emergence time (a proxy of fish boldness) differed between native crucian carp
 866 (*Carassius carassius*) and invasive gibel carp (*C. gibelio*) and declined with experimental duration
 867 (evaluated at 0, 14 and 60 days). PE treatment – 0.1% polyethylene microplastic food content,
 868 TWPs treatment – 0.1% tire wear particles food content, Control – no addition to the food pellets.
 869 Points = individual data; boxplots: thick line = median, box = 50% of interquartile range, whiskers
 870 = outer 25% of interquartile range excluding outliers.

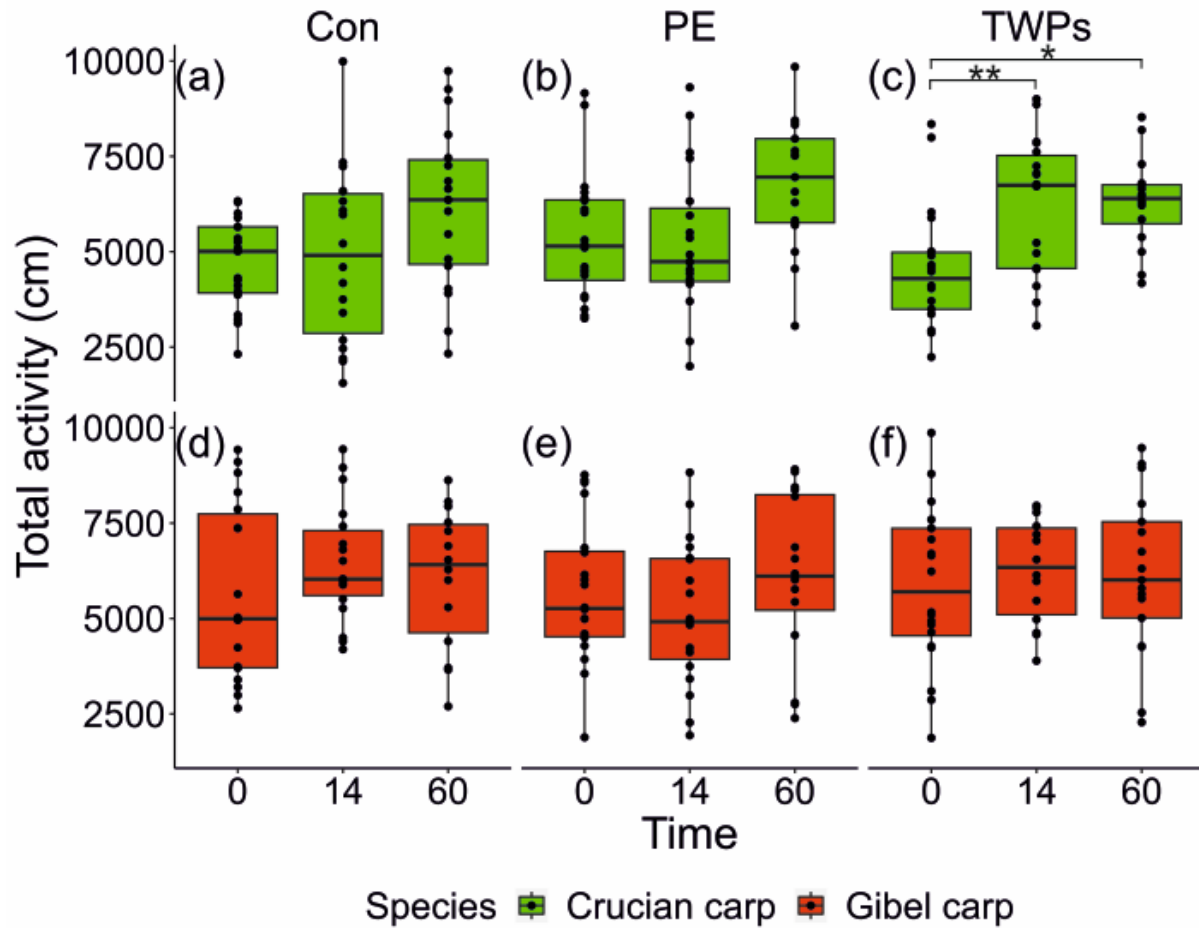
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873 **Figure 2.** Ratio of time spent in the arena centre to the total test time of native crucian carp
 874 (*Carassius carassius*) and invasive gibel carp (*C. gibelio*) during exposure to the control, PE and
 875 TWPs treatments. The native species were more prone to risk at the end of the PE treatment. PE
 876 treatment – 0.1% polyethylene microplastics food content, TWPs treatment – 0.1% tire wear
 877 particles food content, Control – no addition to the food pellets. Points = individual data; boxplots:
 878 thick line = median, box = 50% of interquartile range, whiskers = outer 25% of interquartile range
 879 excluding outliers.

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882 **Figure 3.** Total activity of native crucian carp (*Carassius carassius*) and invasive gibel carp (*C.*

883 *gibelio*) represented as the total distance swam in cm during the behavioural trials. The invasive

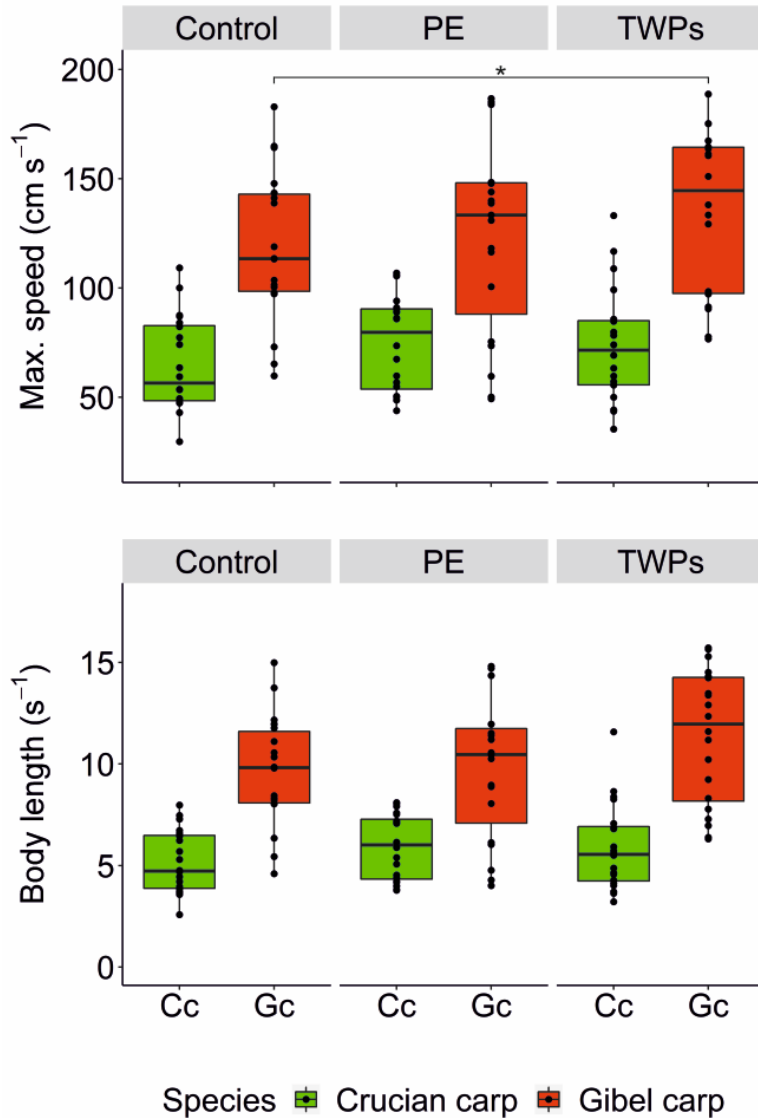
884 species were more locomotive than the native species in all the treatments. PE treatment – 0.1%

885 polyethylene microplastics food content, TWPs treatment – 0.1% tire wear particles food content,

886 Control – no addition to the food pellets. Points = individual data; boxplots: thick line = median,

887 box = 50% of interquartile range, whiskers = outer 25% of interquartile range excluding outliers.

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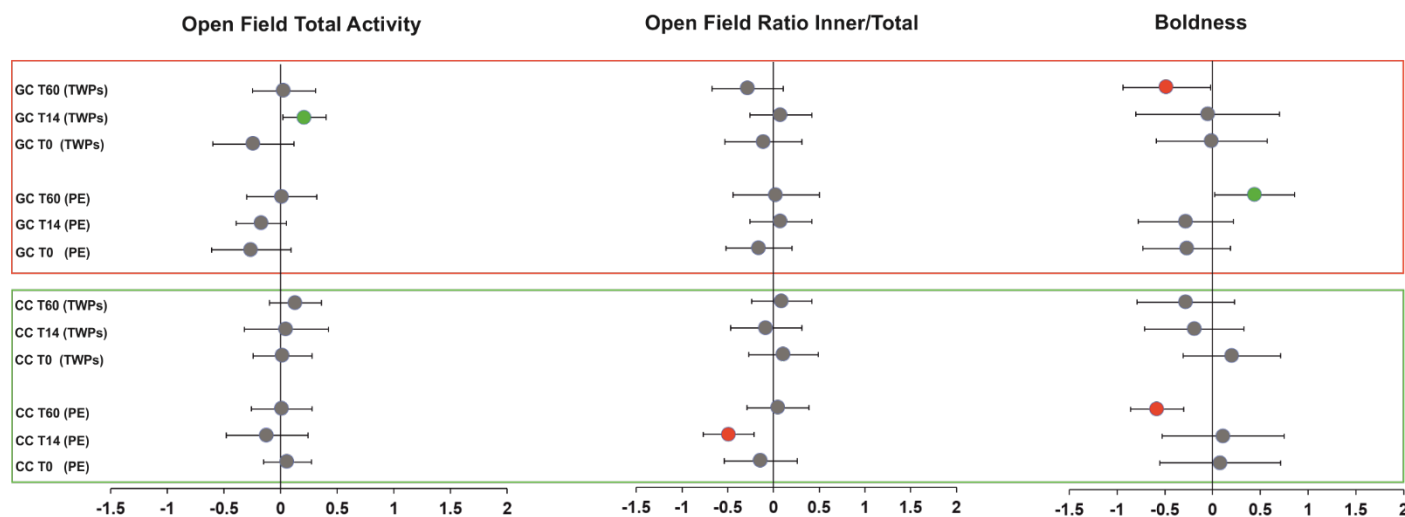
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Figure 4. Swimming activity of native crucian carp (*Carassius carassius*) and invasive gibel carp (*C. gibelio*) represented as the total distance swam in cm s⁻¹ at the end of the 60-day experimental period. The invasive species had better swimming performance than the native species in all the treatments. PE treatment – 0.1% polyethylene microplastics food content, TWPs treatment – 0.1% tire wear particles food content, Control – no addition to the food pellets. Points = individual data; boxplots: thick line = median, box = 50% of interquartile range, whiskers = outer 25% of interquartile range excluding outliers.



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898

899 **Figure 5.** Forest plot showing cumulative impacts the native *Carassius carassius* (CC) and the
 900 invasive *C. gibelio* (GC) due to different treatments, PE and TWPs (ln–response ratio; mean \pm 95%
 901 confidence intervals). Outer rectangular red and green lines around the forest plots indicate a
 902 distinction between the invasive gibel carp (on the top) and native crucian carp (at the bottom).
 903 Red circles: negative ratios; green circles: positive ratios; grey circles: non-significant ratios,
 904 across time 0, 14 and 60 days. Cumulative effects are significant if confidence intervals do not
 905 overlap zero.

906