

Research article

Are microplastics contributing to pollution-induced neurotoxicity? A pilot study with wild fish in a real scenario

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ABSTRACT

Pollution-induced neurotoxicity is of high concern. This pilot study investigated the potential relationship between the presence of microplastics (MPs) in the brain of 180 wild fish (*Dicentrarchus labrax*, *Platichthys flesus*, *Mugil cephalus*) from a contaminated estuary and the activity of the acetylcholinesterase (AChE) enzyme. MPs were found in 9 samples (5% of the total), all of them from *D. labrax* collected in the summer, which represents 45% of the samples of this species collected in that season (20). Seventeen MPs were recovered from brain samples, with sizes ranging from 8 to 96 μm . Polyacrylamide, polyacrylic acid and one biopolymer (zein) were identified by Micro-Raman spectroscopy. Fish with MPs showed lower ($p \leq 0.05$) AChE activity than those where MPs were not found. These findings point to the contribution of MPs to the neurotoxicity induced by long-term exposure to pollution, stressing the need of further studies on the topic to increase 'One Health' protection.

1. Introduction

Worldwide plastic production has been growing faster than the manufacture of any other material [1]. Compared with two decades ago, the plastic production has doubled [2] and the environmental contamination by plastics considerably increased during the SARS-CoV-2 pandemics [3–6]. Due to their long persistence in the environment, plastics became a major pollution problem worldwide.

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Among plastics, debris with size lower than 5 mm commonly designated by microplastics (MPs) are of high concern. MPs present in the wild result from the progressive break down of larger plastic fragments and from the input of particles already with size lower than 5 mm into the environment [7]. Studies have shown that MPs are worldwide distributed in the environment, being present in the atmosphere, freshwater, transitional waters and marine water, soils and sediments, and food webs [8–14]. Such MPs generally have other chemicals (MP-Chem), including additives added during their production and preparation for several applications, and other environmental contaminants that adsorb to their surface when they are in the environment [15–17]. MPs may also carry microorganisms, including multi-resistant pathogenic ones [18].

When MPs enter into the body of animals through ingestion, respiration, or other ways, some of them are internalized into the circulatory system, as shown by several studies with different species [19,20], including humans [21]. Then, they are distributed, some are likely excreted whereas others reach internal organs and tissues where they can be retained and possibly accumulated [5,22–24]. Laboratory studies demonstrated that very small MPs, including nanoplastics (NPs), can cross the blood-brain barrier [25] and cause neurotoxicity through inhibition of the acetylcholinesterase (AChE) enzyme [26–28], brain oxidative stress and damage [29–31], alteration of neurotransmitters' levels and other processes as recently reviewed [32]. In fish, such effects have been related with decreased feeding [33,34], reduced swimming performance [35] and other behaviour alterations [36,37], among other effects that can reduce the individual and population fitness, with potential negative impacts on biodiversity, and human food safety and security [38]. Moreover, MPs interact with the toxicity of many other contaminants in fish often leading to a greater negative impact [29,39,40]. Furthermore, warmer water temperature and other alterations resulting from global warming influence the uptake, accumulation and toxicity of MPs [41,42], and often act synergistically with chemical toxicity in model animals, including in long-term exposures [4,43]. Therefore, there is high concern on the combined long-term effects of MPs, MP-Chem and other stressors on wild organisms and human health, including regarding neurotoxicity.

Data on the potential MP-induced neurotoxicity in wild populations of fish is still very limited [22,32,44]. The effects induced in real scenarios can be different from those obtained in the laboratory due to the influence of several factors, such as individual and population adaptation, properties of the MPs and other pollutants present, seasonal variation in the levels of MPs and environmental conditions (e.g., temperature, water volume, pH), among several others [45]. The differences may be more evident in populations living in highly dynamic ecosystems, such as contaminated estuaries of large rivers, which often have high loads of MPs and are important contributors to the marine pollution by these particles [46]. Considerable concentrations of MPs were documented in fish from estuaries located in different regions of the world, such as the La Plata River estuary, Argentina [47], the Ciliwung River estuary, Indonesia [48], the Tecolutla River estuary, Gulf of Mexico [49], and the Minho River estuary, Iberian Peninsula [5]. Further research with fish in real scenarios is needed to better understand the pollution-induced long-term neurotoxicity in fish, which is also important regarding human health and wellbeing.

The main goal of this pilot study was to investigate the potential presence of MPs in the brain of wild fish in relation to brain AChE activity in a real scenario of long-term exposure to pollution caused by a diversity of chemicals. Two null hypotheses were tested: H_{01} - The samples collected from fish brain do not have MPs; H_{02} - Brain AChE activity is not related with the presence of MPs in fish brain samples.



Fig. 1. Localization of the Douro River estuary with the fish sampling areas. All fish were captured in the area defined by the coordinates $41^{\circ} 8'37.95''N$, $8^{\circ}40'29.27''W$ and $41^{\circ} 8'39.11''N$, $8^{\circ}37'49.35''W$. Sea bass were mainly collected in the downstream area (blue) and grey mullet and flounders in the areas indicated in yellow and green. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

2. Material and methods

2.1. Alternative hypotheses, fish sampling and animal ethical issues

The alternative hypotheses to H_{01} and H_{02} were, respectively: H_{A1} - The samples collected from fish brain have MPs; H_{A2} - Brain AChE activity is related with the presence of MPs in fish brain samples.

Fish were from the estuary of the Douro River, hereafter indicated as Douro estuary, which is located in the Northwest (NW) coast of Portugal and ends into the North East (NE) Atlantic Ocean (Fig. 1). This estuary was selected because the hydrological basin of the Douro River is one of the most important in the Iberian Peninsula, the Douro estuary has an extent of about 22 km, draining into the Atlantic Ocean [50], its environmental conditions variate along the year [51], and it is contaminated with a diversity of MPs [52,53], among many other environmental contaminants [54–56].

Three fish species were investigated: the European seabass (*Dicentrarchus labrax* Linnaeus, 1758), the European flounder (*Platichthys flesus* Linnaeus, 1758) and the flathead grey mullet (*Mugil cephalus* Linnaeus, 1758). They were chosen for this study because they can be found in the Douro estuary along the year, they are fished for human consumption as food, and they were used in bio-monitoring studies in relation to MPs and other contamination in previous studies in estuaries [5,57–60]. A total of 180 fish were investigated, 20 per species and per season (summer 2019, autumn 2019, and winter 2019–2020). All the fish were obtained from local fishery, were captured in the area defined by the coordinates $41^{\circ} 8'37.95''N$, $8^{\circ}40'29.27''W$ and $41^{\circ} 8'39.11''N$, $8^{\circ}37'49.35''W$, and aimed at being sold for human food consumption. Shortly after their capture by fishermen, the fish corps were immediately transported intact to the laboratory to avoid contamination by MPs present in the environment, in thermally isolated boxes in cold conditions.

The study had authorization from the “ORBEA – Orgão Responsável pelo Bem Estar Animal” of ICBAS – School of Medicine and Biomedical Sciences of the University of Porto, reference number P372/2020/ORBEA, and was carried out according the European and Portuguese principles and procedures regarding animal’s ethics and experimentation. L. Guilhermino and L.R. Vieira have accreditation from the Portuguese Authority (“Direção Geral de Alimentação e Veterinária”), to coordinate studies and carry animal experimentation.

2.2. Quality assurance and quality control

Several quality assurance and quality control (QA/QC) measures were implemented during all the stages of sample collection and preparation (dissection, digestion, and filtration), MPs isolation and characterization (primary and chemical) to prevent external and cross-contamination of samples and particles as in previous studies [5,22]. Briefly, researchers used nitrile gloves and 100% cotton laboratory coats covering other clothes. The work was done in spaces with restrict access, the surface areas, materials and instruments were previously carefully cleaned and disinfected, the instruments were washed before and between tissue collection, among other measures. Dissections, preparation of samples and filtration were performed in a flow cabinet, to avoid the risk of external contamination. All the filters were previously observed under a stereo-microscope to remove any plastic debris potentially present. Blanks to control for external contamination were used in all the steps, and controls with potassium hydroxide (KOH) solution in ultra-pure water (10% v/v) were also used.

2.3. Sample collection

In the laboratory, the total body length (length) and the total body weight (wet weight – ww, hereafter indicated as body weight) of each fish were determined, and a basic physical examination of each fish was made. Then, each fish was carefully washed to remove

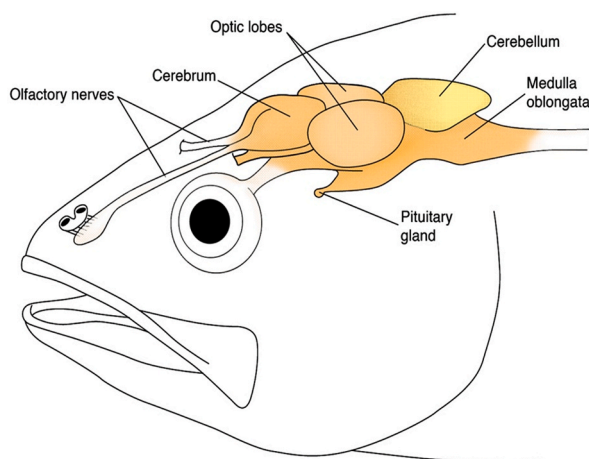


Fig. 2. Illustrative representation of the fish brain structure. Adapted from [64].

external particles, and several tissue/organ samples were collected for different types of analyses to take the best advantage of animal corps. For this study, the whole brain of each fish was isolated and divided into two portions (Fig. 2): the cerebellum was isolated and prepared for AChE activity determination because this region of fish brain has high activity of this enzyme [61,62]; the remaining regions of the brain were put together for MP analyses. Samples for AChE analyses were weighted, put in cold phosphate-buffer (1:10 w/v; 0.1 M, pH = 7.2), and kept at -80°C until further analyses. Samples for MP analyses were weighted (Kern ABS 120–4, Kern & Sohn GmbH, Germany), carefully rinsed with ultrapure water to remove any potential contamination [63], and then transferred to pre-cleaned glass vessels, which were sealed and externally covered with aluminium foil, and kept at -20°C until further analyses.

2.4. Microplastic isolation, visual identification and chemical characterization

The particles isolation and primary characterization were carried out as previously described [5,22,65]. In brief, each brain sample was covered with a volume of a 10% potassium hydroxide solution in ultra-pure water and incubated at 60°C for 24 h in an oven (Drying oven EV50, Raypa, Spain) [65]. After cooling, the product was filtered through 42 mm diameter glass-microfiber filter membranes (pore size $1.2\ \mu\text{m}$, Munktell & Filtrak GmbH, Germany) in vacuum conditions. Subsequently, the filters were put in previously cleaned glass Petri dishes, which were sealed and dried at 40°C for 24 h (Drying oven EV50, Raypa, Spain). All the particles present in the filters were observed and pictures were taken (Nikon SMZ800 Stereo Microscope with integrated camera DS-Fi1, Japan). Each particle was measured using the ImageJ software (<https://imagej.nih.gov/ij/>), and its shape and colour were recorded.

The chemical composition of all the particles isolated from brain samples was determined by Raman spectroscopy, a method used successfully to identify the plastic nature of particles in other studies [66,67]. The Raman spectra of the particles were recorded using a WITec confocal CRM alpha 300 R Micro-Raman spectroscopy (WITec GmbH, Ulm Germany) with an air-cooled solid state laser operating at $532\ \text{nm}$ a CCD detector cooled to -60°C . Different Raman settings for the same spot were used or multiple spectra at different spots on the same sample were collected [67]. Subsequent processing of the data was conducted using a WITec Project software (FIVE 5.2, WITec GmbH, Ulm, Germany). All spectra were compared to reference libraries using the search/match software TrueMatch integrated with the WITec Suite FIVE software. Spectral identification was accepted only after spectra underwent visual confirmation with matched reference spectra and a minimum Hit Quality Index (HQI) of 70%.

After characterization, in addition of being sorted by chemical composition, the particles identified as MPs were quantified by shape, colour and size. For the shape, the following types were considered based on the recommendations of the Technical Subgroup on Marine Litter (TSG-ML) for the European Marine Strategy Framework Directive document: plastic fragments, pellets, filaments/fibres, plastic films, foamed plastic, granules and styrofoam [68]. Regarding colour, the following categories were considered: red, orange, yellow, green, blue, violet, black, white and transparent [69]. Considering size (longest dimension for fibres; Feret's diameter for the other shapes) the following size classes were considered: $<100\ \mu\text{m}$; $101\text{--}150\ \mu\text{m}$; $151\text{--}500\ \mu\text{m}$; $501\text{--}1500\ \mu\text{m}$; $1501\text{--}3000\ \mu\text{m}$; $3001 < 5000\ \mu\text{m}$ [22].

The concentrations of MPs in brain samples were expressed as the number of MP items per fish (MPs/fish) or per weight (ww) of the analysed brain tissue (MPs/g).

2.5. Determination of acetylcholinesterase activity

The activity of AChE was selected for the present study because it is widely used as a neurotoxicity biomarker, and several MPs were found to alter the activity of this enzyme in fish, both in laboratorial [26,29,70,71] and field conditions [22,72].

The procedures for further preparation of samples and determination of AChE activity were described in detail elsewhere [73,74]. Briefly, after defrosting on ice, each cerebellum sample was kept in phosphate-buffer (0.1 M, pH = 7.2), homogenized (1:10 w/v; Ystral GmbH d-7801 homogenizer, Germany) and centrifuged for 3 min at $3,300\ \text{g}$ and 4°C (Sigma 3K30 centrifuge, Germany). The supernatant was carefully collected, its protein content was determined by the Bradford method [75] adapted to microplate [76], using bovine gamma globulin as protein standard and absorbance read at $600\ \text{nm}$ (Power Wave HT 340, BioTek, USA). The protein content of the samples was standardized to $0.3\ \text{mg/mL}$ [74]. The AChE activity was determined by the Ellman technique [77] adapted to microplate [73], using acetylcholine as substrate and the absorbance read at $412\ \text{nm}$ (Power Wave HT 340, BioTek, USA). After enzymatic determinations, the protein content of the remaining samples was determined again (as previously indicated), and used to express the enzymatic activity in nanomoles of substrate hydrolysed per minute per mg of protein (nmol/min/mg protein).

2.6. Statistical analysis

Statistical analyses were carried out in the IBM® SPSS Statistics® package, version 28.0, and the significance level was 0.05.

For each species, the data set of each biological parameter (i.e., total body weight, total body length or AChE activity) was tested for normal distribution and homogeneity of variances through the Shapiro-Wilk test and the Levene's test respectively, and the variables were transformed when necessary [78]. When normal distribution and homogeneity of variances were achieved, a one-way analysis of variance (ANOVA) was used to compare different seasons (summer, autumn and winter), followed by the Tukey's multicomparison post-hoc test when significant differences were found. When the ANOVA assumptions could not be achieved, the Kruskal-Wallis test was used, followed by pairwise comparisons (with significance values adjusted by the Bonferroni correction for multiple test) when significant differences were found.

The Student's t-test was used to compare the AChE activity of fish having MPs in the brain sample analysed with the enzymatic activity of fish where MPs were not found in the analysed brain sample. For text simplicity, these two groups of fish will be hereafter

indicated as fish with MPs and fish without MPs in the brain, respectively. However, it should be noted that we cannot assure that fish with brain samples negative to MPs did not have these items in the brain because the cerebellum was used for AChE determinations and not analysed for MP content.

3. Results

The mean (\pm standard deviation – SD) of the biological parameters determined per species and season are indicated in Table 1, as well as the results of the statistical analysis comparing different seasons. In all the species, significant differences of body weight, length and AChE activity among seasons were found (Table 1). *D. labrax* specimens had significantly lower body weight and higher length in the winter, and lower AChE activity in the summer than in the other seasons. *P. flesus* specimens had significantly lower body weight, length and AChE activity in the winter than in the other seasons. *M. cephalus* specimens had significantly lower body weight in the winter, higher length in the autumn, and lower AChE activity in the autumn than in the other seasons.

Seventeen MPs were recovered from the brain samples, 94% fragments and 6% film (Fig. 3-A), which are exemplified in Fig. 4(A-F). The size of the MPs ranged from 8 μ m to 96 μ m, with 70% of them having less than 50 μ m (Fig. 3-B). Red, black and blue were the colours observed (Fig. 3-D). Three polymer types were identified: the superabsorbent polymers polyacrylamide (71%) and polyacrylic acid (23%), and one biopolymer (Zein) (6%) (Fig. 3-C). Representative spectra are shown in Fig. 5(A-C). Blue and green fibres were found in the blanks, however, no fibres were found in the analysed brain samples.

Among the 180 fish analysed, 9 (5%) had MPs in the brain, all were *D. labrax* specimens collected in the summer. From the total number of *D. labrax* specimens (60), 15% had MPs in the brain, with a mean concentration (\pm standard deviation - SD) in the brain of 0.3 ± 0.8 MPs/fish (2 ± 6 items/g). Considering the samples of *D. labrax* collected in the summer (20), 45% had MPs in the brain, with a mean (\pm SD) concentration of 0.9 ± 1.1 MPs/fish (7 ± 10 MP/g).

Among the specimens of *D. labrax* collected in the summer, significant differences in the activity of AChE between fish with and without MPs were found ($t_{(18)} = 5.383$, $p < 0.001$). The mean (\pm SD) of AChE activity was lower (80 ± 8 nmol/nmol/min/mg protein) in fish with MPs in the brain than in fish where MPs were not found in the analysed samples (117 ± 19 nmol/nmol/min/mg protein) (Fig. 6).

4. Discussion

The values of AChE activity determined in the studied specimens of *D. labrax* and *P. flesus* are in the range values reported in other studies for the same species. For example, AChE activity between 100 and 140 nmol/min/mg protein in wild *D. labrax* specimens from the Arade Estuary, Portugal [79], between 89 and 174 nmol/min/mg protein in wild *P. flesus* from the Minho River estuary, Portugal [80], and between 50 and 121 nmol/min/mg protein in wild *P. flesus* collected in the Gulf of Gdańsk, Poland [81], were documented. The total mean of AChE activity determined in *M. cephalus* from the Douro estuary is lower than the mean of 45.1 ± 26.4 μ mol/min/mg protein documented in wild flathead grey mullets from the Pontine Lakes [82], possibly reflecting biological and environmental differences.

Seasonal variability in fish weight and length was found, which did not have a common pattern to all the species. This is a common situation in estuarine fish and a diversity of factors can account for the differences, such as the biology of distinct species, life-cycle phase of each species, specific habitat and feeding ecology, variation of environmental conditions, such as availability of food, water temperature, concentration of contaminants, among others, and is important to take into consideration such variation in bio-monitoring studies [83].

Table 1

Mean and standard deviation of total body weight, body length and brain acetylcholinesterase activity in *Dicentrarchus labrax*, *Platichthys flesus*, and *Mugil cephalus* from the Douro estuary. Different letters indicate statistically significant differences ($p \leq 0.05$) among seasons per species.

	N	Total body weight (g)	Total body length (cm)	AChE activity (nmol/min/mg protein)
<i>Dicentrarchus labrax</i>				
Summer	20	656 \pm 89 a	31 \pm 4 a	100 \pm 24 a
Autumn	20	625 \pm 95 a	35 \pm 3 b	165 \pm 20 b
Winter	20	532 \pm 61 b	36 \pm 2 b	151 \pm 24 b
Total mean	60	604 \pm 97	34 \pm 4	139 \pm 36
Comparison among seasons	60	H ₍₂₎ = 21.000 $p < 0.001$	H ₍₂₎ = 20.643, $p < 0.001$	F _(2,57) = 46.017 $p < 0.001$
<i>Platichthys flesus</i>				
Summer	20	307 \pm 30 a,b	20 \pm 2 a	91 \pm 29 a
Autumn	20	337 \pm 65 a	30 \pm 2 b	96 \pm 37 a
Winter	20	290 \pm 60 b	29 \pm 2 b	51 \pm 16 b
Total mean	60	311 \pm 56	26 \pm 5	79 \pm 35
Comparison among seasons	60	H ₍₂₎ = 41.721, $p < 0.001$	F _(2,57) = 4.024 $p = 0.023$	F _(2,57) = 25.181 $p < 0.001$
<i>Mugil cephalus</i>				
Summer	20	811 \pm 74 a	33 \pm 4 a	87 \pm 13 a
Autumn	20	1003 \pm 511 a	46 \pm 3 b	43 \pm 17 b
Winter	20	386 \pm 78 b	36 \pm 2 a	81 \pm 25 a
Total mean	60	733 \pm 394	38 \pm 7	71 \pm 27
Comparison among seasons	60	H ₍₂₎ = 33.446 $p < 0.001$	H ₍₂₎ = 29.806 $p < 0.001$	H ₍₂₎ = 31.639 $p < 0.001$

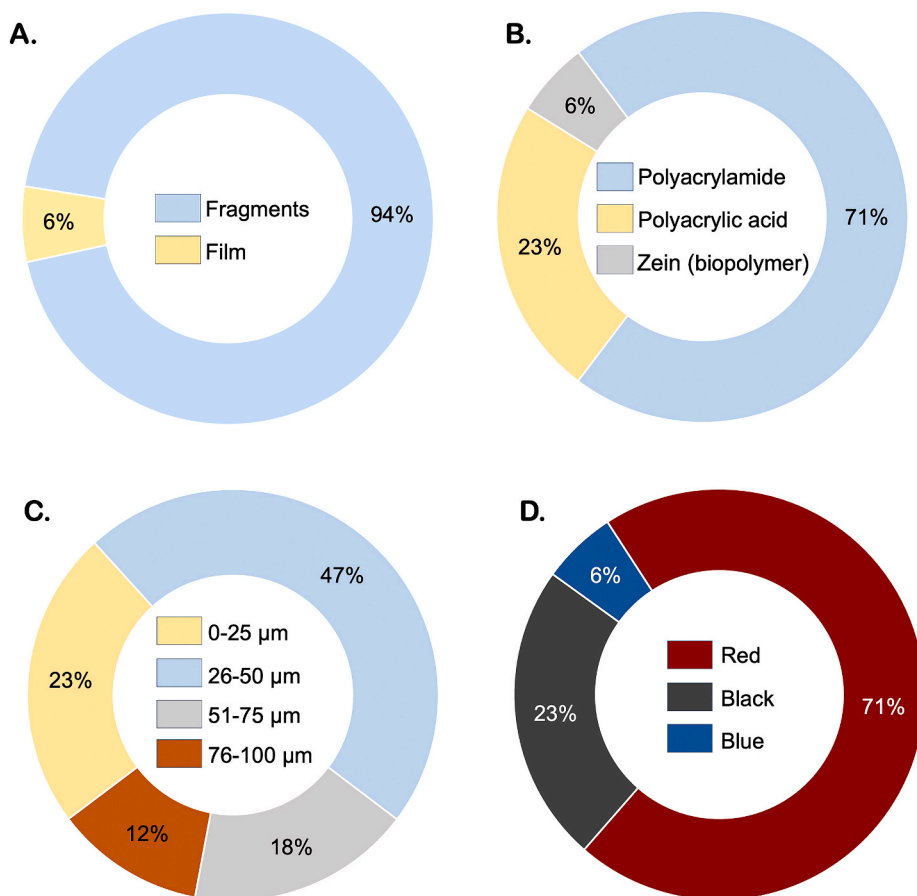


Fig. 3. Characteristics of microplastics recovered from brain samples of wild fish. (A-) shape; (B) polymer composition; (C) size class; and (D) colour. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

4.1. MPs were found in brain samples of fish

MPs were found in brain samples of *D. labrax* fish collected in the summer, leading to the refusal of H_{01} and acceptance of H_{A1} . The presence of MPs in brain samples indicates that fish have uptake MPs, which entered into the blood circulation, crossed the blood-brain barrier and entered into the brain where they were accumulated or at least retained for some time. The presence of NPs and MPs in the brain was documented before in fish and crabs exposed in laboratorial conditions. In fish, particles in the brain were found in Crucian carps (*Carassius carassius*) exposed for 64 days to amino-modified polystyrene NPs with 53 or 180 nm [25], in Nile tilapia (*Oreochromis niloticus*) exposed for 14 days to polystyrene MPs with 0.1 µm [20], 0.3, 5 or 70–90 µm [70], and in zebrafish (*Danio rerio*) exposed for 96 h to polystyrene NPs with 100 nm [84] or after ~7 weeks to polystyrene NPs with 70 nm [85]. In crabs, plastic particles entered and remained within the brain of the velvet swimming crab (*Necora puber*) after 1 h, 24 h, 7 days and 21 days post consumption of mussels contaminated with 0.5 µm polystyrene MPs [86]. Our study shows that this happens also in real scenarios, in agreement with the prior report of MPs in the brain of wild fish from the Black Sea [65].

Our results show that from all the three sampling seasons and studied species, MPs were only found in one species, in the summer period. Such difference could be driven by contrasting seasonal MP abundance and distribution patterns in the Douro estuary [52] or by differences in the feeding features and habitat preferences of the studied fish species [87]. Given these results, further research is needed to understand the factors that control and influence the presence of MP particles in the brain of wild fish populations.

The high observed proportion (70%) of small MP particles (<50 µm in size) in the brain of *D. labrax* specimens indicates that the smaller the particle size is, the easier it is for MPs to penetrate the blood-brain barrier. Our findings also relate to previous research highlighting that smaller MP particles are seemingly able to be retained in the brain of fish more readily than larger particles (70 – 90 µm) [70], although MPs up to 200 µm in size were also found in the brain of fish species caught in the Black Sea [65].

In addition to the size, the internalization of MPs into cells and tissues has been also related to other characteristics, such as shape, surface chemistry, softness and smoothness of the material [88]. Therefore, the results of the present pilot study showing that irregular shaped MPs, such as fragments and films, can reach the brain in fish exposed in real scenarios increases the concerns regarding the long-term exposure of wildlife and humans to a variety of MPs present in the environment.

The mechanisms allowing the MPs to cross the blood-brain barrier are not known. Several mechanisms were proposed, such as the

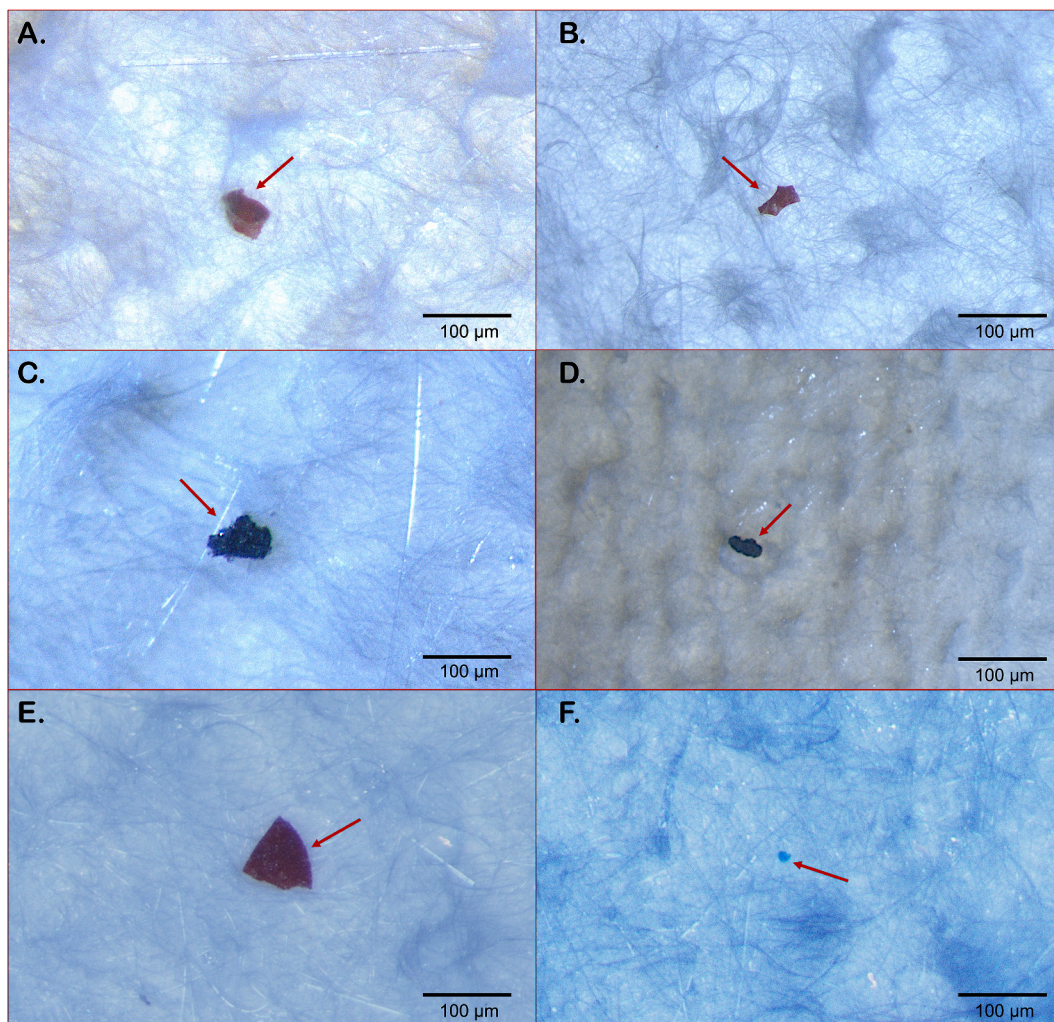


Fig. 4. Examples of microplastics recovered from brain samples of *Dicentrarchus labrax* from the Douro estuary (Portugal). (A–E) fragments; (F) film.

disturbance of the tight junction of the blood-brain barrier facilitating MPs entry [89], microglial phagocytosis [90], and via retrograde transport through olfactory nerve endings [32]. Independently of the mechanisms involved, the presence of MPs in the brain of fish can cause several adverse effects, such as neuronal degeneration and necrosis, cytoplasmic vacuolation, inflammatory cell infiltration and haemorrhage [91], decreased brain mass and morphological changes in the cerebral gyri [25], alterations in neural circuits [71] and alterations of brain enzymes [70], among other adverse effects as recently reviewed [32]. Further studies need to be performed to improve our understanding about the mechanisms involved in the internalization and retention of MPs (including bioplastics) in the brain of fish and other animals, and the influence of several factors, including biological variables (e.g., species, life-cycle phase), properties of the MPs (surface chemistry, size, shape), and environmental determinants (MP concentration, temperature variation, presence of other stressors).

The polymer composition of the MPs found in the brain of fish from the Douro estuary was dominated by two acrylic polymers (polyacrylamide (PAM) and polyacrylic acid (PAA)), followed by one biopolymer (Zein). Considering that the seasonal distribution of MPs in the environment is likely more related to specific plastic types than to the overall plastic concentration [92], one can hypothesise that the predominance of acrylic polymers in the brain of *D. labrax* specimens in the summer can be related with the increasing demand for shipping activities during the summer period (e.g., boat tours) or with the seasonal cleaning of the boats in the boat dock near the area where the fish were captured. These hypotheses can be supported by reports indicating that the presence of acrylic polymer fragments in aquatic environments are normally generated during boat maintenance and cleaning [93,94] or in areas of intense maritime traffic and related activities [95]. In addition, previous studies from the Douro estuary provide supporting evidence of micro-sized acrylic polymers fragments presence near a boat dock/maintenance area [53]. The presence of Zein within fish brain also deserves attention, because it is a main component of bioplastics [96,97]. Products manufactured from Zein can be used to package and/or protect food stuffs [98]. The presence of Zein in fish from the Douro estuary is in agreement with the findings in commercial fish species from the Adriatic Sea [97]. These findings draw attention to the new plastic materials that are now being used,

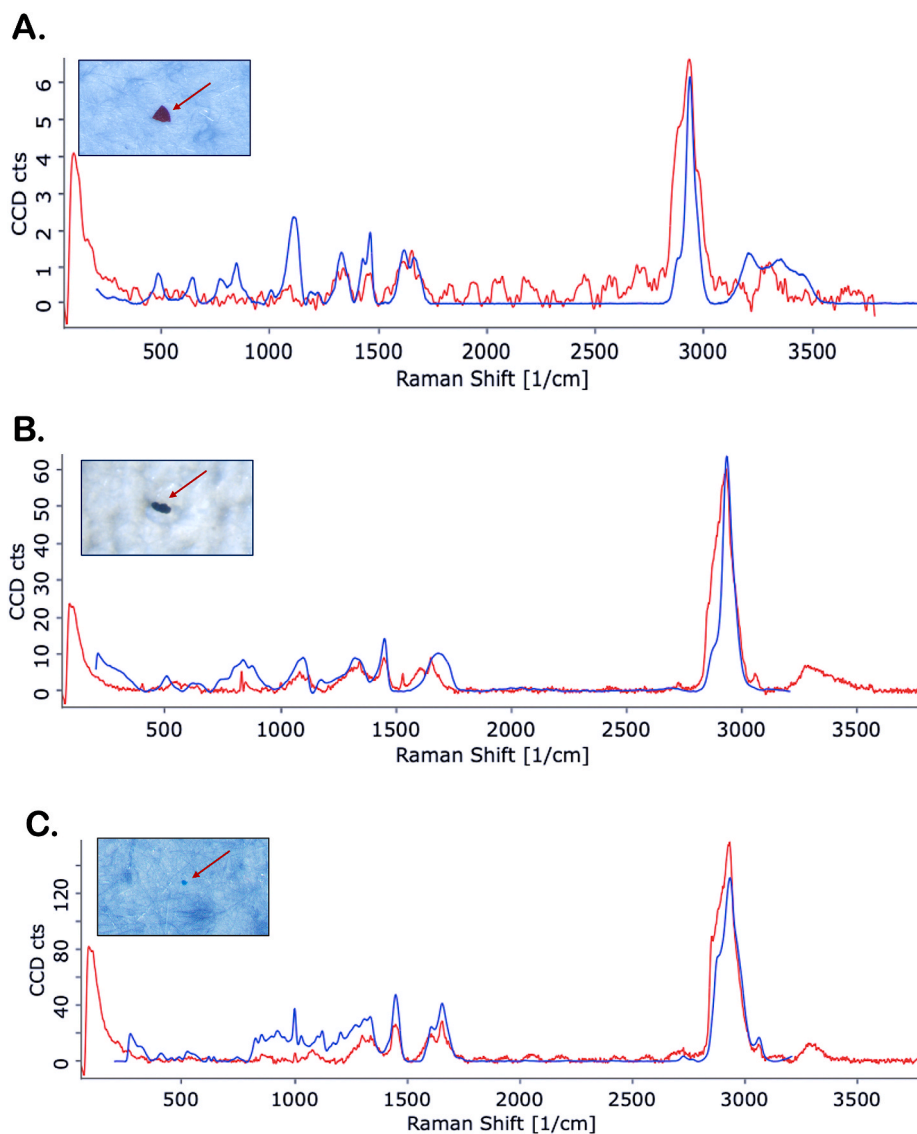


Fig. 5. Representative spectra of the microplastic particles polymers found in brain samples of wild fish. (A) Polyacrylamide; (B) Polyacrylic acid; (C) Zein (biopolymer).

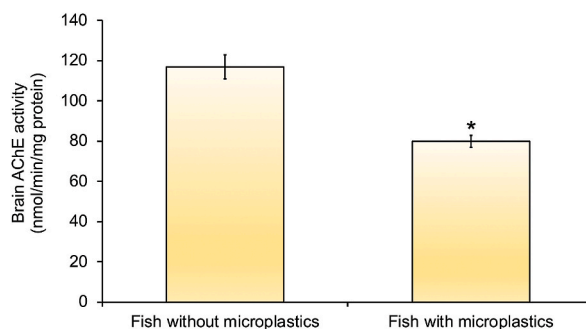


Fig. 6. Acetylcholinesterase activity (AChE) in brain of *Dicentrarchus labrax*, in groups of fish with ($n = 9$) and without ($n = 11$) microplastics in the brain captured during the summer period. Results are expressed as mean values \pm standard errors for each group of fish. * indicates statistical significant differences between groups of fish with and without microplastics (Student's t -test, $p \leq 0.05$).

and the need of more studies on their potential adverse effects.

4.2. Brain AChE activity is related with the presence of MPs in fish brain samples

In the summer, *D. labrax* specimens having MPs in the brain had significantly lower brain AChE activity than specimens of the same species where MPs were not found in the part of the brain analysed. The Douro estuary is contaminated with a diversity of MPs all through the year, and the contamination is considerable including in the summer [52]. All the specimens of *D. labrax* used for the statistical analyses comparing AChE activity in fish with and without MPs in brain samples were collected in the same estuary minimizing the potential influence of previous developmental conditions in different habitats, which may influence some of the MP effects [33,99]. All the fish with MPs in brain samples were collected in the same season (summer), therefore minimizing the potential influence of seasonal variability and other contaminants that may inhibit AChE activity (e.g., some metals, PAHs, organophosphate pesticides) known to be present in the Douro estuary [99,100]. Moreover, in studies carried out in laboratory conditions, AChE inhibition has been documented in several fish species exposed to MPs with a variety of sizes, chemical composition and shape. Just to give some examples, inhibition of brain AChE activity was documented in *Pomatoschistus microps* from wild populations exposed to 1–5 μm polyethylene beds [26,33], in *Argyrosomus regius* exposed to 125 μm irregular low-density polyethylene [34], in *Oreochromis niloticus* exposed to 0.1 μm polystyrene beads [20], in *Symphysodon aequifasciatus* exposed to 70–88 μm polyethylene microspheres [27], in *D. labrax* exposed to 1–5 μm beds with unknown polymer composition [29], in *Clarias gariepinu* exposed to polyvinyl chloride pellets (3.0 g/kg) [28], and in *Etroplus suratensis* from wild populations exposed to 80.72 μm polyvinyl chloride pellets [101]. Furthermore, it is important to highlight that acrylamide, widely employed as a monomer used in the polymer industry, including to produce polyacrylamide, one of the MP types identified in the brain samples of *D. labrax* from the Douro estuary, can cause AChE inhibition, alterations in neurotransmitter levels among other effects in the central nervous system [102–105]. Therefore, the lower AChE activity (~32%) in fish with MPs in the brain than in fish with samples negative for MPs lead to the refusal of H_2O_2 and acceptance of H_2A_2 .

Brain AChE inhibition in fish with MPs points to neurotoxicity induced by long-term exposure to MPs, MP-Chem or both. These findings are in line with previous studies [32]. It may cause a diversity of effects, some of them also documented in fish exposed to MPs in controlled conditions, such as alterations in behavioral patterns, swimming performance decrease, and reduction of feeding activity [27,35,106]. Inhibition of AChE may also influence key neurodevelopmental events, including cell migration, neurite outgrowth, synaptogenesis, among others [32,91]. This highlights the importance of further research on the neurotoxic effects induced by long-term exposure to MPs and also MP-Chem.

5. Conclusions

Among the 180 wild fish (*D. labrax*, *P. flesus*, *M. cephalus*) analysed, 9 had MPs in brain samples (all the brain except the cerebellum that was used for AChE analyses). All the fish with MPs in brain samples were *D. labrax* specimens collected in the summer, accounting for 45% of the 20 seabasses collected in the summer. These findings demonstrate the ability of MPs to cross the blood barrier of fish exposed in a real scenario (Douro River estuary discharging into the NE Atlantic Ocean), likely for a considerable or long time, to a diversity of MPs and other contaminants. Seventeen MPs (16 fragments, 1 film) with sizes between 8 μm and 96 μm were recovered from the analysed brain samples. Two superabsorbent polymers and one biopolymer were identified. Compared with *D. labrax* specimens collected in the summer with brain samples negative for MPs, fish with MPs in brain samples had significant inhibition (by 32%) of brain AChE activity pointing to neurotoxicity induced by MPs. The findings of this pilot study carried out in a real scenario together with previous studies in the laboratorial and field conditions highlight the potential neurotoxicity that long-term exposure to MPs, MP-Chem and other pollutants may have been inducing on wild species, domestic animals and humans, and stress the urgent need of further studies on the topic using ‘One Health’ approaches.

Author contribution statement

L.G.A. Barboza: conceived and designed the experiments; performed the experiments; analysed and interpreted the data; wrote the paper; original draft of the manuscript.

X.L. Otero: contributed reagents, materials, analysis tools or data; analysed and interpreted the data; wrote the paper; drafting the article or revising it critically for important intellectual content.

E.V. Fernández: contributed reagents, materials, analysis tools or data; analysed and interpreted the data.

L.R.Vieira: performed the experiments; analysed and interpreted the data; wrote the paper; drafting the article or revising it critically for important intellectual content.

S.C. Cunha: conceived and designed the experiments; contributed reagents, materials, analysis tools or data; wrote the paper; drafting the article or revising it critically for important intellectual content.

J.O. Fernandes: conceived and designed the experiments; contributed reagents, materials, analysis tools or data; wrote the paper; drafting the article or revising it critically for important intellectual content.

L. Guilhermino: conceived and designed the experiments; analysed and interpreted the data; contributed reagents, materials, analysis tools or data; wrote the paper; drafting the article or revising it critically for important intellectual content.

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Data availability statement

Data will be made available on request.

Declaration of interest's statement

The authors declare no competing interests.

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