



Are we underestimating microplastic abundance in the marine environment? A comparison of microplastic capture with nets of different mesh-size[☆]



Penelope K. Lindeque^{a,*,1}, Matthew Cole^{a,1}, Rachel L. Coppock^a, Ceri N. Lewis^b, Rachael Z. Miller^d, Andrew J.R. Watts^b, Alice Wilson-McNeal^{a,b}, Stephanie L. Wright^c, Tamara S. Galloway^b

^a Plymouth Marine Laboratory, Prospect Place, Plymouth, PL1 3DH, UK

^b College of Life and Environmental Sciences, University of Exeter, Exeter, EX4 4PS, UK

^c MRC-PHE Centre for Environment and Health, Analytical, Environmental and Forensic Sciences, King's College London, London, SE1 9NH, UK

^d Rozalia Project, PO Box 3075, Burlington, VT, 05408, USA

ARTICLE INFO

Article history:

Received 11 February 2020

Received in revised form

30 April 2020

Accepted 1 May 2020

Available online 3 May 2020

Keywords:

Plastic

Microplastics

Pollution

Ocean

Net

Sampling

ABSTRACT

Microplastic debris is ubiquitous and yet sampling, classifying and enumerating this prolific pollutant in marine waters has proven challenging. Typically, waterborne microplastic sampling is undertaken using nets with a 333 μm mesh, which cannot account for smaller debris. In this study, we provide an estimate of the extent to which microplastic concentrations are underestimated with traditional sampling. Our efforts focus on coastal waters, where microplastics are predicted to have the greatest influence on marine life, on both sides of the North Atlantic Ocean. Microplastic debris was collected via surface trawls using 100, 333 and 500 μm nets. Our findings show that sampling using nets with a 100 μm mesh resulted in the collection of 2.5-fold and 10-fold greater microplastic concentrations compared with using 333 and 500 μm meshes respectively ($P < 0.01$). Based on the relationship between microplastic concentrations identified and extrapolation of our data using a power law, we estimate that microplastic concentrations could exceed 3700 microplastics m^{-3} if a net with a 1 μm mesh size is used. We further identified that use of finer nets resulted in the collection of significantly thinner and shorter microplastic fibres ($P < 0.05$). These results elucidate that estimates of marine microplastic concentrations could currently be underestimated.

© 2020 The Authors. Published by Elsevier Ltd. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

1. Introduction

Microplastics are a prolific, persistent and pernicious contaminant, posing an environmental and economic risk to marine ecosystems across the globe (Rochman et al., 2016). Microplastics, encompassing synthetic plastic particulates, fibres and films, here defined as 1–5000 μm in diameter, have been widely identified in marine ecosystems, including estuaries, coastal biomes, the open ocean and polar waters (Lusher, 2015). Microplastics are either directly manufactured (e.g. cosmetic exfoliates, air blasting media),

or derive from the fragmentation of larger plastics over time (Cole et al., 2011). By design, plastics are resistant to degradation and as such are expected to persist in the natural environment for hundreds, if not thousands of years (Andrady, 2015). Owing to their small size, microplastics are bioavailable to a range of organisms across trophic levels, including zooplankton (Steer et al., 2017), bivalves and fish destined for human consumption (Rochman et al., 2015), and marine megafauna (Duncan et al., 2019; Nelms et al., 2019). Exposure studies have highlighted the negative impacts microplastic ingestion can have on marine organisms, including copepods, shellfish, benthic invertebrates and fish, with effects comprising reduced feeding, fecundity, growth and survival, premature moulting, altered behaviour and shifts in ecological functionality (Besseling et al., 2013; Cole et al., 2019; Cole et al., 2015; Cole et al., 2016; Sussarellu et al., 2016; Wegner et al., 2012; Wright et al., 2013). However, it is currently unclear whether such adverse

[☆] This paper has been recommended for acceptance by Eddy Y. Zeng.

* Corresponding author.

E-mail address: pkw@pml.ac.uk (P.K. Lindeque).

¹ P.K.L. and M.C. contributed equally to this work.

health effects are likely to occur in the natural environment due to the mismatch between the size, type and concentration of microplastics that are traditionally sampled during environmental monitoring studies and those used in exposure studies (Burns and Boxall, 2018). At present, the concentration of bioavailable microplastics in the natural environment, a similar size to natural prey and a similar size to those used in effect studies, is relatively unknown (de Sá et al., 2018).

To comprehensively assess the risks that microplastic debris poses to marine ecosystems requires robust estimates of the size, prevalence and distribution of microplastic within the global ocean. However, accurately quantifying and characterising microplastic debris within environmental samples, and subsequently modelling this data, has proven hugely challenging. Microplastics research is still in its infancy, and over the past decade there has been a multitude of methodological approaches applied when sampling, extracting and identifying microplastic debris, with samples taken from different ecological compartments (i.e. sediments, water column, biota) each providing their own unique challenges (Lusher et al., 2016; Stock et al., 2019). Thus far, field sampling has predominantly focussed on the subtropical gyres of the northern hemisphere, with data gaps for large swathes of the open ocean, the southern hemisphere, equatorial regions and coastal waters (Clark et al., 2016). One of the most widely applied methods for collecting microplastics at the sea surface has been to conduct trawls using 330–335 μm nets, hereafter referred to as 333 μm , which have traditionally been used for sampling zooplankton (Hidalgo-Ruz et al., 2012; Lusher et al., 2016). Such environmental data has been used to derive initial estimates of oceanic microplastic budgets: for example, van Sebille et al. (2015) estimates that the accumulated number of microplastic particles in 2014, ranged from 15 to 51 trillion particles, weighing between 93,000 and 236,000 metric tons, with >90% of observations collected using a Manta or Neuston net with 333 μm mesh. A recent review highlighted that over 80% of field studies only sample microplastics >300 μm , and as such microplastics smaller than this size, including 95% of cosmetic microbeads, synthetic microfibres and secondary microplastics with diameters <300 μm , will be absent from datasets (Conkle et al., 2018). As such, we hypothesise current estimates of microplastic pollution at the sea surface are likely to be underestimated.

In this study, we determine the relationship between net mesh size and the abundance and character of captured microplastic, providing an estimate of the extent to which microplastic concentrations may be underestimated using 333 μm nets. Our sampling efforts focus on biologically productive coastal waters on both sides of the North Atlantic (i.e. Gulf of Maine and western English Channel), close to land-based and maritime sources of pollution, where microplastics are predicted to have the greatest influence on marine life (Clark et al., 2016). Microplastic debris was collected via sub-surface trawls using 100, 333 and 500 μm nets to compare microplastic concentrations sampled with nets of differing mesh sizes. The study aims to provide a greater resolution in the determination of global microplastic budgets, allowing for the risk of microplastic debris to marine ecosystems to be more clearly defined.

2. Materials and methods

2.1. Environmental sampling

Field sampling was conducted on both sides of the North Atlantic Ocean, focusing on coastal waters of the Gulf of Maine (USA) and the western English Channel (UK). In all cases, sub-surface sampling focused upon the comparison of microplastic

concentrations collected by nets towed in parallel. For our US sampling, the use of a sailing vessel limited us to using a maximum of two nets at a time, comprising either two 333 μm nets or a 100 and 500 μm net. For our UK sampling, the use of the RV Quest (Maritime and Coastguard Agency Category 2 workboat) allowed 100, 333 and 500 μm nets to be towed in parallel.

2.1.1. Gulf of Maine (USA)

Fieldwork was conducted throughout July 2013 in the Gulf of Maine (USA), with sampling targeted at sites of upwelling and riverine output around Hurricane Island, Boothbay Harbor, Portland, Kittery, Star Island and Boston (Fig. 1; Table S1). Sampling was conducted on-board the RV *American Promise*, with nets deployed from the spinnaker pole to capture sub-surface debris outside of the vessel's wake; nets were maintained half in and half out of the water. Each trawl (250 m transects; 0.7–2.8 knots) used two nets towed in parallel, comprising either: two 333 μm Neuston nets (0.5 m² aperture; rectangular, 1 m × 0.5 m); or 100 μm and 500 μm plankton nets (0.2 m² aperture; circular, 0.5 m ϕ). The nets and cod-ends were thoroughly rinsed down, and samples transferred onto clean nylon mesh of corresponding size. Any large pieces of flotsam (e.g. wood, macroalgae) were rinsed with freshwater to remove adhered microplastics, and then removed from the sample. Meshes were rinsed with freshwater and then folded and secured to retain samples and minimise contamination. Adapting the protocols of Moore et al. (2002), samples were desiccated at 60 °C overnight in a food-dehydrator, and stored in sample bags in a desiccating chamber prior to analysis.

2.1.2. English Channel (UK)

Fieldwork was conducted in the western English Channel off the coast of Plymouth (UK) between July and September 2015 (Fig. 1; Table S2). Sub-surface sampling was conducted on board the RV *Plymouth Quest* using three Neuston nets (100, 333 and 500 μm ; 0.2 m² aperture; circular, 0.5 m ϕ) rigged in parallel and trawled off the beam of the boat (500 m trawl; 0.5–1.5 knots) to avoid downwelling of the debris in the vessel's wake; nets were maintained half in and half out of the water. Each net and cod end were rinsed into a clean bucket with surface seawater collected using the boat's intake system. Any large pieces of flotsam (e.g. wood, macroalgae, feathers) were rinsed with filtered seawater (0.2 μm) to remove adhered microplastics, and then removed from the sample. The bucket contents were poured through a nylon mesh matching the mesh size of the net and rinsed with filtered seawater (0.2 μm). Meshes were folded and secured and then temporarily wrapped in aluminium foil during transit to avoid contamination. Samples were stored at –80 °C and subsequently freeze-dried prior to analysis.

2.2. Enzymatic digestion

To reveal any microplastics obscured by biotic material within the samples, we employed enzymatic digestion per the protocols of Cole et al. (2014). Samples were transferred individually into a pre-cleaned porcelain mortar and the weight of the pestle was used to gently break down large structures. Each sample was weighed, transferred to an acid-washed glass vial, and homogenising solution added at a ratio of 15 mL to 0.2 g dry weight sample. Samples were physically homogenised using a 19G needle and 10 mL syringe then incubated at 50 °C in an orbital shaker at 100 rpm for 30 min. Proteinase K was added to a concentration of 500 $\mu\text{g mL}^{-1}$, and samples incubated at 50 °C again at 150 rpm for 2 h. Digested samples were visually examined, and any still containing large quantities of organic material were incubated for a further 2 h. Sodium perchlorate (5 M) was then added and each sample

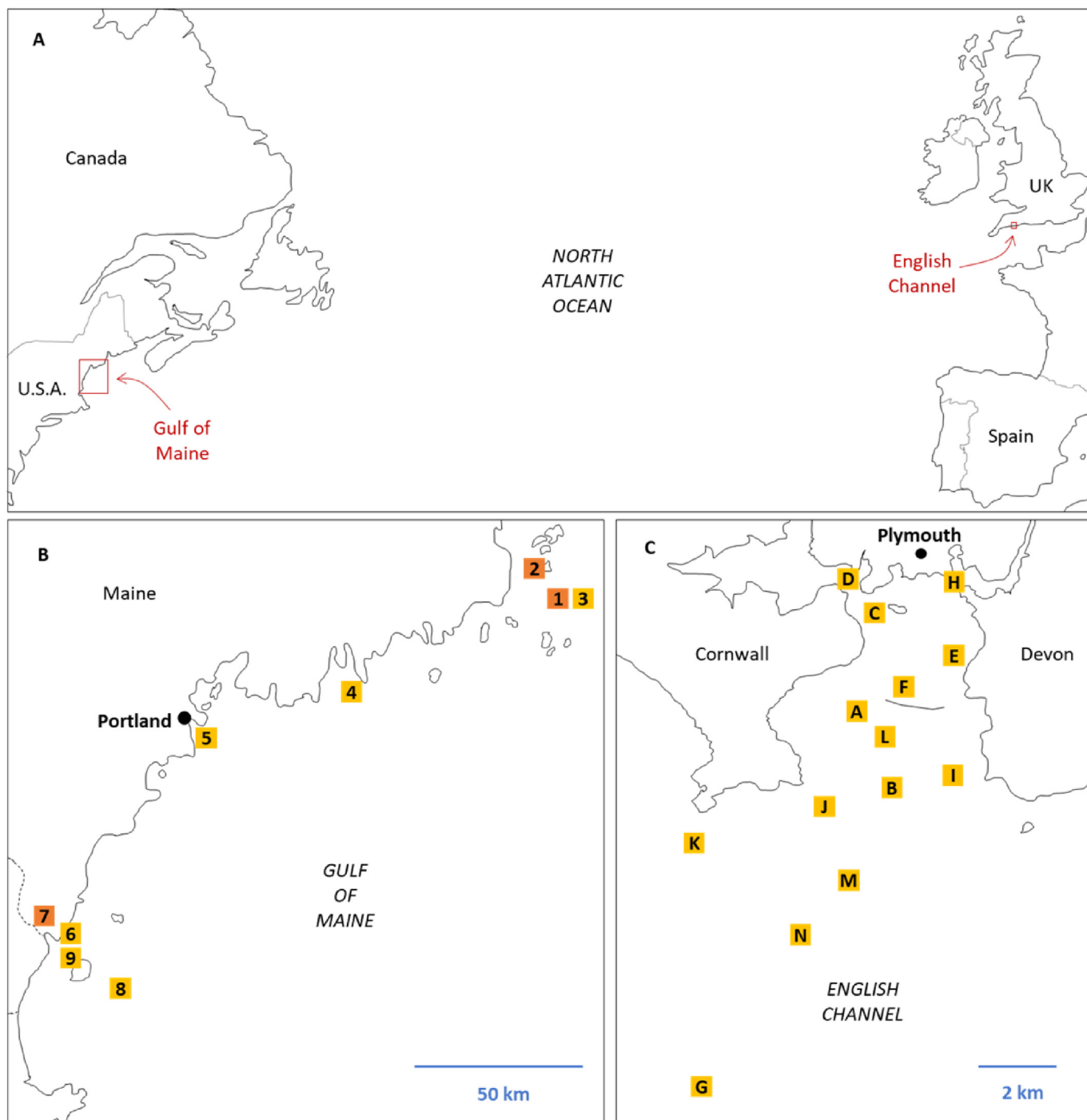


Fig. 1. Charts showing locations of sampling sites. (A) North Atlantic Ocean, noting locations of the Gulf of Maine and English Channel. (B) North-eastern US seaboard, relative to Portland (ME), with 50 km scale; yellow boxes denote sites where samples were taken using 100/500 μm nets and 333/333 μm nets, and orange boxes denote where samples were taken using 100/500 μm nets only. (C) Plymouth Sound and western English Channel, with 2 km scale; yellow boxes denote sites sampled with 100/333/500 μm nets. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

homogenised using a 21G needle before mixing at 150 rpm at room temperature for 20 min. Finally, samples were incubated at 65 °C for a further 20 min. Digested samples were vacuum filtered through 50 μm nylon mesh filters. Samples containing large volumes of material were sub-divided over multiple meshes. All samples were treated identically, irrespective of net size.

2.3. Characterisation

Per the proposed categorisation framework of Hartmann et al. (2019), we look to characterise microplastics by their chemical composition, size, shape and colour. Mesh filters were systematically analysed under a dissection microscope (Olympus SZX16; x40–100 magnification), using a sterilised needle to tease apart the

sample. Suspected microplastics were visually identified by their uniformity, colour and form per the guidance of Norén et al. (Norén, 2007). The shape (fibre, fragment or sphere) and colour of all particles was recorded immediately. Owing to the large number of particles present, for each sample 15 particles were randomly selected for sizing and polymeric analysis. Particles were randomly selected by: (1) dividing the mesh into 9 (3 rows x 3 columns); (2) using a random number generator (Microsoft Excel) to determine which section to first select a microplastic from; (3) 15 particles were picked from this first section; (4) where <15 particles were available, a binary random number was used to determine which section to next sub-sample from (i.e. go sequentially up or down through the grid). Sizing was conducted using CellSens software and light microscope (Olympus SX16) with two-dimensions recorded. Polymeric analysis was conducted on randomly selected particles using either Attenuated Total Reflectance Fourier Transform Infrared spectroscopy (Bruker Alpha ATR-FTIR) or micro ATR (μ ATR) in Reflectance mode (PerkinElmer Spotlight 400 FTIR). Owing to the limitations of the Bruker ATR-FTIR, the particles identified using this instrument ($n = 355$) required one dimension >100 μm for spectral analysis, the remainder of selected particles analysed (PerkinElmer, $n = 416$) required a minimum dimension of 11 μm . Spectra were analysed using OPUS 6.5 software (Bruker) and Spectra software (PerkinElmer). Spectra showing no defined peaks (i.e.; <60% match) were dismissed, otherwise particles were classified as either 'natural' (e.g. chitin, cellulose), or 'microplastic', with further sub-division by polymer: acrylic, polyamide, polyester, polyethylene, polypropylene, polyvinylchloride, biopolymer (e.g. rayon), elastomer (e.g. neoprene, rubber), or other (i.e. copolymers, polystyrene).

2.4. Quality control

Prior to fieldwork and analysis, all participants were instructed on minimising sample contamination via atmospheric deposition, clothing and equipment. During sample collection, nets were trawled to the side of the research vessel to avoid any paint or material from the boat contaminating the sample. Samples were handled by personnel wearing cotton clothing and latex gloves, and procedural blanks using filtered seawater were conducted at each sampling station on each cruise to account for contamination. Samples were enclosed in meshes and stored in sealable containers prior to analysis. To minimise contamination in the laboratory, all analyses were conducted by trained researchers. Further, samples were covered wherever feasible, glassware was used in place of plastic where possible, and all reusable equipment was cleaned thoroughly with ethanol and rinsed twice with Milli-Q water (0.2 μm filtered) prior to use. Sample processing was conducted in positive-pressure (i.e. laminar flow) hoods to prevent airborne contamination. Procedural blanks ($n = 14$ for UK samples; $n = 6$ for US samples), containing no sample, but otherwise treated as per the given methodology, were used to quantify contamination of samples during processing.

2.5. Microplastic concentrations

The waterborne concentration of microplastics (microplastics m^{-3}) from each net at each site was calculated using our data adjusted for volume sampled, contamination and mis-identification. The mean number of particles identified in the procedural blanks was subtracted from the total number of particles picked out from each sample; this data was then adjusted to account for the proportion of particles confirmed as plastic following FT-IR. The approximate volume of water sampled (m^3)

was calculated by multiplying 50% of the net aperture (m^2), noting nets were half submerged, by length of tow measured as distance (m) over the ground (therefore taking boat speed and tidal stream into consideration), assuming a 95% sampling efficiency (Skjoldal et al., 2013).

2.6. Statistical analyses

Statistical analyses were conducted using R statistical software v3.5.1 (R Core Team, 2019). Normality of data was tested using the Shapiro-Wilk test, and non-parametric data log transformed where applicable. Comparisons between datasets were assessed using a student's t -test or ANOVA with post-hoc Tukey test, or a Kruskal-Wallis test for non-parametric data. Significant difference is attributed where $P < 0.05$. A power law regression analysis was conducted using pooled mean microplastic concentrations across all UK sites for each net size.

3. Results

3.1. Environmental data

3.1.1. Gulf of Maine (USA)

In total 2,755 particles were isolated from the 100, 333 and 500 μm net samples taken from 9 sites along the coast of the Gulf of Maine. The samples predominantly consisted of fibres (84%), with a smaller quantity of fragments identified (16%); only 12 beads were observed (Fig. 2A). Fibres ranged from 5 to 282 μm in diameter and from 164 μm to >13 mm in length; the diameter of beads and fragments ranged from 57 to 3585 μm . The majority of fibres were black (62%), blue (15%), red (13%) or transparent (10%; Fig. 2B); fragments were predominantly blue (32%) or white/grey (24%), with an otherwise even distribution of colour (Fig. 2C). An ATR-FTIR analysis of a randomised sub-sample ($n = 254$, excluding particles providing a poor spectral signature) revealed that 85% of the isolated particles were 'microplastic', per the classification criteria set out by Hartmann et al. (2019) (Fig. 3A). Almost a third of the plastics identified were biopolymers (30%), of which the majority were rayon, with co-polymers (21%), polyethylene (13%) and polyester (13%) also well represented in the samples (Fig. 3B).

3.1.2. English Channel (UK)

In total 22,666 particles were isolated from the 100, 333 and 500 μm net samples taken from 14 sites in the western English Channel and Plymouth Sound. Across all samples, fibres (77%) were the most common, with smaller quantities of fragments (19%) and beads (4%) identified (Fig. 2A). Fibres ranged from 5 to 350 μm in diameter and from 55 μm to >8 cm in length; the feret diameter of beads and fragments ranged from 15 to 12,500 μm . Fibres were predominantly black (37%) or blue (32%), with substantial numbers of transparent (15%) and red (10%) filaments (Fig. 2B); the vast majority of fragments were blue (73%; Fig. 2C). Of the randomised sub-sample of isolated particles ($n = 517$, excluding particles providing a poor spectral signature), 94% were microplastic (Fig. 3A). The majority of these microplastics were made up of polyester (22%), biopolymers (22%), polypropylene (18%) and acrylic (14%). Also present in substantial quantities was polyethylene (9%) and polyamide (8%), with PVC (2%), elastomers (1%) and others (5%) making up the total (Fig. 3B).

3.1.3. Procedural blanks

Owing to the strict protocols in place, contamination of procedural blanks was relatively low. For the procedural blanks conducted alongside our Gulf of Maine analysis, we identified a mean of 1.5 particles per sample (89% fibres, 11% fragments). For

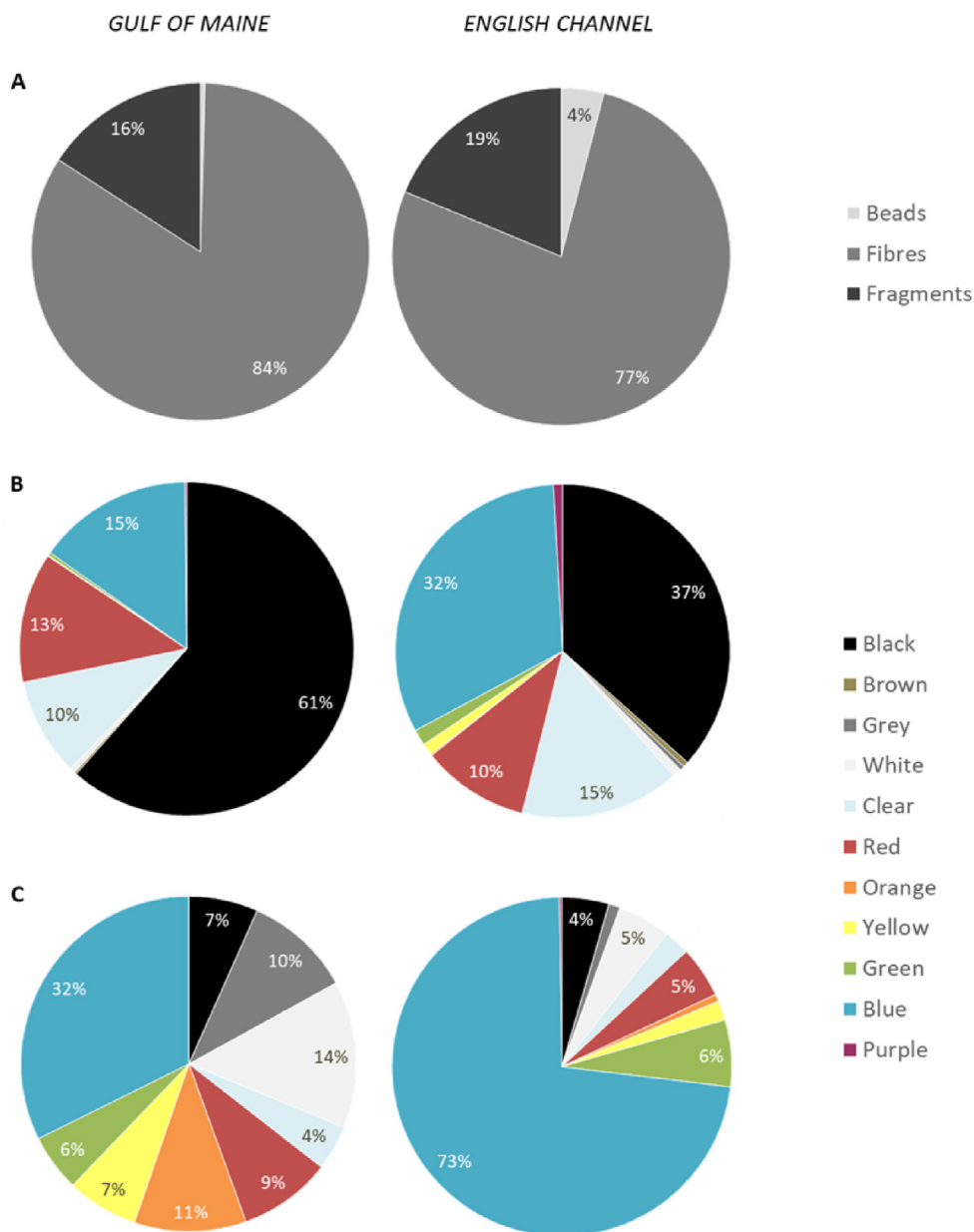


Fig. 2. Composition of particles identified in Gulf of Maine (left column; $n = 2,755$) and English Channel (right column; $n = 22,666$) samples. (A) Breakdown of particles by shape, i.e. fibres, fragments or beads. (B) Colour breakdown of fibres. (C) Colour breakdown of fragments. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

procedural blanks conducted in parallel with the English Channel sampling and analysis, we identified a mean of 9.4 particles per sampling station (75% fibres, 25% fragments).

3.2. Net comparisons

3.2.1. Gulf of Maine (USA): 333 μm nets

Average microplastic concentrations (mean \pm standard error) collected via two 333 μm nets, towed in parallel at five sites in the Gulf of Maine, were 0.54 ± 0.2 and 0.46 ± 0.3 microplastics m^{-3} , with no statistically significant difference in microplastic concentrations identified (t -test; $P = 0.406$; Fig. 4A). However, looking at individual site data (Fig. 4B), it is evident that there can be clear differences in microplastic concentrations collected using two nets towed in parallel (i.e. Site 5).

3.2.2. Gulf of Maine (USA): 100 and 500 μm nets

Based on parallel tows conducted at nine sites in the Gulf of Maine, we identified average microplastic concentrations of 6.03 ± 1.03 microplastics m^{-3} (100 μm net) and 0.60 ± 0.25 microplastics m^{-3} (500 μm net). On average, sampling with a 100 μm net revealed 10-fold higher microplastic concentrations compared with using a 500 μm net (t -test; $P < 0.001$; Fig. 5A). Highest microplastic concentrations, as sampled using a 100 μm net, were identified at Site 1 (Outer Penobscot Bay; 10.0 microplastics m^{-3} ; Fig. 5B).

3.2.3. English Channel (UK): 100, 333 and 500 μm nets

Sampling efforts across 14 sites in the western English Channel and Plymouth Sound revealed mean microplastic concentrations of 10.03 ± 2.21 microplastics m^{-3} (100 μm net), 4.08 ± 1.32

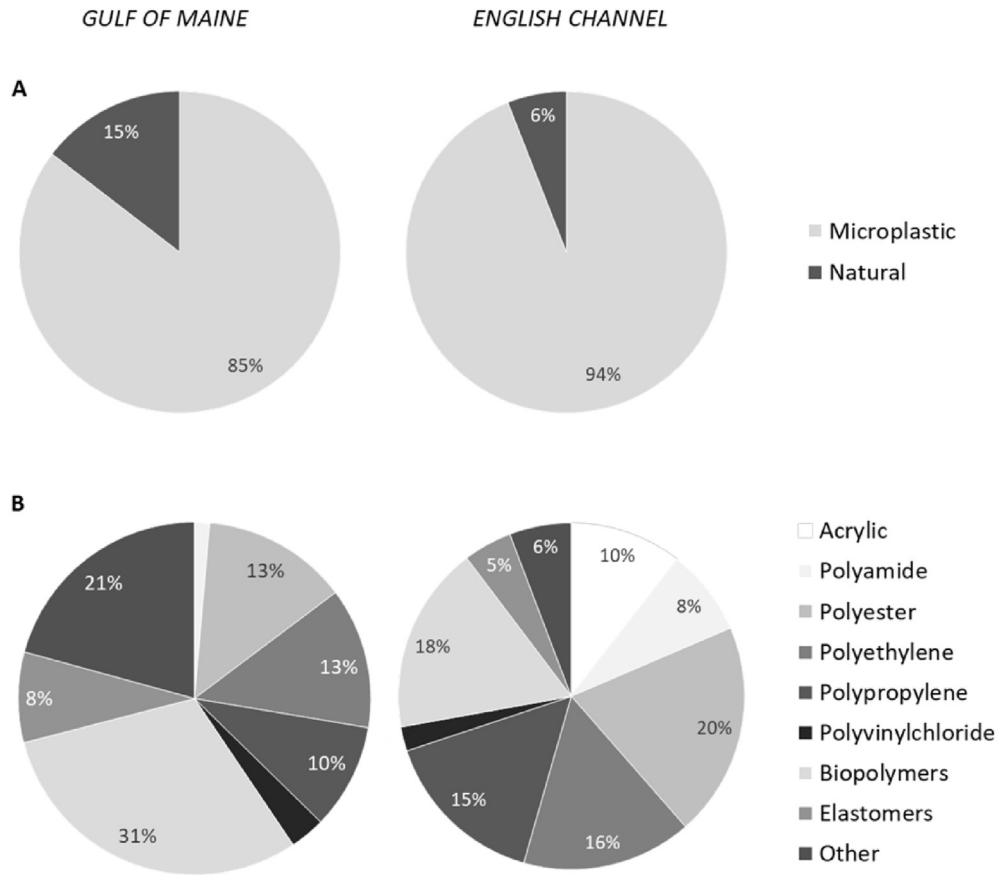


Fig. 3. Composition of particles picked-out in Gulf of Maine (left column; $n = 254$) and English Channel (right column; $n = 517$) samples. (A) Composition of material, i.e. naturally occurring or plastic. (B) Breakdown of plastics by polymer type, including biopolymers and elastomers.

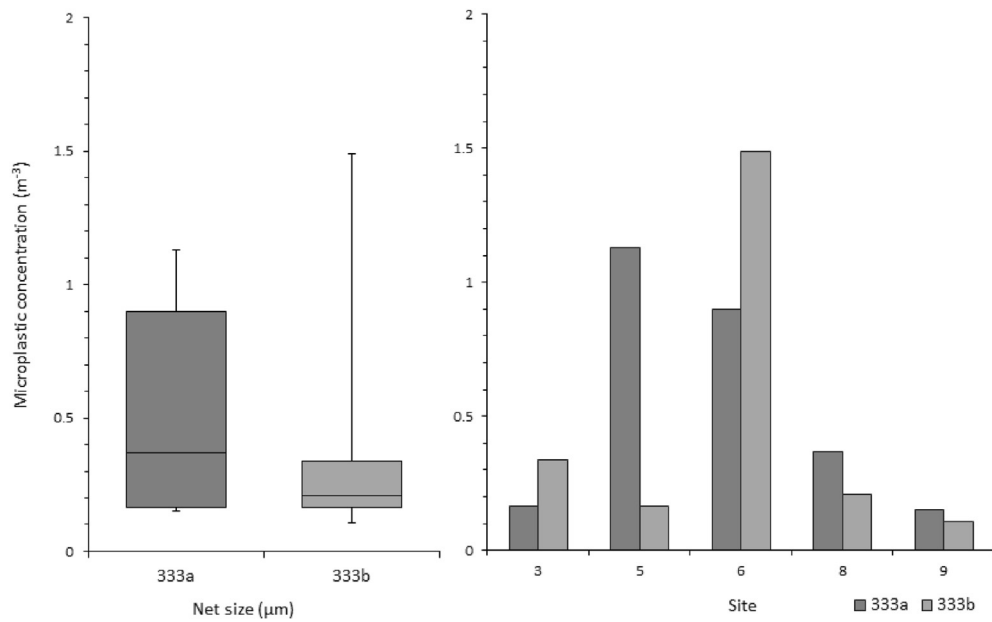


Fig. 4. Waterborne concentrations of microplastics (items m^{-3}) in the Gulf of Maine using two 333 μm nets towed in parallel. (A) Box and whisker plots showing median concentrations across sites and (B) bar chart displaying concentrations found at each site.

microplastics m^{-3} (333 μm net) and 1.03 ± 0.16 microplastics m^{-3} (500 μm net). Mesh size was a significant factor in resulting

microplastic concentrations (ANOVA, $P < 0.001$; Fig. 6A, displaying median and interquartile values), with no significant influence of

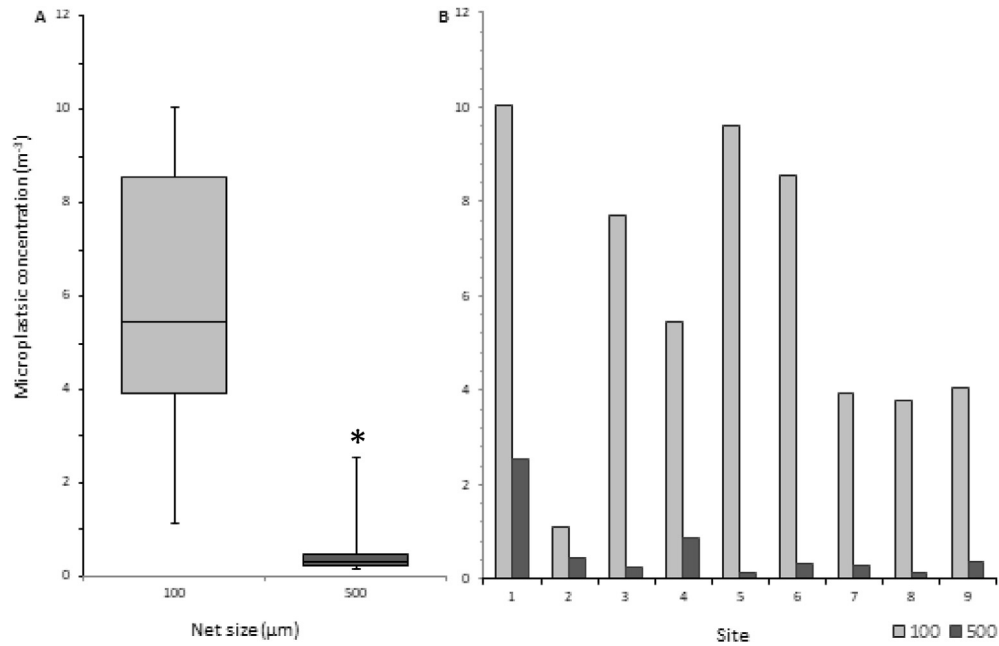


Fig. 5. Waterborne concentrations of microplastics (items m⁻³) in the Gulf of Maine using 100 µm and 500 µm nets towed in parallel; *denotes significant difference (*t*-test $p < 0.05$). (A) Box and whisker plots showing median concentrations across sites and (B) bar chart displaying concentrations found at each site.

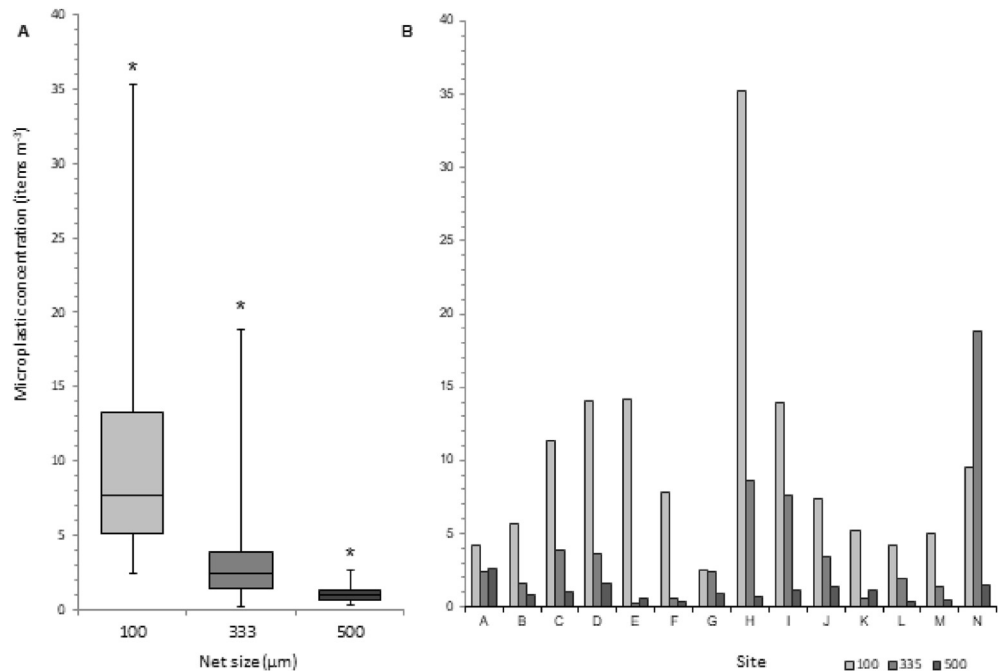


Fig. 6. Waterborne concentration of microplastics (items m⁻³) in the western English Channel, as sampled using 100, 333 or 500 µm nets. (A) Box and whisker plots showing median concentrations across sites; *denotes significant difference (ANOVA, $p < 0.05$). (B) Bar chart displaying microplastic concentrations for each net found at each site.

Site (ANOVA, $P = 0.79$). On average, a 100 µm net revealed 2.5-fold higher microplastic concentrations than using a 333 µm net (ANOVA, $P < 0.05$) and 10-fold greater microplastic concentrations than using a 500 µm net (ANOVA $P < 0.001$); using a 333 µm net resulted in sampling 4-fold greater microplastic concentrations as when a 500 µm net was employed (Tukey Post-hoc; $P < 0.05$). However, at some sites this trend was not apparent, for example: at Site N (Outside Breakwater 4, 7 km offshore; Fig. 6B) microplastic concentrations collected using a 333 µm net exceeded those

collected via 100 µm net by two-fold; and at Site A (seaward side of Plymouth breakwater) and Site K (Rame Head), use of a 500 µm net revealed marginally greater microplastic concentrations than collected via 333 µm nets. The highest waterborne microplastic concentration, collected using a 100 µm net, was found at Site H (mouth of the River Plym; 35.5 microplastics m⁻³).

Fibres captured with a 100 µm net were significantly shorter than those sampled with a 333 and 500 µm net, with a significantly smaller diameter than those sampled with a 500 µm net (Kruskal-

Wallis, $P < 0.05$; Fig. 7). Mean fragment/bead diameter was far greater in the 500 μm net samples (575 μm) than the 100 μm (121 μm) or 333 μm (133 μm) net samples, however these differences were not statistically significant (Kruskal-Wallis, 100v500, $P = 0.07$; 333v500, $P = 0.08$). Fibres were the dominant particle shape characterised across all nets, comprising 75% in the 100 μm net, 81% in 333 μm net and 83% in 500 μm net (Fig. 7). Beads were only observed in the 100 μm net whilst fragments made up the remaining particle shape across all nets. Blue, black, clear and red

were the predominant particle colours across all net sizes, recording similar concentrations in each net size. Extrapolation of mean microplastic concentrations from pooled data across all sites provided estimates of concentrations using different mesh sizes (Fig. 8), estimating a mean concentration of 11.4 microplastics m^{-3} when using a 100 μm mesh size, 207.1 microplastics m^{-3} with a 10 μm mesh and increasing to 3700 microplastics m^{-3} if using a 1 μm mesh.

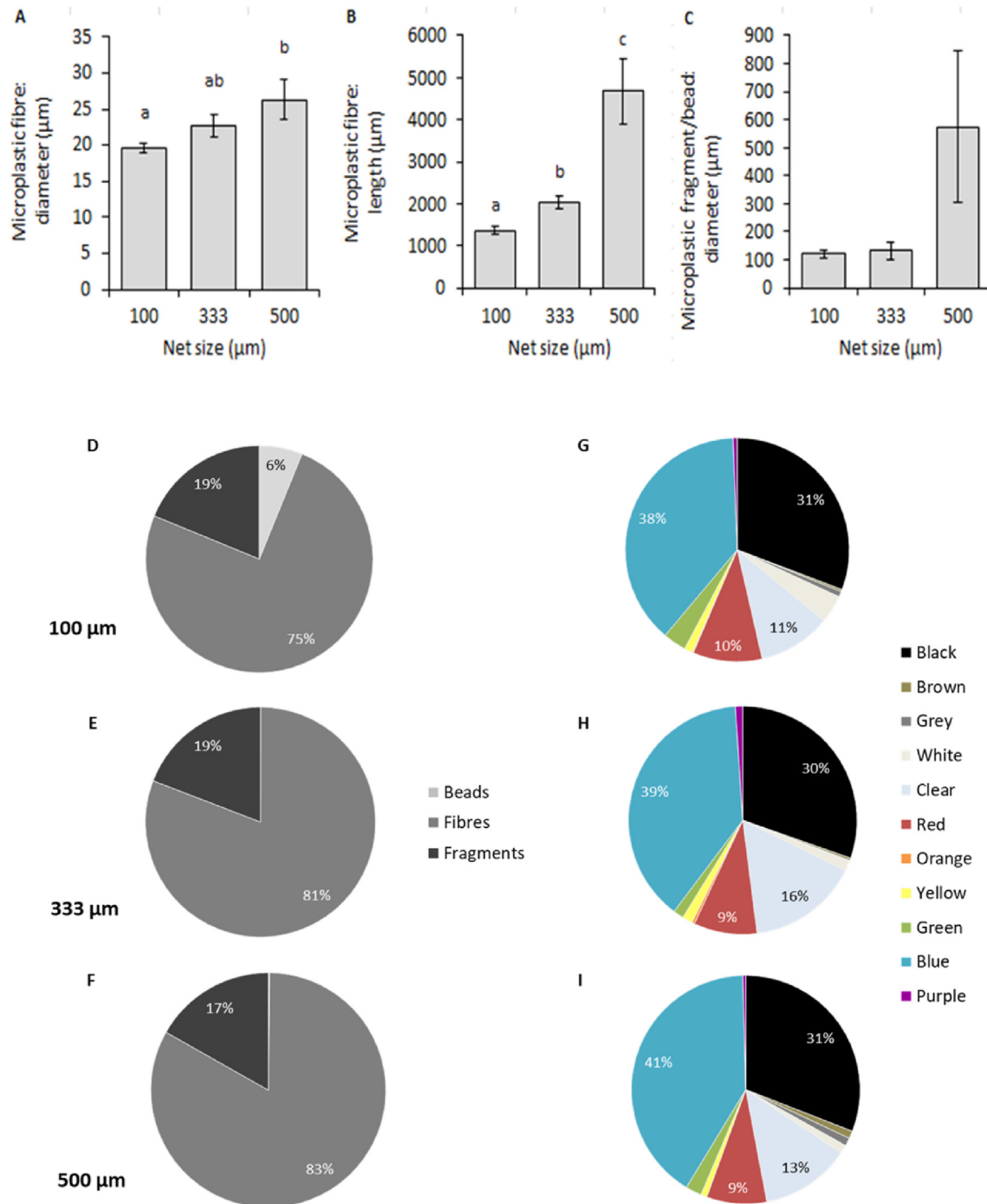


Fig. 7. Average size of microplastics identified in UK coastal samples collected using nets with different mesh size. (A) Microplastic fibre diameter; (B) Microplastic fibre length; (C) Fragment/bead diameter. Data presented as mean \pm standard error. A Kruskal-Wallis test was applied to compare datasets, with significance attributed where $P < 0.05$. Proportion of UK characterised particles by shape (D,E,F) and colour (G,H,I) for each net size; 100 μm (D,G), 333 μm (E,H), 500 μm (F,I). (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

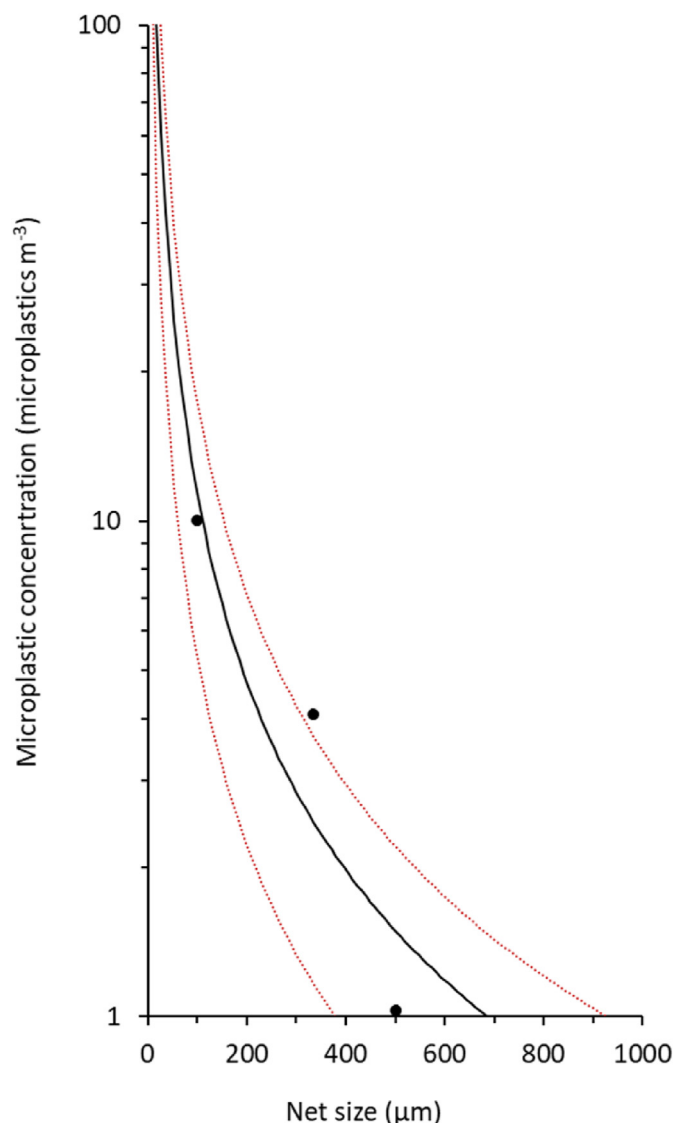


Fig. 8. Extrapolation of microplastic concentrations (logarithmic scale) based on our UK coastal samples collected using nets with 100, 333 or 500 μm mesh (black dots), using a power law (black line); 95% confidence intervals shown with dotted red lines. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

4. Discussion

Our results demonstrate that sampling with a smaller sized mesh yields a significantly higher concentration of microplastics compared to sampling with larger mesh sizes; a consistent result seen across a series of biologically productive coastal stations on both sides of the North Atlantic. Both our US and UK datasets reveal that sampling with a 100 μm net results in the capture of 10-fold greater microplastic concentrations compared with using a 500 μm net. Further, our UK sampling regime revealed a 2.5-fold increase in microplastic concentrations sampled with a 100 μm mesh compared to a 333 μm mesh. We believe this to be the first study directly comparing microplastics captured with different size mesh using nets towed concurrently. Our results demonstrate that using a traditional 333 μm mesh can result in the underestimation of waterborne microplastic concentrations owing to smaller microplastics and microfibrils being missed. Several other studies

have indicated this trend, for example: [Enders et al. \(2015\)](#) identified a greater abundance of smaller microplastic particles sampled in the smaller fraction of a staggered underway intake filtration set-up in the North Atlantic ocean; comparing discrete water samples with towed nets [Norén \(2007\)](#) found concentrations of microplastics up to 1,000 times higher when water column samples were concentrated onto an 80 μm mesh, as opposed to using a 450 μm mesh Neuston net; in the Nakdong River mouth in the Southern Sea of Korea, [Kang et al. \(2015\)](#) identified 0.62–860 microplastics m^{-3} using a 330 μm Manta trawl, and 21–15,560 microplastics m^{-3} using a 50 μm hand net; and [Barrows et al. \(2017\)](#) demonstrated that a surface grab collected over three orders of magnitude more microplastic per volume of water than sampling with a Neuston tow net; and lastly, a study by [Covernton et al. \(2019\)](#) demonstrated microplastic concentrations determined by filtering a 1 L bulk sample through an 8 μm filter was on average approximately 5.8 times greater (per L of water) than a 10 L bucket sample sieved through 63 μm mesh. All the above recent studies concur that microplastic concentration increases significantly with decreasing mesh size. As 80% of microplastic sampling campaigns focus only on the collection of >300 μm plastic debris ([Conkle et al., 2018](#)), we conclude that current estimates of marine microplastic pollution is being vastly underestimated.

Global estimates of floating microplastic debris, modelled on data primarily ascertained from 333 μm net samples, is in the order of 5–50 trillion particles ([Eriksen et al., 2014](#); [van Sebille et al., 2015](#)). Based on the relationship between microplastic concentrations identified with 100 and 333 μm nets as detailed in this study, we surmise that for buoyant microplastics >100 μm , the global plastic reservoir is in the order of 12.5–125 trillion particles. We can further extrapolate our data using a power law as prescribed elsewhere ([Cózar et al., 2014](#); [Lenz et al., 2016](#)), to estimate how many microplastics might be sampled by nets with even smaller mesh sizes ([Fig. 8](#)). Based on this extrapolation, in the waters around Plymouth (UK) we estimate the use of a 10 μm mesh net would yield on average approximately 207 microplastics m^{-3} , and by using a 1 μm mesh microplastic concentrations could exceed 3700 microplastics m^{-3} . Appreciably there are wider considerations to any such extrapolation; for example, we know microplastics can be “removed” from surface waters through coastal deposition ([Hinata et al., 2017](#)), rapid nano-fragmentation ([Andrady, 2015](#)), ingestion by biota ([Cole et al., 2013](#)), and repackaging of microplastics in faeces ([Cole et al., 2016](#); [Coppock et al., 2019](#)) and marine snow ([Porter et al., 2018](#)). However, such a model supports our hypothesis that smaller plastics are underestimated based on traditional sampling. Such a model may also be useful in providing estimates of bioavailable microplastic concentrations for exposure studies ([Lenz et al., 2016](#)). A more accurate description of the size and number of microplastics present in the environment, is essential to guide the concentration, shape and size of particles used in exposure experiments in order to identify the mechanisms of interaction between microplastics and organisms, to yield more realistic estimates of sub-lethal effects, and better understand the risk of microplastic pollution to aquatic ecosystems. On average, our results show an increase in microplastic particles sampled with a smaller mesh size, however inconsistencies to this trend are evident at individual sites. This was most notable at site N (UK), where the 333 μm net sample contained twice as many microplastics as the 100 μm net. A small variation in the general trend was also observed at sites A, E, and K (UK), with the 500 μm nets collecting slightly more microplastics than the 333 μm nets, however the differences here are negligible. Potentially, in these highly productive waters, this was a consequence of the 100 μm net becoming clogged with organic material (e.g. localised *Phaeocystis* blooms), thereby decreasing the

efficiency of the net and resulting in a decrease of water volume sampled (personal observations). Alternatively, highly localised spatial variation may have resulted in these discrepancies. On average, there was no difference in the concentration of microplastics collected by two 333 μm nets towed in parallel, however there were clear discrepancies between individual samples, highlighting the heterogeneity of microplastic concentrations at such small spatial scales; for example in Outer Portland Bay (Site 5) microplastic concentrations were 0.2 and 1.1 microplastics m^{-3} between nets trawled just metres apart. Reasons for this heterogeneity may include aggregation of microplastics around or within biological material or small scale local eddies and currents. Further, the high-density sampling around Plymouth Sound provides further evidence of the spatial and temporal variability in microplastic concentrations within localised waters, with values of 2.5–35.3 microplastics m^{-3} identified within a region of just 50 km^2 . This calls into question how frequently in time and space one must sample to gain an accurate picture of localised microplastic concentrations. Sampling practices may also influence the accuracy of collected data; for example, sea state and primary productivity can both influence the position of the net in the water, causing inaccuracies in estimating the volume of water sampled. While not applied here, sea state data can be used to compensate for wind-driven mixing of microplastics (Kooi et al., 2016; Kukulka et al., 2012).

Considering the geographical distance between our US and UK sampling sites, the number of microplastics sampled on both sides of the north Atlantic with a 100 μm mesh net were remarkably similar, with average concentrations of 6.03 ± 1.03 microplastics m^{-3} in the US and 10.03 ± 2.21 microplastics m^{-3} in the UK. All samples were taken from coastal waters, influenced by run-off from land and riverine input (Smyth et al., 2015). The slightly higher concentration of microplastics sampled in the UK is likely due to the sites' proximity to the coast, with the furthest site sampled in the UK being 6.5 km from shore and the furthest site sampled in the US being 24 km from the shore. A previous study in the same UK region showed that the concentration of microplastics decreased with distance from the shore (Steer et al., 2017). Highest microplastic concentrations in our US samples were associated with the outflows of the Penobscot and Piscataqua rivers, and in our UK samples the greatest abundance of microplastics (35.3 particles m^{-3}) was found at the mouth of the River Plym (Site H). Rivers, which receive inputs from agriculture, industry, storm water drains and sewage outflow, are hugely important transport pathways of plastic from land to sea (Lebreton and Andrady, 2019; Lebreton et al., 2017). Sampling at site H occurred after a storm event, and we hypothesise that the high microplastic concentrations observed were associated with high rainfall potentially resulting in the flushing out of roads, drainage systems and agricultural land, and the possible overflow of wastewater treatment works (Horton et al., 2017; Moore et al., 2002).

In addition to sampling a greater number of microplastics with a smaller size mesh, the fibres that were sampled were also significantly smaller. Sampling with a smaller mesh net therefore not only gives a better indication of the microplastic budget but also gives a better estimation of the abundances of microplastic particles of a size that are bioavailable to small marine organisms such as zooplankton (Botterell et al., 2018). Microplastics can be ingested by a range of marine organisms, including zooplankton (Desforges et al., 2015; Steer et al., 2017; Sun et al., 2017), deep sea invertebrates (Courteney-Jones et al., 2019), bivalves, and fish destined for human consumption (Rochman et al., 2015; Walkinshaw et al., 2020), with the capacity to impact upon the health of the organism and potentially their ecosystem functionality (Galloway et al., 2017; Green, 2016). Using smaller meshed nets will allow

researchers to better sample and estimate the abundance and bioavailability of microplastics, in turn allowing more accurate evaluations of the risks microplastics pose to biota, biodiversity, ecosystem function and productivity. The fact that microplastics less than 100 μm in size were sampled with a 100 μm mesh net is indicative of some of these plastics becoming trapped in organic material (e.g. exopolymeric agglomerations, phytoplankton; Long et al., 2015; Summers et al., 2018).

Fibres were the predominant type of microplastic identified in all our environmental samples (84% USA; 77% UK), being principally black or blue in colour. Microplastic fibres can stem from the breakdown of larger plastic items (e.g. rope) (Welden and Cowie, 2017) or the release of microfibrils from synthetic garments during washing cycles (Napper and Thompson, 2016). Abrasion from clothing is also likely to be a significant source of fibre pollution, demonstrated by high quantities observed in atmospheric fallout (Dris et al., 2016) and run off from snow melts (Bergmann et al., 2019). Rayon (biopolymer), polypropylene and polyester are widely used in textiles, providing further evidence that wastewater effluent (containing microfibrils from clothes washing (Napper and Thompson, 2016)) and degradation of fishing gear (Welden and Cowie, 2017) are substantial sources of microplastics in coastal waters (Murphy et al., 2016; Napper and Thompson, 2016). The elastomers identified in the UK samples may be associated with vehicle tyre wear (Kole et al., 2017), with inputs stemming from highway drainage (e.g. A38, Tamar bridge). A better understanding of the detailed characteristics of microplastics in the marine environment may help elucidate the origin of these particles, as discussed above, which in turn can help influence societal behaviour and drive future policy intervention.

In recent years there have been calls for harmonisation of microplastic sampling methods (Frias and Nash, 2019; Hartmann et al., 2019; Hidalgo-Ruz et al., 2012), to facilitate comparability between data sets. For example, collection may be via discrete sampling such as using a Niskin bottle (Courteney-Jones, 2017) or via a more continuous sampling method such as a Manta trawl (Sadri and Thompson, 2014) or ships underway system (Lenz et al., 2015), all with differences in error rate and sampling efficiency. Differences in laboratory processing such as methods to digest biotic material, sub-sampling, characterisation and polymeric analysis further serve to make comparisons challenging. Despite these harmonisation calls however, a huge range of different techniques for sampling and quantifying plastics, each championed by different research groups, continue to be used. Furthermore, polymeric analysis of samples would ideally be carried out using automated detection of particles, such as Focal Plane Array (FPA) or image mapping using FT-IR. Whilst this is the clear way forward in microplastic research, when these methods have been used to date, samples have tended to be very 'clean', and not yet suitable for complex, biologically rich samples such as those obtained in this study.

5. Conclusion

We have demonstrated that the 333 μm nets commonly used for microplastics sampling underestimate microplastic abundance, particularly for <333 μm microplastics that are within the optimal prey size range of numerous marine organisms. Where possible, sampling should aim to collect the fullest range of microplastics present, with an appreciation that sampling with larger mesh size nets will not give an accurate estimate of abundance or a full account of the microplastics present within the water column. However, we also appreciate that when sampling there needs to be a balance between efficiency, accuracy and detail. We surmise that sampling with smaller sized mesh nets (i.e. 100 μm) gives a better

representation of microplastic concentrations in the natural environment and helps to ascertain more reliable estimates of microplastic budgets. In turn this effort allows for better assessment of the current level of risk posed to the marine environment, better guiding monitoring efforts, and providing a clearer benchmark against which to judge the effectiveness of future management scenarios.

CRedit authorship contribution statement

Penelope K. Lindeque: Conceptualization, Methodology, Investigation, Writing - original draft, Writing - review & editing, Supervision, Project administration, Funding acquisition. **Matthew Cole:** Conceptualization, Methodology, Investigation, Writing - original draft, Writing - review & editing, Supervision, Formal analysis. **Rachel L. Coppock:** Validation, Formal analysis, Writing - review & editing. **Ceri N. Lewis:** Investigation, Writing - review & editing. **Rachael Z. Miller:** Investigation, Writing - review & editing. **Andrew J.R. Watts:** Investigation, Writing - review & editing. **Alice Wilson-McNeal:** Investigation, Formal analysis, Writing - original draft, Writing - review & editing. **Stephanie L. Wright:** Investigation, Writing - review & editing. **Tamara S. Galloway:** Funding acquisition, Investigation, Writing - original draft, Writing - review & editing.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgments

The research was funded by the Natural Environment Research Council (NERC Grant NE/L003988/1 and NE/L007010/1) and the University of Exeter and Plymouth Marine Laboratory collaboration fund, with in-kind contributions from the 'Rozalia Project'. We would like to thank March Limited of Bermuda for philanthropic support. We thank the crews of the American Promise and Plymouth Quest for their enthusiasm, patience and hard work in helping collect the environmental samples, Lara Mesquita for assisting with FT-IR analyses, and Dr. James Clark and Dr. Victor Vicente-Martinez for their helpful comments on the manuscript.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.envpol.2020.114721>.

References

- Andrady, A.L., 2015. Persistence of plastic litter in the oceans. In: Bergmann, M., Gutow, L., Klages, M. (Eds.), *Marine Anthropogenic Litter*. Springer, pp. 57–72.
- Barrows, A.P., Neumann, C.A., Berger, M.L., Shaw, S.D., 2017. Grab vs. neuston tow net: a microplastic sampling performance comparison and possible advances in the field. *Anal Methods* 9, 1446–1453.
- Bergmann, M., Mützel, S., Primpke, S., Tekman, M.B., Trachsel, J., Gerds, G., 2019. White and wonderful? Microplastics prevail in snow from the alps to the arctic. *Sci Adv* 5, eaax1157.
- Besseling, E., Wegner, A., Foekema, E.M., van den Heuvel-Greve, M.J., Koelmans, A.A., 2013. Effects of microplastic on fitness and PCB bioaccumulation by the lugworm *Arenicola marina* (L.). *Environ. Sci. Technol.* 47, 593–600.
- Botterell, Z.L., Beaumont, N., Dorrington, T., Steinke, M., Thompson, R.C., Lindeque, P.K., 2018. Bioavailability and effects of microplastics on marine zooplankton: a review. *Environ. Pollut.* 245, 98–110.
- Burns, E.E., Boxall, A.B., 2018. Microplastics in the aquatic environment: evidence for or against adverse impacts and major knowledge gaps. *Environ. Toxicol. Chem.* 37, 2776–2796.
- Clark, J.R., Cole, M., Lindeque, P., Fileman, E., Blackford, J., Lewis, C., Lenton, T.M., Galloway, T.S., 2016. Marine microplastic debris: a targeted plan for understanding and quantifying interactions with marine life. *Front. Ecol. Environ.* 14, 317–324.
- Cole, M., Lindeque, P., Halsband, C., Galloway, T.S., 2011. Microplastics as contaminants in the marine environment: a review. *Mar. Pollut. Bull.* 62, 2588–2597.
- Cole, M., Lindeque, P., Fileman, E., Halsband, C., Goodhead, R., Moger, J., Galloway, T.S., 2013. Microplastic ingestion by zooplankton. *Environ. Sci. Technol.* 12, 6646–6655.
- Cole, M., Webb, H., Lindeque, P., Fileman, E., Halsband, C., Galloway, T.S., 2014. Isolation of microplastics in biota-rich samples and marine biota. *Sci. Rep.* 4, 4528.
- Cole, M., Lindeque, P., Fileman, E., Halsband, C., Galloway, T., 2015. The impact of polystyrene microplastics on feeding, function and fecundity in the marine copepod *Calanus helgolandicus*. *Environ. Sci. Technol.* 49, 1130–1137.
- Cole, M., Lindeque, P.K., Fileman, E., Clark, J., Lewis, C., Halsband, C., Galloway, T.S., 2016. Microplastics alter the properties and sinking rates of zooplankton faecal pellets. *Environ. Sci. Technol.* 50, 3239–3246.
- Cole, M., Coppock, R., Lindeque, P.K., Altin, D., Reed, S., Pond, D.W., Sørensen, L., Galloway, T.S., Booth, A.M., 2019. Effects of nylon microplastic on feeding, lipid accumulation, and moulting in a coldwater copepod. *Environ. Sci. Technol.* 53, 7075–7082.
- Conkle, J.L., Del Valle, C.D.B., Turner, J.W., 2018. Are we underestimating microplastic contamination in aquatic environments? *Environ. Manag.* 61, 1–8.
- Coppock, R.L., Galloway, T.S., Cole, M., Fileman, E., Queirós, A.M., Lindeque, P.K., 2019. Microplastics alter feeding selectivity and faecal density in the copepod, *Calanus helgolandicus*. *Sci. Total Environ.* 687, 780–789.
- Courteney-Jones, W., Quinn, B., Gary, S.F., Mogg, A.O., Narayanaswamy, B.E., 2017. Microplastic pollution identified in deep-sea water and ingested by benthic invertebrates in the Rockall Trough, North Atlantic Ocean. *Environ. Pollut.* 231, 271–280.
- Courteney-Jones, W., Quinn, B., Ewins, C., Gary, S.F., Narayanaswamy, B.E., 2019. Consistent microplastic ingestion by deep-sea invertebrates over the last four decades (1976–2015), a study from the North East Atlantic. *Environ. Pollut.* 244, 503–512.
- Covernton, G.A., Pearce, C.M., Gurney-Smith, H.J., Chastain, S.G., Ross, P.S., Dower, J.F., Dudas, S.E., 2019. Size and shape matter: a preliminary analysis of microplastic sampling technique in seawater studies with implications for ecological risk assessment. *Sci. Total Environ.* 667, 124–132.
- Cózar, A., Echevarría, F., González-Gordillo, J.L., Irigoien, X., Úbeda, B., Hernández-León, S., Palma, A.T., Navarro, S., García-de-Lomas, J., Ruiz, A., 2014. Plastic debris in the open ocean. *Proc. Natl. Acad. Sci. Unit. States Am.* 111, 10239–10244.
- de Sá, L.C., Oliveira, M., Ribeiro, F., Rocha, T.L., Futter, M.N., 2018. Studies of the effects of microplastics on aquatic organisms: what do we know and where should we focus our efforts in the future? *Sci. Total Environ.* 645, 1029–1039.
- Desforges, J.-P.W., Galbraith, M., Ross, P.S., 2015. Ingestion of microplastics by zooplankton in the northeast Pacific Ocean. *Arch. Environ. Contam. Toxicol.* 69, 320–330.
- Dris, R., Gasperi, J., Saad, M., Mirande, C., Tassin, B., 2016. Synthetic fibers in atmospheric fallout: a source of microplastics in the environment? *Mar. Pollut. Bull.* 104, 290–293.
- Duncan, E.M., Broderick, A.C., Fuller, W.J., Galloway, T.S., Godfrey, M.H., Hamann, M., Limpus, C.J., Lindeque, P.K., Mayes, A.G., Omeyer, L.C., 2019. Microplastic ingestion ubiquitous in marine turtles. *Global Change Biol.* 25, 744–752.
- Enders, K., Lenz, R., Stedmon, C.A., Nielsen, T.G., 2015. Abundance, size and polymer composition of marine microplastics > 10 µm in the Atlantic Ocean and their modelled vertical distribution. *Mar. Pollut. Bull.* 100, 70–81.
- Eriksen, M., Lebreton, L.C., Carson, H.S., Thiel, M., Moore, C.J., Borerro, J.C., Galgani, F., Ryan, P.G., Reisser, J., 2014. Plastic pollution in the world's oceans: more than 5 trillion plastic pieces weighing over 250,000 tons afloat at sea. *PLoS One* 9, e111913.
- Frias, J., Nash, R., 2019. Microplastics: finding a consensus on the definition. *Mar. Pollut. Bull.* 138, 145–147.
- Galloway, T.S., Cole, M., Lewis, C., 2017. Interactions of microplastic debris throughout the marine ecosystem. *Nat. Ecol. Evol.* 1, 1–8.
- Green, D.S., 2016. Effects of microplastics on European flat oysters, *Ostrea edulis* and their associated benthic communities. *Environ. Pollut.* 216, 95–103.
- Hartmann, N.B., Hüffer, T., Thompson, R., Hasselöv, M., Verschoor, A., Daugaard, A., Rist, S., Karlsson, T.M., Brennholt, N., Cole, M., Herrling, M., Heß, M., Ivleva, N., Lusher, A., Wagner, M., 2019. Are we speaking the same language? Recommendations for a definition and categorization framework for plastic debris. *Environ. Sci. Technol.* 53, 1039–1047.
- Hidalgo-Ruz, V., Gutow, L., Thompson, R.C., Thiel, M., 2012. Microplastics in the marine environment: a review of the methods used for identification and quantification. *Environ. Sci. Technol.* 46, 3060–3075.
- Hinata, H., Mori, K., Ohno, K., Miyao, Y., Kataoka, T., 2017. An estimation of the average residence times and onshore-offshore diffusivities of beached microplastics based on the population decay of tagged meso-and macrolitter. *Mar. Pollut. Bull.* 122, 17–26.
- Horton, A.A., Walton, A., Spurgeon, D.J., Lahive, E., Svendsen, C., 2017. Microplastics in freshwater and terrestrial environments: evaluating the current understanding to identify the knowledge gaps and future research priorities. *Sci. Total Environ.* 586, 127–141.
- Kang, J.-H., Kwon, O.Y., Lee, K.-W., Song, Y.K., Shim, W.J., 2015. Marine neustonic microplastics around the southeastern coast of Korea. *Mar. Pollut. Bull.* 96,

- 304–312.
- Kole, P.J., Löhr, A.J., Van Belleghem, F., Ragas, A., 2017. Wear and tear of tyres: a stealthy source of microplastics in the environment. *Int. J. Environ. Res. Publ. Health* 14, 1265.
- Kooi, M., Reisser, J., Slat, B., Ferrari, F.F., Schmid, M.S., Cunsolo, S., Brambini, R., Noble, K., Sirks, L.-A., Linders, T.E., 2016. The effect of particle properties on the depth profile of buoyant plastics in the ocean. *Sci. Rep.* 6, 33882.
- Kukulka, T., Proskurowski, G., Morét-Ferguson, S., Meyer, D., Law, K., 2012. The effect of wind mixing on the vertical distribution of buoyant plastic debris. *Geophys. Res. Lett.* 39.
- Lebreton, L., Andrady, A., 2019. Future scenarios of global plastic waste generation and disposal. *Palgrave Commun* 5, 6.
- Lebreton, L.C., Van der Zwet, J., Damsteeg, J.-W., Slat, B., Andrady, A., Reisser, J., 2017. River plastic emissions to the world's oceans. *Nat. Commun.* 8, 15611.
- Lenz, R., Enders, K., Stedmon, C.A., Mackenzie, D.M., Nielsen, T.G., 2015. A critical assessment of visual identification of marine microplastic using Raman spectroscopy for analysis improvement. *Mar. Pollut. Bull.* 100, 82–91.
- Lenz, R., Enders, K., Nielsen, T.G., 2016. Microplastic exposure studies should be environmentally realistic. *Proc. Natl. Acad. Sci. Unit. States Am.* 113, E4121–E4122.
- Long, M., Moriceau, B., Gallinari, M., Lambert, C., Huvet, A., Raffray, J., Soudant, P., 2015. Interactions between microplastics and phytoplankton aggregates: impact on their respective fates. *Mar. Chem.* 175, 39–46.
- Lusher, A., 2015. Microplastics in the marine environment: distribution, interactions and effects. In: Bergmann, M., Gutow, L., Klages, M. (Eds.), *Marine Anthropogenic Litter*. Springer, pp. 245–307.
- Lusher, A., Welden, N., Sobral, P., Cole, M., 2016. Sampling, isolating and identifying microplastics ingested by fish and invertebrates. *Anal. Methods* 9, 1346–1360.
- Moore, C.J., Moore, S.L., Weisberg, S.B., Lattin, G.L., Zellers, A.F., 2002. A comparison of neustonic plastic and zooplankton abundance in southern California's coastal waters. *Mar. Pollut. Bull.* 44, 1035–1038.
- Murphy, F., Ewins, C., Carbonnier, F., Quinn, B., 2016. Wastewater treatment works (WwTW) as a source of microplastics in the aquatic environment. *Environ. Sci. Technol.* 50, 5800–5808.
- Napper, I.E., Thompson, R.C., 2016. Release of synthetic microplastic plastic fibres from domestic washing machines: effects of fabric type and washing conditions. *Mar. Pollut. Bull.* 112, 39–45.
- Nelms, S., Barnett, J., Brownlow, A., Davison, N., Deaville, R., Galloway, T., Lindeque, P., Santillo, D., Godley, B., 2019. Microplastics in marine mammals stranded around the British coast: ubiquitous but transitory? *Sci. Rep.* 9, 1075.
- Norén, F., 2007. Small Plastic Particles in Coastal Swedish Waters. KIMO Sweden.
- Porter, A., Lyons, B.P., Galloway, T.S., Lewis, C.N., 2018. The role of marine snows in microplastic fate and bioavailability. *Environ. Sci. Technol.* 52, 7111–7119.
- R Core Team, 2019. R: A Language and Environment for Statistical Computing, 0 ed, 3.6. R Foundation for Statistical Computing, Vienna, Austria.
- Rochman, C.M., Tahir, A., Williams, S.L., Baxa, D.V., Lam, R., Miller, J.T., Teh, F.-C., Werorilangi, S., Teh, S.J., 2015. Anthropogenic debris in seafood: plastic debris and fibers from textiles in fish and bivalves sold for human consumption. *Sci. Rep.* 5, 14340.
- Rochman, C.M., Browne, M.A., Underwood, A., Franeker, J.A., Thompson, R.C., Amaral-Zettler, L.A., 2016. The ecological impacts of marine debris: unraveling the demonstrated evidence from what is perceived. *Ecology* 97, 302–312.
- Sadri, S.S., Thompson, R.C., 2014. On the quantity and composition of floating plastic debris entering and leaving the Tamar Estuary, Southwest England. *Mar. Pollut. Bull.* 81, 55–60.
- Skjoldal, H.R., Wiebe, P.H., Postel, L., Knutsen, T., Kaartvedt, S., Sameoto, D.D., 2013. Intercomparison of zooplankton (net) sampling systems: results from the ICES/GLOBEC sea-going workshop. *Prog. Oceanogr.* 108, 1–42.
- Smyth, T., Atkinson, A., Widdicombe, S., Frost, M., Allen, I., Fishwick, J., Queiros, A., Sims, D., Barange, M., 2015. The Western channel observatory. *Prog. Oceanogr.* 137, 335–341.
- Steer, M., Cole, M., Thompson, R.C., Lindeque, P.K., 2017. Microplastic ingestion in fish larvae in the western English Channel. *Environ. Pollut.* 226, 250–259.
- Stock, F., Kochleus, C., Bansch-Baltruschat, B., Brennholt, N., Reifferscheid, G., 2019. Sampling techniques and preparation methods for microplastic analyses in the aquatic environment—A review. *Trac. Trends Anal. Chem.* 113, 84–92.
- Summers, S., Henry, T., Gutierrez, T., 2018. Agglomeration of nano- and microplastic particles in seawater by autochthonous and de novo-produced sources of exopolymeric substances. *Mar. Pollut. Bull.* 130, 258–267.
- Sun, X., Li, Q., Zhu, M., Liang, J., Zheng, S., Zhao, Y., 2017. Ingestion of microplastics by natural zooplankton groups in the northern South China Sea. *Mar. Pollut. Bull.* 115, 217–224.
- Sussarellu, R., Suquet, M., Thomas, Y., Lambert, C., Fabioux, C., Pernet, M.E.J., Le Goïc, N., Quillien, V., Mingant, C., Epelboin, Y., 2016. Oyster reproduction is affected by exposure to polystyrene microplastics. *Proc. Natl. Acad. Sci. Unit. States Am.* 113, 2430–2435.
- van Sebille, E., Wilcox, C., Lebreton, L., Maximenko, N., Hardesty, B.D., Franeker, J.A.v., Eriksen, M., Siegel, D., Galgani, F., Law, K.L., 2015. A global inventory of small floating plastic debris. *Environ. Res. Lett.* 10, 124006.
- Walkinshaw, C., Lindeque, P.K., Thompson, R., Tolhurst, T., Cole, M., 2020. Microplastics and seafood: lower trophic organisms at highest risk of contamination. *Ecotoxicol. Environ.* 190, 110066.
- Wegner, A., Besseling, E., Foekema, E., Kamermans, P., Koelmans, A., 2012. Effects of nanopolystyrene on the feeding behavior of the blue mussel (*Mytilus edulis* L.). *Environ. Toxicol. Chem.* 31, 2490–2497.
- Welden, N.A., Cowie, P.R., 2017. Degradation of common polymer ropes in a sub-littoral marine environment. *Mar. Pollut. Bull.* 118, 248–253.
- Wright, S., Rowe, D., Thompson, R.C., Galloway, T.S., 2013. Microplastic ingestion decreases energy reserves in marine worms. *Curr. Biol.* 23, 1031–1033.