Microplastics in Freshwater Biota: A Critical Review of Isolation, Characterization, and Assessment Methods

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1. Introduction

Freshwater systems are closely coupled with the terrestrial environment, and as such, provide key pathways for the influx of basal resources as well as pollutants.[1,2] Headwater streams serve as important vectors for the transport of materials which may extend into the downstream reaches of rivers, lakes, and estuaries,[3,4] while longitudinal physical gradients can regulate biotic interactions with such materials by controlling community assemblages.[1,3] All of which are assumed relevant for microplastic (MP; 1 µm to 5 mm) movement within freshwater food webs. Freshwaters play an important role in the overall lifecycle of microplastics (MPs) in the environment, functioning as receiving waters for waste water discharges and landfill leachates where the majority of MPs are sourced,[5] while also prolonging the longevity of MPs by retaining them in their sediment[6,7] and exporting them during high flow or flood events.[8] Similar to the marine environment, MPs in freshwater are widespread and pervasive, having been reported in the surface waters of lakes[9–11] and rivers,[12–16] as well as in river and lake shore sediments.[11,17]

Unlike the marine environment however, and until quite recently, MPs in freshwater systems were comparatively understudied,[18–20] with a level of growth in marine scientific literature, in the context of MPs, estimated to be five times that of freshwater ecosystems.[20] Resultant of which, the hydrodynamics influencing MP behavior in freshwater as well as biological interactions with MPs in nature are relatively undetermined, particularly among lower trophic levels (e.g., primary consumers). The novelty of freshwater MP research, combined with the comparatively low number of field studies that focus on freshwater biota thus far, presents challenges in MP isolation and characterization, especially in methodological standardization, despite the evident parallels with marine MP...
research and the experimental studies conducted to date. To our knowledge, there are no reviews that focus on the challenges faced in analyzing freshwater biota.

Up to now, most studies on effects in biota were conducted with MP particle concentrations far exceeding those measured in the environment, with an apparent mismatch between particle sizes used to determine effect thresholds in laboratory experimentation and those reported in nature, something which is difficult to mitigate given the current technological limitations in the verification of polymers collected from the field. It is noted however that while maintaining ecological relevance is important, what is relevant at present may not be relevant in the future, and we therefore argue that higher concentrations should also be included to account for future risk.

Here, we present a critical review of the current level of literature pertaining to isolation, characterization, and assessment of MPs in freshwater biota, outlining the challenges and limitations currently faced by freshwater MP researchers, which may well apply to the MP research field as a whole. We provide recommendations for methodological standardization, particularly with regard to particle characterization, quality assurance and quality control (QA/QC) procedures and reporting, as well as considerations for the assessment of MPs in freshwater biota as a means of improving comparisons between studies.

2. Microplastics in Freshwater Biota: Occurrence and Methodological Challenges

2.1. Microplastics in Freshwater Field Samples

Attention given toward detecting and monitoring MPs in biotic (and abiotic) components of freshwater environments has increased in recent years, expanding our knowledge on susceptibility of species to ingestion, potential translocation within body tissues, and trophic transfer of MPs. Freshwater MP studies undertaken to date on biota have focused on higher taxa, namely fish and birds, with fewer environmental studies assessing lower taxa (Table 1). All studies showed evidence of MP ingestion within the studied species, either through direct consumption and/or secondary ingestion (i.e., trophic transfer). Though for the most part, the specific pathways were not determined. Lower taxa were primarily assessed to ascertain if the prevalence of MPs within individuals, including macroinvertebrates, tadpoles of four species of frog and/or toads, and the Asian clam Corbicula fluminea (Müller, 1774), was related to the type (abundance, shape, and polymer distribution) and relative abundance of MPs in water and/or sediment samples, thus assessing their potential as bioindicators of MP pollution in freshwater environments. Species susceptibility to MP ingestion due to their ecological niche and functional feeding group (grouping feeding habits, traits, and behavior) was also considered in order to assess the relative processes related to the uptake of MPs; processes such as particle selection.

Fish and bird species were also assessed as sentinel taxon for MP exposure. Due to the higher trophic level of freshwater fish and the potential for trophic transfer of MPs, most studies have assessed the presence, occurrence, and type of

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MPs in gastrointestinal (GI) tracts, identifying sampling methodologies for doing so,
scant at present, have also analyzed liver and muscle tissue for translocation of MPs,
however potential translocation routes and impacts were not assessed. Studies on fish also
included other parameters for assessing the variability in individual MP burdens, including feeding habitats and behaviors (demersal vs pelagic), fish size (assessing consumption and potential bioaccumulation), and the environmental availability of MPs.

### Table 1. Isolation protocols described in field studies reviewed for analyzing MP particles in freshwater biota along with units, shapes, and size ranges.

| Author(s)          | Organism a) | Component b) | Treatment | Exposure c) | Reporting units d) | Shapes e) | Size range(s) |
|--------------------|-------------|--------------|-----------|-------------|---------------------|-----------|---------------|
| Sanchez et al. 2014 | Fish        | GI tracts    | –         | –           | –                   | –         | –             |
| Phillips and Bonner 2015 | Fish        | GI tracts    | –         | –           | –                   | Films, fragments, and filaments (fibers not recorded) | Max 5.5 mm (linear length) |
| Faure et al. 2015 | Fish and birds | GI tracts    | –         | –           | –                   | Mostly fibers and fragments | –             |
| Holland et al. 2016 | Birds       | GI tracts    | –         | –           | –                   | Mostly fragments | 50 µm to 5 mm |
| Peters and Bratton 2016 | Fish        | GI tracts    | –         | –           | –                   | 96% threads (fibrous) | 4% > 5 mm |
| Biginagwa et al. 2016 | Fish        | GI tracts    | NaOH (10 mol L⁻¹) | 24 h (60 °C) | –                   | –         | –             |
| Silva-Cavalcanti et al. 2017 | Fish       | GI tracts    | –         | –           | –                   | Fibers only shape described (46.6%) | 1–12 mm |
| Campbell et al. 2017 | Fish        | GI tracts    | 10% NaClO and HNO₃: NaClO (1:10 v/v) | Overnight and until dissolved (room) | Mean MPs fish⁻¹ | Predominantly fibers and fragments | –             |
| Pazos et al. 2017 | Fish        | GI tracts    | 30% H₂O₂ | Until dissolved (60 °C) | MPs fish⁻¹ | 96% fibers | 60 µm to 4.7 mm |
| Vendel et al. 2017 | Fish        | GI tracts    | –         | –           | –                   | 90% fibers, remainders films and fragments | –             |
| McGoran et al. 2017 | Fish        | GI tracts    | –         | –           | –                   | Mostly fibers | –             |
| Jabeen et al. 2017 | Fish        | GI tracts    | 30% H₂O₂ and density separation NaCl (1.2 g cm⁻³) | 24–72 h (65 °C) | Items g⁻¹, items individual⁻¹ | Fibers, fragments, and film | 40 µm to 5 mm |
| Horton et al. 2018 | Fish        | GI tracts    | –         | –           | –                   | Particles fish⁻¹ | Most fibers (75%) | <5 mm |
| Collard et al. 2018 | Fish        | GI tracts, liver, and muscle tissue | NaClO (14 g L⁻¹) | Overnight (room) | AP g⁻¹ of stomach content | Fibers dominant | 390 µm to 7.38 mm |
| McNeish et al. 2018 | Fish        | GI tracts    | Dried 24 h (75 °C) 30% H₂O₂: Fe(II) (0.05 mol L⁻¹) | Until dissolved (75 °C) | No. MPs fish⁻¹ | Fibers dominant | <1.5 mm, 1.6–3.2 mm, >3.3 mm |
| Hurley et al. 2017 | Invertebrates | Whole        | 10% KOH | ≤10 min (60 °C) | Particles g⁻¹ (wet weight) | 87% fibers, remainders fragments | 55 µm to 4.1 mm |
| Hu et al. 2018 | Tadpoles    | Whole        | 30% H₂O₂  | ≤72 h (65 °C) | Items individual⁻¹ | Mostly fibers | <0.5 mm |
| Nel et al. 2018 | Invertebrates | Whole        | HNO₃  | 6 h (room) + 1 h (100 °C) | Particles mg⁻¹ (wet weight) | –         | –             |
| Su et al. 2018 | Molluscs     | Soft tissue  | 30% H₂O₂ | ≤72 h (65 °C) | Items g⁻¹ | Fibers | 0.021–4.02 mm |
| Windsor et al. 2019 | Invertebrates | Whole        | Density separation NaCl (1.2 g cm⁻³) and 15% H₂O₂ | 48 h (25°C) | MP mg⁻¹ | –         | 500 µm to 5 mm |

a) Broad classification of organisms studied (invertebrates = benthic macroinvertebrates); b) Component is the target component analyzed for MPs; c) Exposure = time of digestion (temperature); d) Units used to report microplastic burden in terms of abundance or concentration.
Feeding behaviors, i.e., whether fish are benthic feeders or pelagic predators, may influence MP ingestion, due to different encounter rates and/or different types of MPs (size, density, color, and morphology) found in the benthic sediment and water column.[36,37] Trophic roles may also influence MP burdens in fish, where MPs have been consumed via trophic transfer—detritivores and predators may indirectly ingest MPs when consuming prey or scavenging detrital matter.[36,42]

2.2. Extracting Microplastics from Biological Samples: Challenges in Isolation

Lack of standardization in isolation methods for MPs internalized in body cavities/tissues or entangled (externally) in biota is currently recognized as a foremost concern across all MP research.[43–46] The lack of harmonization across studies means that it is difficult to facilitate comparisons or assess MP exposure levels between similar species.[44,47] An increasing number of methods have been modified and devised in recent years that include dissection, depuration, homogenization, and digestion of tissue, with the selection of one or more methods largely determined by the research question being explored.[48] A number of isolation approaches have been observed to damage polymers (e.g., acid digestion)[48] or under represent smaller MP particles (e.g., visual sorting without digestion).[44]

Post dissection visual observation methods or sorting do not utilize chemicals and are often a preferred method when identifying and separating MPs from freshwater biota, particularly where dietary remains are required to remain intact. Dissection followed by rinsing of GI tracts with water is a well-practiced protocol in freshwater field studies, especially among fish[9,32,36,37,41] and birds,[31] but can lead to an underrepresentation of MPs resultant of challenges with identifying and enumerating smaller particles that may adhere to dietary remains and other biological material within the GI tract of an organism.[31,42,49] Moreover, the lack of a digestion pretreatment means that where organs are damaged due to freezing or drying out, particle abundance may be underestimated as GI tracts cannot be analyzed in their entirety.[9]

While enzymatic digestion (e.g., trypsin, proteinase-k)[50,51] is an emerging isolation technique in MP research, chemical treatments (e.g., oxidizers, acids, bases), due to their less expensive nature and relatively high recovery rates,[52] are generally the more applied method. Although recently used to extract MPs from freshwater invertebrate samples (i.e., Chironomus spp.),[27] strong mineral acids, such as nitric acid (HNO₃), have been reported to cause degradation and damage to sensitive polymers (e.g., polyamide (PA), polystyrene (PS), polyethylene (PE)) and are often substituted for bases (e.g., 10% KOH) or oxidizers (e.g., 30% H₂O₂) depending on the organ or organism under analysis. Alkaline hydrolysis is easy to perform on soft tissue organisms such as tubificid worms[26] or on the GI tracts of freshwater fish,[19] and can be achieved at room temperature.[53] Wet peroxide oxidation (WPO) using H₂O₂, often in the presence of a Fe(II) catalyst (Fenton’s reagent), is a preferred method for organisms that contain hard parts such as benthic macroinvertebrates,[24,54] which usually require an application of heat to break down the chitin within their exoskeleton (personal observation), but has also been used as a method to digest the soft tissue of C. fluminea[28] tadpoles,[49] and the GI tracts of fish.[34,35,55] However, such treatments have limitations, with NaOH (10 mol L⁻¹) reported to inflict damage or discoloration on PA and PE particles.[49] Biginagwa et al.[38] who tested this method prior to digesting the GI tracts of Nile perch Lates niloticus (Linnaeus, 1758) and Nile tilapia Oreochromis niloticus (L.), report a high digestion efficiency. Similar observations have been made for WPO, which has been found to be an efficient method for breaking down biogenic material despite discoloring (bleaching) MP particles. This could possibly affect the determination of colors rather than leading to an underestimation of particle abundance.[28,56]

2.3. Validating Techniques for Microplastic Isolation

As many chemical digestions are known to incur negative effects on the characteristics of certain pH sensitive polymers[48,49] the choice of digestion method used to isolate MP particles, along with the manner in which it is applied (i.e., heat, level of exposure), can impact significantly upon MP recovery rates.[57] As well as this, difficulties in determining whether all MPs within a field sample are effectively recovered, renders the validation of such isolation techniques a useful practice within MP research.[45] Extraction efficacy is typically assessed via “spiking,” whereby MPs of a predetermined abundance, type, size range, and morphology are incorporated into a control sample, subjected to a specific treatment, and the number of MPs recovered expressed as a percentage[58] (the number of particles remaining in a sample following exposure to an extraction method).[57] This is generally performed on biological tissue, such as fish,[58] or wild mussels,[51] and as recommended by Hermsen et al.,[45] should be included as both positive samples (i.e., in tissue) and blank samples (negative controls), treated in parallel to field samples.

To our knowledge, there are no freshwater biological studies, from the field at least, that report the testing of such MP isolation protocols, with exception to Biginagwa et al.,[38] who assessed the digestion efficiency of NaOH in breaking down biogenic material. It is expected however that most biological field studies follow validated techniques. Windsor et al.[24] applied a method on freshwater benthic macroinvertebrate samples previously tested by Avio et al.[58] who report an average extraction yield from laboratory-acclimatized mullet Mugil cephalus (L.) GI tracts of 95% ± 2 (mean% ± SEM), while Hurley et al.[26] used a 10% KOH solution to digest Tubifex tubifex (M.) tissue following a series of protocols previously validated by Karami et al.[59] on African catfish Clarias gariepinus (Burchell, 1822). Some authors, such as Hu et al.,[59] even replicated extraction methods previously tested by themselves in other studies, whereby a 95% recovery efficiency of spiked microfibers was observed using 30% H₂O₂ at temperatures of 65 °C for an exposure period of up to 48 h.[9] A shortcoming of this however is possible discrepancies in the recovery rates between studies, resultant of the variation in the nature of
target components (e.g., soft tissue of GI tracts vs hard macroinvertebrate exoskeleton) and chemicals used.

2.4. Considerations for Controlling and Accounting for Microplastic Contamination

The prevalence of contamination in MP research means that QA and QC are paramount throughout all stages of the sample process. As such, rigorous precautions must be adopted while processing field samples, so as to account for any background contamination and provide accurate data. The inclusion of procedural blanks, which follow the same analytical protocol, the use of nonplastic tools where possible, as well as natural fiber clothing when sampling and processing are crucial.

Incidental contamination from airborne particles is a major risk within a MP laboratory, particularly from clothing. While many authors report the use of latex or nitrile gloves and cotton laboratory coats,[24,25,37] only one study from those reviewed, describe the wearing of natural fiber attire. Hermens et al. suggest that the wearing of a 100% cotton laboratory coat alone may not be enough to eradicate contamination should synthetic garments be worn underneath. It is imperative therefore that samples are covered during and between processing steps so as to keep air exposure to a minimum. In the laboratory, field samples are typically covered with aluminum foil or foil lids but may also be covered with watch glasses. Some authors also report keeping air contamination to a minimum by working under positive pressure conditions in laminar flow cabinets, which stream filtered air vertically through the work space, while McHoran et al. reduced the possibility of airborne contamination by opening fish GI tracts one section at a time.

Due to the high background levels of MPs in the laboratory, particularly microfibers, it is vital that workspaces and tools are inspected and cleaned prior to coming into contact with samples. Many studies report cleaning workspaces and tools with ethanol, some using lint free paper, as well with samples. Many studies report cleaning workspaces and tools are inspected and cleaned prior to coming into contact with samples. Many studies report cleaning workspaces and tools are inspected and cleaned prior to coming into contact with samples.

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Procedural blanks are a highly recommended way of accounting for background contamination and isolating steps in the processing stage where contamination can occur. Many of the field studies reviewed have reported running procedural blanks as part of processing to account for exogenous MP particles. Mean MP contamination is generally accounted for based on unique characteristics and may be subtracted from results or rather, taken into consideration when interpreting results.

3. Characterizing Microplastics in Field Studies

3.1. The Role of Microplastic Characteristics in Uptake and Trophic Transfer

As evident from the literature reviewed, a considerable number of organisms are exposed to the possibility of MP ingestion. Not only does the morphology of a particle give indication as to the source of MP pollution, surface characteristics (e.g., polymer, shape, size), combined with organism physiology (e.g., feeding trait and mouth part morphology as well as aperture), may have significant influence on the uptake and residence of MP particles.

Internalization of MP particles has been shown for a variety of freshwater species in laboratory studies including selective filter feeders like Daphnia magna (Straus, 1820) and C. fluminea, as well as fish (e.g., zebrafish Danio rerio (Hamilton, 1822)) Kolandhasamy et al. also found a new pathway of MP uptake, albeit in a marine experiment, via adherence of particles to soft tissues of blue mussels Mytilus edulis (L) without being ingested, something which was further demonstrated by Gutow et al. for the transfer of particles via seaweed to the benthic herbivore Littorina littorea (L). The adherence or entanglement of MPs may therefore play an additional role in the trophic transfer of MP particles through the freshwater food web.

As seen, uptake and prevalence of various polymer types, colors, and shapes (i.e., fragments, fibers, sheets, and spheres) have been detected in a range of freshwater organisms and are therefore considered valuable criteria when assessing potential pathways of MPs and transfer through aquatic food webs. Particle shape is especially pertinent to residence time, given that specific shapes (i.e., fibers) of 1–5 mm particle length have been shown to aggregate in the gut of certain biota (e.g., crustacea), and is reported in all but three studies reviewed. Polymer color is also considered an important descriptor for MP characterization, given that previous research infers a preference for the ingestion of certain colors that potentially resemble prey, and has been described among the majority of freshwater field studies reviewed.

It is noted however that weathering can alter polymer color in the field as can certain oxidation treatments (e.g., H2O2) during isolation, which could possibly result in a certain level of subjectivity in color differentiation during MP characterization.

Therefore, the classification of polymers found in biota in natura should be performed where possible, to further evaluate possible effects of those particles under experimental conditions. It appears however that research groups are often limited, at least from a technological standpoint (e.g., costs, expertise), in the detection or verification of smaller size ranges within field samples. It should also be noted, that immediately after plastic enters the environment biomolecules interact with them forming an ecocorona on surfaces, changing the plastic properties, while a biofilm can also grow on surfaces, affecting ingestion rates among biota. For example Vroom et al. showed a higher uptake probability of experimentally aged MPs containing an ecocorona than pristine particles. This indicates that experimental studies using pristine MP particles...
might underestimate uptake and effects of MPs on biota. Furthermore, as MPs present in the environment undergo transformation processes such as hydrological and UV degradation that influence their surface morphology and behavior,\(^6\) it is important to prioritize which physical (i.e., surface morphology, size, color), as well as chemical characteristics are ecologically relevant.\(^{46,82}\)

3.2. Characterizing Microplastics in Biota: Size Ranges and Associated Challenges

While laboratory experimental set ups are a good way of ascertaining the relative risk imposed by MPs on an organism, as well as the associated impacts thereof, the physical characteristics of MPs ingested in these scenarios were often observed, as outlined earlier, to be significantly different than those documented in natura.\(^83\) Additionally, experimental exposure concentrations were generally at least two orders of magnitude higher than those described in the field,\(^84\) or at best, inclusive of the upper most concentrations reported for freshwater subcompartments (e.g., sediment).\(^54\)

MPs collected in the field are largely characterized by their morphological characteristics such as color, shape, and size, the latter of which is usually measured along the longest axis.\(^42\) Size categories have been used, but discrepancies in the reporting of particle size ranges among freshwater biota combined with an overall lack of standardization in defining micro-, meso-, and macroplastics\(^{20}\) renders comparisons between studies, regions, and biota challenging. McNeil et al.\(^{55}\) investigating the role of species’ trait in MP uptake within riverine fish, categorized MPs into three groups (e.g., small \(<\ 1.5 \text{ mm, medium } 1.6–3.2 \text{ mm and large } > \ 3.3 \text{ mm,} \) while Hu et al.\(^{29}\) in order to facilitate comparisons in MP ingestion among tadpoles, classified MPs into groups according to the size of the mouth aperture (<0.5, 0.5–1, 1–2, or 2–5 mm). In some cases the size of all particles was deemed to be \(< 5 \text{ mm,} \) while in many other studies it was not recorded,\(^{27,36,38}\) or simply not reported.\(^{27}\) While the verification of smaller size ranges from field samples is desired, a number of authors report polymer identification too difficult with available instruments.\(^{29,37,38,55}\) This is not surprising given the current technological limitation in identifying MPs \(< 20 \text{ or } < 100 \ \mu \text{m} \) with most micro-Fourier transformed infrared (µFT-IR) spectrometry systems.\(^{22}\) Tough recent advances (e.g., µ-Raman) do allow for polymer verification of particles \(< 10 \ \mu \text{m.} \)\(^{27,85}\) Until such systems are readily accessible to research groups however, and the technology is improved (e.g., measurement time), accurate reporting of MPs at the smaller size range, as well as the development of suitable experimental studies looking to incorporate relevant MP characteristics at such size limits remain impeded.

3.3. Reporting Microplastics in Environmental Samples

One of the main problems currently within MP research is the reporting of different units for MP quantification which complicates comparisons between studies.\(^{52}\) Within freshwater biotic studies, it is generally the case that the percentage occurrence (\%) of MP items is reported, followed by either the number of items ingested per organism,\(^{9,29–13,34–37,39,41,55}\) MP mg\(^{-1}\) for either dry or wet weight of organism,\(^{24,26–28}\) or number of items per gram of stomach contents (e.g., in fish)\(^{23}\) (see Table I). Some initial studies do not report MP burden in the context of abundances or concentrations at all however.\(^{33,38,40}\) If biota are to be employed as bioindicators of MP pollution within the EU Water Framework Directive and the EU Marine Strategy Framework Directive, units employed should be standardized, at least for comparisons within individual trophic levels. For instance, while it might be feasible for larger consumers (e.g., fish), it is often difficult to report on the number of MPs ingested per individual for smaller organisms (e.g., benthic macroinvertebrates) mainly due to their size and variability thereof, and thus analyzing a subsample comprising of a number of individuals is generally a more practical approach. MP burden in this scenario can be expressed as a concentration of MPs per unit mass (e.g., MPs mg\(^{-1}\)), which is the preferred unit used among most invertebrate studies reviewed.\(^{24,26–28}\) The authors concur with previous recommendations for the reporting of additional units,\(^{52}\) where possible, to ensure that comparison among studies can be undertaken, at least until such time as a consensus can be reached with regard to MP quantitation. Moreover, the (average) size of particles, as well as the relative abundance of the colors and shape of MPs should be reported.\(^{86}\)

4. Field versus Laboratory Studies: Consistencies and Mismatches

Laboratory based studies describe a wide range of effects of MPs on different taxa including effects on morphology and life history traits, such as reduction in food intake and somatic growth, increased mortality, behavior, or altered enzymatic activity.\(^{67–70,87–92}\) However, the lack of observed effects on measured parameters are also described.\(^{54,71,93,94}\) When looking at plastic particle concentrations, morphology as well as the polymer types used in experimental studies, a discrepancy between those used and what has been observed in field studies is apparent. Out of the above-mentioned experimental studies, a discrepancy between those used and what has been observed in field studies is apparent. Of the above-mentioned experimental studies, beads/spheres were the most commonly used morphology type (53%), followed by fragments (35%) and fibers (6%). Whereas field studies assessing MP burden in freshwater biota have reported that fibers are the predominant particle type ingested (ranging from 46.6% to 100%).\(^{28,32,41}\) Reasons for the use of spherical beads in experimental studies are multifarious, though being more easily available compared to other morphology types may be a leading factor. As Ogonowski et al.\(^{89}\) showed that the effects caused by MPs can be attributed to particle morphology, it is suggested that future experimental studies include particle shapes found in higher concentrations in natura to allow for greater ecological relevance.

Further discrepancies in the reporting of background particle concentrations within field studies and those utilized in experimental studies, as well as the units employed, are apparent. Concentrations from field data are reported as number of particles per surface area (e.g., particles m\(^{-2}\)) or volume (e.g., particles m\(^{-3}\)), while laboratory results in most cases are stated as mass per volume, which makes comparison of...
concentrations difficult. Particle concentrations in laboratory experiments ranged from environmentally relevant concentrations (such as 400 particles L\(^{-1}\)) to high concentrations (10\(^8\) particles L\(^{-1}\)), though the latter studies were focused on detecting effect-based thresholds.\(^{[94,95]}\) Whereas field studies have reported concentrations of 0.05 particles m\(^{-3}\) to 3.2 \(\times\) 10\(^5\) particles m\(^{-3}\), with the latter being detected in waste water.\(^{[12,14]}\) This mismatch in the actual concentrations used and units employed has already been addressed by Lenz et al.\(^{[84]}\) for marine systems. The aspect of concentration gains even more importance when one considers that observed effects might not be caused by toxicity of polymers, but either by the sheer number of particles used or by the replacement of nutritious food particles by plastic particles, thus causing a dilution of available food concentration. Animals in aquatic environments are exposed to a diverse array of natural particulate matter. Studies examining the effects by natural particulate matter show similar effects, such as altered feeding rate, survival rate or population growth, which are thought to be caused by MP ingestion.\(^{[96,97]}\) Therefore, interpreting results in order to conclude a reliable and realistic risk assessment of MPs is challenging as most studies did not carry out a particle control with natural particulate matter. The only study we reviewed using such a particle control, such as kaolin or other particulate substances, was implemented by Ogonowski et al.\(^{[89]}\) We therefore suggest the use of naturally occurring suspended solids without a nutritional value as particle controls for pelagic organisms in future studies.

4.1. Recommendations for Assessing Microplastics in Biota within Experimental Setups

MP ingestion has been evaluated under laboratory conditions for a variety of freshwater species.\(^{[54,89,94,95,98–100]}\) some of which being standard toxicity text species.\(^{[101]}\) As mentioned in previous sections, the use of different approaches makes it difficult to establish clear conclusions on the susceptibility to MP ingestion by aquatic biota and the effects derived from this ingestion.\(^{[23,102,103]}\) Here we summarize the most important factors that need to be taken into account during the assessment of MPs in biota and we propose ways to improve the comparison among studies.

4.1.1. Units

Most studies report the amount of MPs in biota as number of particles (ingested/egested) per organism. However, several studies have suggested that MP ingestion can cause energy depletion due to a lower food assimilation.\(^{[54,95,100,104,105]}\) Therefore, ingestion and egestion can also be presented as the dry weight of MPs/weight of biota using the volume and density of the MPs.\(^{[84]}\) As mentioned earlier, we suggest the use of both units when assessing MPs in biota.

4.1.2. Time

The performance of short-term ingestion tests can help to screen a wider number of organisms and identify the ones with a higher capacity to ingest MPs. However, while the ingestion of MPs can rapidly occur, the full gut depuration can often take more than 24 h.\(^{[99,106,107]}\) Therefore, laboratory studies should aim at quantifying ingestion and egestion rates over longer exposure times.\(^{[108]}\) Time will also allow for biofilm growth, which can affect the ingestion rates of aquatic biota and renders more ecologically relevant tests.\(^{[81]}\)

4.1.3. Concentrations

The use of a wide range of concentrations, including environmentally realistic concentrations but also higher concentrations, can provide information on dose-dependent effects and ingestion.\(^{[23]}\) In fact, several studies have shown a positive linear relationship between the amount of MPs ingested and present in the surrounding medium (water/sediment).\(^{[54,98]}\) This relationship can be quantified using trophic transfer factors (TTF), which can be used to compare species, MPs or exposure conditions.\(^{[109]}\)

4.1.4. Bioavailability

MP ingestion is directly related with the bioavailability of the particles, which depend on MP properties such as size, shape and density, as well as on the species traits and experimental conditions, in particular the exposure medium. When it comes to MP properties, ingestion and effects have been found to be size and shape dependent for several aquatic species.\(^{[54,81,98,99,105,110,111]}\) MP density and biofilm (i.e., the composite particle) will determine whether MPs will be found floating, in the water column or in the sediment, which will also depend on the exposure medium and the exposure time. Laboratory studies ideally should therefore use size, shape and density distributions to determine species ingestion preferences and to minimize environmental realism. In the case of size, the lower and upper size limits could be identified and used to correct for the bioavailable sizes, if detection limits allow.\(^{[84]}\) Species traits such as the feeding behavior, mobility, size, and developmental stage can affect MP bioavailability.\(^{[54,94,98]}\) Therefore, the use of a variety of organisms with different traits can provide information on which species are most susceptible to ingest, and be affected, by MP exposure. Additionally, the environmental conditions will also affect the bioavailability of MPs, as the presence of natural particles such as food or sediment has been found to decrease ingestion.\(^{[54,89,98]}\) For this reason, the addition of natural particles to laboratory systems mimicking environmentally realistic situations is of utmost importance.\(^{[103,108]}\)

5. Conclusions

While procedures for the isolation of MPs in some higher-level taxa have been largely established (e.g., GI tracts in fish), little standardization exists for the diverse range of freshwater taxa among lower trophic levels. Standard operating procedures (SOPs) for these processes are needed to optimize extraction efficiencies and increase the representativeness of samples. SOPs could be developed through comparative studies testing a number of digestion techniques on a range of biota that share similar physical characteristics.
It is of equal importance that SOPs for contamination control are implemented. QC/QA measures should include the use of controls (positive and negative), cotton lab coats, reduction or elimination of artificial fibers in the laboratory setting where possible, with further reductions achieved by minimizing air-exposure times and working in laminar flow cabinets.

While laboratory ingestion experiments have provided valuable information regarding the ingestion of MPs, further improvements are required to improve the efficacy of these tests including increasing running times of ingestion/egestion tests to align with complete gut depuration while allowing for biofouling. Though the incorporation of environmentally relevant MP concentrations is deemed increasingly important in laboratory assessment, higher concentrations are essential for the assessment of dose-dependent effects and insight into potential future impacts. In any case, the creation of environmentally realistic situations in these studies, including the inclusion of natural particles will provide the optimal outcome.

By using a wide spectrum of MP characteristics during experimentation (i.e., size, shape, polymer type), combined with taxa that exhibit varied physiological traits (e.g., feeding mechanism), factors previously deemed critical for bioavailability can be accounted for. As well as this, standardization of units reported will further enhance comparisons of studies and improve the pertinence of specific measurements (e.g., mass/volume) in effect threshold and feeding studies.

Technological limitation in the identification of MPs within size ranges relevant to lower trophic levels (e.g., macroinvertebrates), as well as lengthy processing times, necessitates improvements in nondestructive and accessible verification methods within MP research (e.g., focal plane array sensors). These technological advances, in addition to methodological standardization are essential in providing policy makers with tools and measures necessary to determine the distribution of MPs in freshwater ecosystems, while allowing comparisons and providing compliance for future monitoring requirements.

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Conflicts of Interest

The authors declare no conflict of interest.

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