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Using mussel as a global bioindicator of coastal microplastic pollution☆

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ABSTRACT

The ubiquity and high bioavailability of microplastics have an unknown risk on the marine environment. Biomonitoring should be used to investigate biotic impacts of microplastic exposure. While many studies have used mussels as indicators for marine microplastic pollution, a robust and clear justification for their selection as indicator species is still lacking. Here, we review published literature from field investigations and laboratory experiments on microplastics in mussels and critically discuss the suitability and challenges of mussels as bioindicator for microplastic pollution. Mussels are suitable bioindicator for microplastic pollution because of their wide distribution, vital ecological niches, susceptibility to microplastic uptake and close connection with marine predators and human health. Field investigations highlight a wide occurrence of microplastics in mussels from all over the world, yet their abundance varies enormously. Problematically, these studies are not comparable due to the lack of a standardized approach, as well as temporal and spatial variability. Interestingly, microplastic abundance in field-collected mussels is closely related to human activity, and there is evidence for a positive and quantitative correlation between microplastics in mussels and surrounding waters. Laboratory studies collectively demonstrate that mussels may be good model organisms in revealing microplastic uptake, accumulation and toxicity. Consequently, we propose the use of mussels as target species to monitor microplastics and call for a uniform, efficient and economical approach that is suitable for a future large-scale monitoring program.

1. Introduction

Environmental presence and accumulation of plastic debris has become a widespread scientific and social concern due to the dramatic increase in the production of plastics, with an estimate of an additional 335 million tonnes of world plastic production in 2016 alone (PlasticsEurope, 2017). Microplastics (particles less than 5 mm; Arthur et al., 2009) are reported to account for 92.4% of marine plastic debris (Eriksen et al., 2014) and have been identified in many environmental matrices globally. This includes surface waters of every major ocean, the water column, beaches, sea ice, deep sea sediment, marine biota and consumables sourced from the sea (Ng and Obbard, 2006; Browne et al., 2011; Van Cauwenberghe et al., 2013; Cózar et al., 2014; Eriksen et al., 2014; Nor and Obbard, 2014; Lusher et al., 2014, 2015; Van Sebille et al., 2015; Yang et al., 2015; Wesch et al., 2016).

Microplastic ingestion has been identified in a range of species

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from mussels to mammals, with over 220 species from different trophic levels consuming microplastic debris *in natura*, and 99% of all seabird species are predicted to ingest microplastic by 2050 (Wilcox et al., 2015; Hu et al., 2016; Lusher et al., 2017a; Ter Halle et al., 2017). Microplastic ingestion by marine organisms can accelerate microplastics' transference from the sea surface through the water column to the sea floor via feces and marine snow, or between trophic chains via predation (Farrell and Nelson, 2013; Setälä et al., 2014; Katija et al., 2017; Santana et al., 2017). Additionally, microplastics are subject to biofouling leading to colonization by microorganisms and invertebrates, which in turn can contribute to long-range transport of alien species, and serve as reservoirs for pathogen transmission, which broadens the risks of microplastic pollution to marine organisms and ecosystems (Barnes, 2002; Andrady, 2011; GESAMP, 2015, 2016). In addition, environmental weathering of microplastics may also cause release of harmful monomers and additives from the polymer into the associated media (Rochman et al., 2014; Nobre et al., 2015; Gandara e Silva et al., 2016). Together, these aspects represent some of the primary and emerging problems associated with microplastics to date but are by no means the only issues.

Since microplastics are ubiquitous and bioavailable, the associated environmental and health impacts have received an increasing amount of attention amongst the scientific community, regulatory agencies, the public, media and policy makers. Nevertheless, consequences of wild biota interacting with microplastic have not been established, although the current body of evidence from laboratory studies suggests that microplastic exposure may lead to a suite of negative health effects for marine biota; for example, increased immune response, decreased food intake and growth rate, weight loss, energy depletion, apoptosis, upregulation of stress and damage repair pathways and negative impacts on subsequent generations (e.g., Von Moos et al., 2012; Besseling et al., 2013; Canesi et al., 2015; Sussarellu et al., 2016). However, to date most exposure studies have tested unrealistically high doses, and used plastic polymers that are less environmentally-relevant (Phuong et al., 2016), making extrapolation challenging in terms of the microplastic associated risk to the environment. In addition, microplastics' capacity to adsorb, act as vectors of, and leach toxic substances to marine biota may also pose further health risks (Engler, 2012; Browne et al., 2013; Gandara e Silva et al., 2016; Frère et al., 2017).

Despite uncertainties regarding ecological and health risks of microplastic pollution, knowledge based on the wide occurrence of microplastics in the environment has led to calls to classify microplastics as hazardous, and plastic pollution has been compared with climate change in terms of scale and degree of severity by the United Nations Environment Programme (Rochman et al., 2013; UNEP, 2016; Borrelle et al., 2017). From a risk assessment perspective, it is necessary to develop a comprehensive and harmonizated evaluation method of microplastic pollution for inclusion in routine monitoring programs. Traditionally, three marine compartments including water column, sediment and biota could be used to monitor spatial and temporal trends of microplastic abundance. However, microplastic abundances in water and sediment tend to be affected by a variety of environmental factors such as biofilms, bioturbation, tides, winds, currents and wave fronts; all these parameters giving a stochastic pattern, which can complicate the interpretation of impacts on biota (Gibson and Bowman, 2000; Turra et al., 2014; Eriksen et al., 2014; GESAMP, 2015; Moreira et al., 2016a, b; Fisner et al., 2017). In addition, sediment is a more complicated compartment to analyze than water and most biota, including mussels, since sample processing requires multiple steps to degrade organic material and separate microplastics from natural particles. Biomonitoring can be used to address unknowns in terms of risk related to microplastics (Gibson and Bowman, 2000; Wesch et al., 2016).

To have a robust bioindicator for environmental monitoring the following criteria should be fulfilled: a wide distribution range, a well known biology, immobility, an ability to provide an early alert, a key function in the ecosystem, a homogeneous response to pollutants, and the existence of identifiable toxic effects associated with the degree of pollution (Hilty and Merenlender, 2000; Goodsell et al., 2009). Seabirds and sea turtles have been selected as bioindicators for monitoring ingestion of plastic debris (>1 mm) for the land-ocean interaction. For instance, fulmar (*Fulmarus glacialis*) is used as an indicator species in Northern Europe, and its digestive content are currently utilized as an indicator for regional plastic pollution under the OSPAR Convention (Van Franeker et al., 2011). Loggerhead turtles (*Caretta caretta*) have been chosen as a target species to monitor litter presence in the Mediterranean Sea under UNEP-MedPol Convention and Descriptor 10 of the European Union (EU)'s Marine Strategy Framework Directive (MSFD) (Galgani et al., 2014). The suitability of loggerhead turtles as a bioindicator for marine litter >1 mm has been confirmed and is widely supported (Campani et al., 2013; Matiddi et al., 2017; Pham et al., 2017). Although some studies have addressed their proposal for indicator species in microplastic investigation, a robust and clear justification for their selection as indicator species is still scarce (Wesch et al., 2016). Furthermore, the methods currently used for seabirds and turtles are not appropriate for the study of the ingestion of smaller microplastics (<1 mm).

Mussels have been utilized extensively as ideal biological indicators in monitoring of anthropogenic pollution trends in coastal waters due to their special characteristics (Farrington et al., 2016; Beyer et al., 2017). As one of the first animals used to assess the environmental quality of seawater (Goldberg, 1975), mussels meet almost all required criteria for a useful bioindicator species. Firstly, mussels are globally distributed, easily accessible and have a high tolerance to a wide range of environmental parameters including salinity, temperature, oxygen levels and food availability (Bayne, 1976; O'Connor, 1998). Furthermore, as representative benthic filter feeders, mussels can efficiently accumulate chemical pollutants from seawater to provide an integrative measure of the concentration and bioavailability of seawater pollutants *in situ* (Beyer et al., 2017). Mussels provide food (Kautsky, 1981) and habitat (Norling and Kautsky, 2007) to a lot of other species, forming important links between pelagic and benthic ecosystems (Dame, 1993). They also act as a transport route of marine pollutants to higher trophic levels in the coastal marine food chain (Meador et al., 1995; Strand and Jacobsen, 2005). Importantly, mussels have been an important seafood for humans for thousands of years (Beyer et al., 2017). Hence, mussels also attract attention regarding assessing human health risks associated with marine pollution (Van Cauwenberghe and Janssen, 2014; UNEP, 2016). Up to now, mussel has been widely used in many regional environmental monitoring programs such as the U.S. Mussel Watch Project, Assessment and Control of Pollution in the Mediterranean region (MEDPOL), and OSPAR's Coordinated Environmental Monitoring Program (CEMP) (Beyer et al., 2017).

In this review, both the suitability and challenges related to mussels as bioindicator for microplastic pollution will be discussed. We aim to address (i) why mussels lend themselves as good indicators of microplastics; (ii) the extent to which mussel can provide useful information regarding microplastics pollution in the marine environment; and (iii) how to improve current methodology, with an emphasis on standardization of techniques to allow cross calibration between studies worldwide.

2. Global field investigations on microplastic pollution in mussels

Environmental risks associated with microplastics are primarily focused on their suspected bioavailability for marine organisms (Wright et al., 2013; Desforges et al., 2015). Bivalves are of particular interest because their extensive filter-feeding activity exposes them directly to microplastics present in the environment. Globally, microplastic occurrences in wild caught mussels have been extensively investigated and reported (Table 1).

2.1. Selected species and geographic coverage

Mussels (*Mytilus* spp.) are currently the dominant species used for field investigations of microplastics. *Mytilus* has seven subspecies that can interbreed with each other and are widely distributed around the world (Beyer et al., 2017). For instance, *M. galloprovincialis* has become an invasive species and is widely spreaded in South America, South Africa, Japan, California, New Zealand, and Australia (Beyer et al., 2017). Different species within the genus *Mytilus* have different genomic composition and gene expression profiles, which may lead to differences in the way they deal with stress as well as microplastic uptake (De Witte et al., 2014; Lusher et al., 2017b). *Mytilus* spp. have been investigated in all the involved countries except Brazil and Indonesia, which investigated *Perna viridis* and *P. perna* instead (Table 1, Fig. S1).

Spatially, field investigations of microplastics in mussels are currently spread over 16 countries (Fig. S1), especially in European countries including Germany, France, Belgium, the Netherland, Italy, Greece, Portugal, Spain, Denmark, Finland, Norway and the U.K. In addition, research from China, Indonesia, Canada and Brazil also contribute to the available field data. Research on microplastic can be traced back to 1970s when the occurrence of small plastic particles in coastal environment was first reported (Bowmer and Kershaw, 2010). At that time, small polystyrene beads in New England (Carpenter et al., 1972), Sargasso Sea (Carpenter and Smith, 1972) and Bristol Channel (Morris and Hamilton, 1974) attracted researchers's attention. Afterwards, the term “microplastic” were put forward for the first time by Thompson in Europe (Thompson et al., 2004). Currently, the monitoring of marine litter is required as part of the EU Marine Strategy Framework Directive (MSFD) (Hanke et al., 2013) and many projects fund research on microplastic pollution in Europe such as Marine Litter Projects Funded under FP7 and Horizon 2020, which likely accounts for the increased number of studies from Europe.

2.2. Characteristics of microplastic pollution

It is indisputable that microplastics are widespread in both wild and farmed mussels in many countries (Table 1). Regarding the morphotypes of microplastics observed in such mussels, fibers are dominant in 13/27 of the current filed investigations compared with fragments which account for 5/27. Only one paper reported the prevalence of pellets (Murphy, 2018), although some researchers have highlighted the possibility for misinterpretations of pearls in place of beads and/or pellets (Bråte et al., 2018b). The remaining studies counted one type of microplastics due to methodological limitations or omitted to report the proportion of different types. Polyethylene, polypropylene, polystyrene, polyester, polyethylene terephthalate, polyamide, polyvinyl chloride and cellophane were the most reported polymers. Out of the studies conducted, nine of them did perform a corresponding investigation of the microplastic level in the associated sediment or seawater (Table 1). From these, it appears that the main morphotype and polymeric composition in mussels tend to be consistent with their surrounding environmental media (Leslie et al., 2017; Li et al., 2018; Qu et al., 2018; Digka et al., 2018a; Railo et al., 2018). Furthermore, Qu et al. (2018) observed consistency of their proportion in mussels and in seawater. These results suggest that the microplastics in mussels can reflect the real pollution status in the environment in terms of morphotype and polymer types.

Currently, the lower size limit for microplastic detection in mussels is 5 μm, yet some studies fail to provide this information (Table 1). The lower size limit often depends on the methodology employed by research teams. Selected research to date have adopted a classified size range approach and in doing so have highlighted a dominant smaller size range (e.g., 5–250 μm, 10–300 μm, 50–100 μm, 50–250 μm,100–500 μm, 0.25–1 mm) that reveals mussel's uptake incidences for specific size ranges (Leslie et al., 2017; Li et al., 2018; Phuong et al., 2018a; Kolandhasamy et al., 2018; Qu et al., 2018; Digka et al., 2018b). However, the lack of a unified classification standard for reporting the size range complicates efforts to compare these results. In addition, smaller size of microplastics seems to take up a larger proportion in mussels compared to the surrounding environmental media (Li et al., 2018; Qu et al., 2018; Digka et al., 2018a). For example, the smaller microplastics (<1 mm) account for 62.3%, 96.9%, 100% in seawater, sediments and mussels from the Northern Ionian Sea respectively (Digka et al., 2018a) and the mussels from U.K. contained 44%–83% smaller microplastics (less than 250 μm) compared to seawater with only 30%–40% (Li et al., 2018). Another interpretation is thus that the microplastics in mussels indicates the size range in the surrounding environment partially as a factor of their selective feeding behavior (Ward and Shumway, 2004).

Microplastic abundance varies between different studies, ranging from 0.05 items/g to 259 items/g (Table 1). These levels may either represent regional variation in microplastic abundance, the diversity of method used, or in some case background contamination which has not be accounted for. On a broad scale, research has demonstrated a positive correlation between coastal microplastic concentrations and human population density (Browne et al., 2010, 2011). Furthermore, microplastic abundance in mussels is closely related to human activity, and mussels from areas with intensive human activities contain significantly higher numbers (Li et al., 2016), or in areas suggested to have accumulation zones of microplastics such as the Barents Sea (Lusher et al., 2017b). There are indications that microplastics can accumulate because significantly higher concentrations have been found in mussels (3.7×10^4 items/kg dry weight) compared to surrounding sediment (48 items/kg dry weight) and seawater (27 items/L) (Karlsson et al., 2017). When we unify the units of the abundance in mussels as items/g.w and in seawater as items/L, similar abundances can be found in mussels and ambient seawater (Table 1, Van Cauwenberghe et al., 2015; Karlsson et al., 2017; Li et al., 2018; Qu et al., 2018), which is further supported by a recent study that showed a positive and quantitative correlation of microplastics in mussels and in their surrounding waters (Qu et al., 2018). This indicates that microplastic pollution in mussels is closely correlated with the degree of pollution in coastal habitats and can reflect the real abundance of microplastics in the environment within certain size range. However, one study does not show the quantitative correlation between microplastics in mussels and their ambient seawaters (Li et al., 2018), this may be due to limited sampling sites and outliers derived from contingency. More studies are still needed to verify this outcome.

2.3. Methodological challenges

Procedures for investigating microplastic pollution in mussels involve a series of steps and details that must be taken into

Table 1
Summary of global field investigations on microplastics in mussels.

Species & Location	Digestion method	Identification technique	Classification	Abundance (items/g.ww)	Size (μm)	Environmental media	Reference
Mytilus. edulis							
Canada	30% H_2O_2	visual sorting	fiber	2.79–7.42 [a]	no data	sediments: 2–8 items/g.dw	Mathalon and Hill, 2014
Germany	69% HNO_3	micro-Raman	particle	0.36 ± 0.07	no data	no data	Van Cauwenberghe and Janssen, 2014
Belgium	HNO_3:$HClO_4$	visual sorting	fiber, fragment, film, sphere	0.26–0.51	200 –1500	no data	De Witte et al., 2014
France, Belgium, Netherlands	69% HNO_3	micro-Raman	particle	0.2 ± 0.3	20–90	seawater: 0.4 ± 0.3 items/L sediments:6 ± 5.7 items/kg.dw	Van Cauwenberghe et al., 2015
UK	trypsin	FTIR	fiber, bead, fragment, film	1.05–4.44	200 –10670	no data	Courtene-Jones et al., 2017
UK	Corolase 7089 enzyme	FTIR	fiber, particle, film	2.5	no data	no data	Catarino et al., 2017
UK	30% H_2O_2	micro-FTIR	fiber, fragment, sphere, flake	0.7–2.9	8–4700	seawater:1.5–6.7 items/L	Li et al., 2018
Netherlands	proteinase K and 30% H_2O_2	Raman	fiber, particle	37 (items/g.dw)	30–2000	seawater: 27 items/L sediments: 48 items/kg.dw	Karlsson et al., 2017
Netherlands	HNO_3, NaOH & 30% H_2O_2	FTIR	fibre, sphere, foil	19-105 (items/g.dw)	10–5000	sediments: 100 –3600 items/kg.dw	Leslie et al., 2017
France	10% KOH	micro-FTIR	filament, fragment	0.23 ± 0.20	20–400	no data	Phuong et al., 2018a
France	10% KOH	micro-FTIR	fiber, fragment	0.23 ± 0.09	30–200	no data	Phuong et al., 2018b
Canada	68–70% HNO_3	FTIR	fiber, fragment, pellet	wild:138 ± 202 farmed:259 ± 114	<530	no data	Murphy, 2018
China	30% H_2O_2	micro-FTIR	fiber, fragment, sphere, flake	2.2	5–5000	no data	Li et al., 2016
China	30% H_2O_2	micro-FTIR	fiber, sheet, fragment, sphere,	9.2 [b]	50–5000	no data	Kolandhasamy et al., 2018
China	30% H_2O_2	micro-FTIR	fiber, fragment, pellet	1.52–5.36	5–4000	seawater: 0.68 –6.44 items/L	Qu et al., 2018
M. galloprovincialis							
Italy	30% H_2O_2	visual sorting	filament, fragment	0.05 (items/g.dw)	60.01 ± 38	no data	Bonello et al., 2018
Italy	30% H2O2	visual sorting	filament	6.2–7.2[c]	750 –6000	no data	Renzi et al., 2018
Italy, Portugal, Spain	69% HNO_3	visual sorting	fiber, particle	0.12 ± 0.04	no data	no data	Vandermeersch et al., 2015b
Italy, Portugal, Spain	HNO_3:$HClO_4$	visual sorting	fiber, particle	0.18 ± 0.14	no data	no data	Vandermeersch et al., 2015b
Greece	30% H_2O_2	FTIR	filament, fragment, film	46.25% ingested microplastics	<5000	seawater: 0.41 items/m^2 sediments: 1816.7 items/m^2	Digka et al., 2018a
Greece	30% H_2O_2	FTIR	fiber, fragment	wild:5.3 ± 0.5[d] farmed:2.5 ± 0.3[d]	40–737	no data	Digka et al., 2018b
China	30% H_2O_2	micro-FTIR	fiber, fragment, pellet	2.39 ± 1.32	5–5000	no data	Li et al., 2015
M. trossulus							
Finland	Sodium Dodecyl Sulphate (SDS) and detergent enzymes	FTIR	fiber, fragment, sphere, flake	0.4 ± 1.9	>20	seawater: 11.4 –23.5 items/m^3	Railo et al., 2018
***Mytilus* spp.**							
Norway	10% KOH	micro-FTIR	fiber, foam, fragment, film	1.85 ± 3.74	150 –8010	no data	Lusher et al., 2017b
Norway	10% KOH	micro-FTIR	fiber, foam, fragment, film	0.97 ± 2.61	70–3870	no data	Bråte et al., 2018b
UK	Corolase® 7089 enzyme	Nile Red staining and FT-IR	fiber, film, sphere, other particle	3 ± 0.9	200 –2000	no data	Catarino et al., 2018
Italy, Netherlands France, Denmark, Spain, Portugal	HNO_3:$HClO_4$	visual sorting	fiber, particle	0.13 ± 0.14	no data	no data	Vandermeersch et al., 2015b

(*continued on next page*)

Table 1 (*continued*)

Species & Location	Digestion method	Identification technique	Classification	Abundance (items/g.ww)	Size (μm)	Environmental media	Reference
Modiolus modiolus							
UK	Corolase® 7089 enzyme	Nile Red staining and FT-IR	fiber, film, sphere, other particle	0.086 ± 0.031	200 −2000	no data	Catarino et al., 2018
Perna perna							
Brazil	22.5 M HNO_3	visual sorting	fiber, irregular particle	75% ingested microplastics	no data	no data	Santana et al., 2016
P. viridis							
Indonesia	30% H_2O_2	SEM/EDX[e]	fiber, fragment, sphere, flake	4–20	51.31 −232	no data	Khoironi and Anggoro, 2018
China	30% H_2O_2	micro-FTIR	fiber, fragment, pellet	1.52–5.36	5–4000	seawater: 0.68 −6.44 items/L	Qu et al., 2018

[a] The microplastic level was transferred by dividing total microplastics per individual by the shelled weight.
[b] The abundance of microplastics in intestine.
[c] The abundance of microplastics in hepatopancreas and gills.
[d] The abundance of microplastics in digestive glands and gills.
[e] Scanning Electron Microscopy/Electron Dispersive X-Ray.

consideration including: sampling sites and strategy, sample size (number of individuals per site), individual condition, sample storage, digestion solution, filter pore size, chemical identification techniques, classification of microplastics, reporting units, and contamination control. Although many reviews have systematically and critically discussed existing microplastic extraction methods and identification techniques, there is still a lot of debate and many knowledge gaps surrounding choices of an optimal method (Hidalgo-Ruz et al., 2012; Lusher et al., 2017a; b; Elert et al., 2017; Shim et al., 2017). Variations in methods make it hard to compare microplastic contamination among different studies and locations (Vandermeersch et al., 2015b).

Hence, a major challenge for monitoring microplastic pollution within mussels is the lack of uniform methods from extracting to identifying microplastics. Call for the standardization or harmonization of methods are repeatedly highlighted by the International Council for the Exploration of the Sea (ICES) and researchers working within the field (Hidalgo-Ruz et al., 2012; ICES, 2015; Wesch et al., 2016; Lusher et al., 2017a; b; Rochman et al., 2017). Since these methods always have a tension between accuracy, precision and feasibility, different approaches should be chosen according to the sampling sites, media, equipment, replicates request and the specific scientific questions of interest (Rochman et al., 2017). In this situation, we suggest that both standardization and intercalibration of different methods should be adopted at the same time for improving the comparability of different studies. Some factors could be united while other variables should be intercalibrated and selected according to the actual situation in the specific procedure.

Sampling strategy represents a challenge in designing a representative and adequately replicated monitoring scheme. Patchiness of microplastics in different spatial (Browne et al., 2011; Moreira et al., 2016a; Fisner et al., 2017) and temporal (Moreira et al., 2016b) scales may lead to variable amounts within mussels. Phuong et al. (2018a) showed the season was not a relevant influencing factor on the quantitative and qualitative analysis of microplastics in mussels. However, a different conclusion revealed the similarity of microplastic types and significant differences of abundance in mussels collected in different seasons (Catarino et al., 2018). That is to say, some factors changing with season (e.g., wind, currents, rainfall, temperature, human activity) may affect microplastic distribution. The extent to which these factors change microplastic abundance or type in the environment varies with sampling sites. Sampling time and sites should be variable factors considered during the investigation; such that harmonization of sampling strategy should take these complex environmental and anthropogenic factors that shows temporal and spatial differences into consideration. Additional factors such as sample number and preservation method must also be standardized. Both ICES and MSFD recommend 50 individuals per species, although research suggest 20 individuals could also be a suitable number for large-scale spatial investigations (Lusher et al., 2017b). Finally, but definitely most importantly, procedural contamination should be minimized throughout the sample preservation and identification processes.

For the extraction method, common agents used to digest biotic tissues include acid (HNO_3, HNO_3:$HClO_4$), alkaline (NaOH, KOH), oxidizing (H_2O_2) and enzymatic (trypsin, proteinase K, Corolase 7089) approaches. However, drawbacks of these digestion methods have been widely reported, such as structural damage, dissolution and discoloration caused by acid, basic and H_2O_2; incomplete soft tissue digestion by enzyme; production of foam caused by H_2O_2; expensive price and time-consuming nature of some of the solvents (Table 1, Lusher et al., 2017b). This might lead to underestimations of microplastic loads, especially smaller particles, or limit their adaptability for large scale monitoring. Hence, selection of a digestion solution requires further testing and optimization.

In future investigations, different digestion agents could be chosen under the premise that the selected agent does not destroy the main polymer types in the objective environment, which requires consulting literature or preliminary research. In addition, the digestion efficiency and recovery rate should be provided for the intercalibration of methods. However, only ten published studies report corresponding recovery rate and five tested polymer alterations by digestion treatment (Table 1). Low digestion efficiency and recovery rate may lead to underestimations of microplastics, therefore, a threshold for both efficiency is required.

The pore size of the filter, the magnification and resolution of microscopy employed determine the size resolution of observed microplastic. ICES has recommended the use of filter with 5 μm pore size for mussel (Vandermeersch et al., 2015b). In the current literature, 5 μm pore size of filter has been the most frequently used (9/27). Other studies had finer (0.45, 0.7, 0.8, 1.2, 2.5, 2.7 μm) or bigger (12, 20 μm) size. Among all the given size ranges of microplastics detected in mussels, 5 μm is the minimum size (Table 1). Although smaller sized microplasctics are present in mussels, their observation and identification are still limited by current instrumentation and methodology. For example, 20 μm seems to be the smallest size that could be identified using μFTIR in the reflection mode under manual inspection (Phuong et al., 2018b). Hence, 5 μm

could be used for the unity of pore size of filter. The detection limit of current methods will not hamper the use of mussels as a bio-indicator of microplatic pollution since a quantitative correlation of microplastics within certain size range in mussels and in their surrounding waters has been demonstrated (Qu et al., 2018).

Current methods for microplastic identification involve visual sorting (with the aid of polarized light microscopy), Nile Red staining, Fourier transformed infrared spectrometry (FT-IR), attenuated total reflectance (ATR), Raman spectrometry, pyrolysis-gas chromatography combined with mass spectroscopy (Pyr-GC-MS), high temperature gel-permeation chromatography (HT-GPC) with IR detection, SEM-EDS, thermal extraction desorption gas chromatography mass spectrometry (TED-GC-MS) and liquid extraction. FT-IR is the most commonly used technique in recent literature (Table 1). Each applied technique has some drawbacks including size limitations, time constraints and interference factors and we refer the readers to published literature on the advantages and limitations of these methods (Shim et al., 2016, 2017; Elert et al., 2017; Lusher et al., 2017b). Since no single method is able to obtain the physical (size, shape and colour) and chemical (polymer type) characteristics of particles in a single step, the combination of several parallel approaches should be applied and considered in future research. Meanwhile, intercalibration between different methods is necessary to understand the extent to which each method differs and compare the data already collected with that in future studies.

Preliminary visual sorting is still needed for a quantification analysis. Nevertheless, the result is largely dependent on personal experience which may result in underestimation or overestimation of real results to different degree. A library matching the photos of environmental samples with their spectrograms should be established to help reduce error rates and misidentification and improve this method. For future, small-scale investigations, FT-IR and Raman are strongly recommended with 70% match rate as a standard threshold which has been applied in most research. However, spectra libraries still require intercalibration. For future large-scale investigation, Nile Red staining and thermo-analytical technique could be combined to obtain both qualitative and quantitative information efficiently. However, the accuracy of Nile Red staining should be calibrated using spectroscopy methods simultaneously.

The variability in the way the results are characterized further hampers the comparision among different studies. These factors such as reporting units, classification of type and size range should be standardized in future studies. Both items individual^{-1} and items gram^{-1} as reporting units are required. The latter is a more appropriate unit to compare different studies and it has been used most commonly in current research (Table 1). For the classification of type, four kinds including fiber (filament), fragment, sphere (pellet, bead), film (flake, sheet) could be adopted which almost covers all the types in current studies (Table 1). An optimal classification of size range still requires more research to determine. In addition, contamination control is a crucial factor during the whole procedure. Procedural blanks must be carried out to monitor contamination and correct empirical data. Most of the current investigations (25/27) set procedural blanks. Two studies even tested limit of detection of airbore fibers (De Witte et al., 2014; Bråte et al., 2018b).

3. Laboratory exposures of microplastics in mussels

3.1. Uptake, accumulation and clearance of microplastics

In addition to field studies, mussels have been widely used in laboratory exposure experiments to study uptake, accumulation, clearance characteristics and impact of microplastics. Microplastic uptake has been demonstrated in all exposure concentrations (Table 2), and egestion as feces and pseudofeces has also been observed (Ward and Kach, 2009; Wegner et al., 2012; Khan and Prezant, 2018; Santana et al., 2018). During active feeding, mussels can continuously pump and filter seawater through coordinated action of cilia localized at the gill epithelium surface, at a rate of 50 ml of seawater per minute (Famme et al., 1986).

According to mussel feeding strategies and laboratory exposure studies, we can hypothesize pathways of microplastic intake and accumulation as follows. When microplastics in seawater encounter gill surfaces, they may be captured and trapped into mucus and subsequently assimilated over the gill epithelium or transported into the mouth and digestive system (Von Moos et al., 2012; Beyer et al., 2017; Bråte et al., 2018a; Kolandhasamy et al., 2018). Not every particle captured by gills is ingested (Santana, 2015; Santana et al., 2018) since mussels are able to separate and reject nonnutritive particles as pseudofeces as a way to defend organisms against high quantities of suspended particulate matter (Ward and Shumway, 2004).

Von Moos et al. (2012) demonstrated that mussels can ingest and accumulate microplastics (0–80 μm) in digestive system epithelial cells within hours. It appears that smaller particles are ingested and retained in mussels more easily compared to the larger particles (Van Cauwenberghe et al., 2015). However, behavior of PVC particles in an emulsion/microsuspension (E/M PVC; size range of 0.1–1.0 μm in diameter; Rodolfo et al., 2006) was different, with larger particles proportionally better represented in mussel digestive glands (0.8–0.96 μm) in comparison to surrounding water (mean size, 0.6 μm). Van Cauwenberghe et al. (2015) found that larger sized (15–500 μm) microplastics were detected in mussel's faeces compared to mussel tissue (20–90 μm). These findings indicate mussel's selection for a specific size range of microplastics during ingestion and egestion process, which is consistent with the results of the field investigations discussed in section 2.2. However, this selectivity characteristic poses an obstacle to the reflection of size distribution of microplastics in the environment through biomonitoring. More research is needed to test selectivity of mussels for larger scope and more gradient sizes of microplastics.

In addition to size variation, environmentally aged microplastics are differentially ingested with pre-weathered microplastic ingested to a higher extent by mussels compared with virgin microplastic (Bråte et al., 2018a). In most exposure studies, only particles or spheres were used for the exposure (Table 2), which ignores the selectivity of mussels for microplastics of different shapes. Qu et al. (2018) showed fibers were dominant in mussels from field investigation while beads were most ingested by mussels after five-day indoor exposure. One explanation is that fibers in mussels result from long-term accumulation in the marine environment, while beads are more easily ingested by mussels in short time periods. Once ingested, beads could be egested more quickly than fibers. The delay in egestion of synthetic fibers has been addressed since only fibers were detected in mussels after gut clearance period (De Witte et al., 2014). Moreover, fibers trapped into gills and hepatopancreas cannot be easily removed by individuals (Renzi et al., 2018).

It has been suggested that microplastics accumulating in mussels will achieve a dynamic balance between ingestion and clearance and become stable (Li et al., 2016; Setälä et al., 2016), despite their selective feeding strategy and varying retention time of different size/morphotype particles (Ward and Kach, 2009; Farrell and Nelson, 2013). Not only has a positive and quantitative correlation of microplastics in mussels and in their surrounding waters from field investigations been reported (Qu et al., 2018), but similar results from laboratory exposure experiments have been found.

Table 2
Uptake and accumulation of microplastics by mussels in laboratory exposures.

Exposure microplastic			Exposure concentration	Exposure time	Uptake and accumulation organs	Reference
Types	Shapes	Sizes				
Mytilus edulis						
PS	spheres	3, 9.6 μm	42 particles/L	3 h-48 d clearance	gut, haemolymph	Browne et al., 2008
PS	particles, beads	100 nm, 10 μm	1.3×10^4 particles/ml and 1000 beads/ml	45 min-72 h clearance	digestive gland	Ward and Kach, 2009
HDPE	powders	0–80 μm	2.5 g/L	96 h	gill, stomach, digestive gland	Von Moos et al., 2012
PS	beads	30 nm	0.1,0.2,0.3 g/L	8 h	foot	Wegner et al., 2012
PS	spheres	10, 30, 90 μm	110 particles/ml	14 d-24 h clearance	whole soft tissue	Van Cauwenberghe et al., 2015
	beads, fragments and fibers		100,1000 particles/L	5 d	whole soft tissue	Qu et al., 2018
	fibers		2000 microfibers/L	48 h-48 h clearance	gill, intestine, foot, stomach, mantle, gonad, adductor visceral tissue	Kolandhasamy et al., 2018
PS, PE, PP	beads, fibers	7–30 μm (beads) or 23 × 3000 μm (fibers)	50 beads/ml or 0.1 fibers/ml	60 min	whole soft tissue	Porter et al., 2018
M. galloprovincialis						
PS, PE	powders	<100 μm	1.5 g/L	7 d	haemolymph, gill digestive gland	Avio et al., 2015
LDPE	particles	20–25 μm	2.34×10^7 particles/L	28 d	hemolymph, gills, digestive glands, intestine	Pittura et al., 2018
PE	fragments (derived from toothpaste)	50–590 μm	0.01 g/L	21 d	digestive tract, whole body	Bråte et al., 2018a
PS	spheres	3 μm	50-1 × 10^4 particles/ml	24 h-192 h clearance	gut of larva	Capolupo et al., 2018
***Mytilus* spp.**						
PS	beads	2, 6 μm	32 μg/L/day = 2000 beads/ml/day	7 d-7 d clearance	digestive tract intestine, gills	Paul-Pont et al., 2016
Dreissena polymorpha						
PS	beads	1, 10 μm	1×10^6 or 4×10^6 particles/L	6 d	gut, digestive gland, haemolymph	Magni et al., 2018
Geukensia demissa						
PS, PE	spheres	5, 250–300 μm	3.467 g/L	2 h-24 h clearance	stomach, digestive tubules, intestine	Khan and Prezant, 2018
Perna perna						
PVC	spheres	0.1–1 μm	0.5 g/L	3 h-12 d clearance	gut, haemolymph	Santana et al., 2017

Abbreviations: PS, polystyrene; PE, polyethylene; HDPE, high-density polyethylene; LDPE, low-density polyethylene; PP, polypropylene; PVC, polyvinyl chloride.

The abundance of microplastics in mussels was significantly higher in the high concentration exposure group than the low concentration group (Qu et al., 2018) and a significant and linear increase of microplastic uptake in mussel larva with increasing exposure concentrations was observed (Capolupo et al., 2018).

Microplastics can be taken up over the digestive surface of mussels gastrointestinal tracts by endocytosis and granulocytomas and then transferred to lysosomes and circulatory system or eliminated as pseudofaeces particles, which contributes to microplastic adherence to the foot and mantle (Browne et al., 2008; Von Moos et al., 2012; Wegner et al., 2012; Beyer et al., 2017; Kolandhasamy et al., 2018; Khan and Prezant, 2018). Browne et al. (2008) showed the ability of mussel to ingest polystyrene microspheres between 3 and 10 μm in size and to transfer them to the circulatory system, where smaller particles appeared to undergo translocation more readily than larger ones. Assimilation of very small particles of emulsion/microsuspension PVC (~1 μm) was also recorded for *P. perna* (Santana, 2015; Santana et al., 2017). Assimilation of small particles contributes to their accumulation in mussels relatively steadily. This may explain why, after a three day gut clearance, only larger particles (>20 μm) were egested completely, whilst smaller particles (5–20 μm) were still present (Van Cauwenberghe and Janssen, 2014).

Theoretically, small particles or beads should account for a larger proportion due to their assimilation. However, fibers were always dominant in field investigations as mentioned in section 2.2. This could be explained by the limitation of current methodology. Van Cauwenberghe et al. (2015) demonstrated that only microplastics of the smallest size (10 μm) was detected in mussles although three sizes (10 μm, 30 μm, 90 μm) of microplastics were used in the exposure experiment. Furthermore, the size of microplastics reported to occur in haemolymph (e.g., 0.1–1 μm, 3 μm, 9.6 μm, 10 μm, 20–25 μm, Table 2) tend to be close to or smaller than the detection limit of field investigation method. Therefore, a large proportion of these small particles are unlikely to be detected in field surveys. Even so, laboratory exposures of these smaller microplastics contribute to our understanding of accumulation of microplastics in mussels and relative toxicology effects.

The total body burden of microplastics in mussels goes beyond ingestion. Besides uptake through the gut and across the gills, microplastics adhered to mussel's soft tissue (mantle, gonad, adductor, visceral tissue and foot) can further contribute to microplastic presence within individuals. This has been verified in both field and laboratory environments (Von Moos et al., 2012; Kolandhasamy et al., 2018). Since mussels are eaten whole by both animals and humans, Microplastics can also be passed to higher trophic levels following predation, as demonstrated in laboratory exposure experiments (Farrell and Nelson, 2013; Watts et al., 2014; Santana et al., 2017).

At present microparticle behavior within mussel tissues is still largely unknown; this includes translocation into, and from, haemolymph to other tissues as well as depuration and egestion rates.

Studies have shown that microplastics may be retained for extended periods of time, for example, complete clearance of microplastics was not achieved after a seven-days depuration period under laboratory conditions with microbeads (2,6 μm) being retained within the digestive tracts (Paul-Pont et al., 2016). In addition, microplastics were remained in the haemolymph of *M. edulis* 48 days after exposure (Browne et al., 2008), however, there was a reduce in microplastic numbers over time which suggested egestion was occurring. These results suggest that mussels are effective indicators of recent exposure. Although efficient gut clearance and selective feeding behavior of mussels limit their quantitative ability as indicators of microplastic. For example, the only available data on retention refers to those that have been selected by mussels, especially in terms of size. Microplastics in mussels can still reflect the abundance, polymer type and morphotype of microplastics in the environment when sampling and thereby come a bit closer to the risk assessment.

3.2. Toxic effects of microplastics

One of the criteria for a sentinel species is that adverse effects can be observed following exposure. In terms of microplastic toxicity, a number of effects have been reported. It should be noted that most of these studies use unrealistically high exposure levels. Notable histological changes in mussel digestive cells, strong inflammatory responses with formation of granulocytomas, and lysosomal destabilization which increases with exposure time, have all been observed (Von Moos et al., 2012). Avio et al. (2015) demonstrated cellular effects including alterations of immunological responses, lysosomal compartmentalisation, peroxisomal proliferation, antioxidant system, neurotoxic effects, onset of genotoxicity, and changes in gene expression profile associated with microplastic exposure. Bråte et al. (2018a) found histological alterations in gills and digestive tissue, and hemocytic aggregates in the digestive gland following exposure to PE fragments (ranging from 50 to 590 μm) extracted from toothpaste. On a nanoplastic scale, mussels showed reduced filtering activity, and the total weight of the feces and pseudofeces increased with the increase of nano PS (30 nm, Wegner et al., 2012). Furthermore, PS-NH2 particles stimulated increase in extracellular reactive oxygen species and nitric oxide production and induced apoptotic process of hemocytes (Canesi et al., 2015). Finally, Gandara e Silva et al. (2016) showed the toxic effect of leachates of virgin PP and beached plastics pellets caused mortality and abnormal embryos of *P. perna*.

In summary, the reported effects of microplastic uptake include histological changes, inflammatory response, lysosomal membrane destabilization, reduced filtering activity, neurotoxic effects, alterations of antioxidant system, increase in hemocyte mortality, dysplasia, genotoxicity and transcriptional responses (Table S1). These research results lay a good foundation for the exploration of specific biomarkers for microplastic pollution.

3.3. Optimization of laboratory exposures

It should be highlighted that in many laboratory studies, organisms are exposed to unrealistically high doses of microplastics with uniform size or shape, in virgin condition, and for relatively short time frames (Rochman et al., 2016; Koelmans et al., 2017; Lambert et al., 2017). Whereas, environmentally exposed plastics are subject to weathering, abrasion and photodegradation, therefore comprising of a broad size distribution and various shapes (Phuong et al., 2016; Lambert et al., 2017). In addition, weathering processes may weaken the plastic surface, enhance chemical leaching and change the outcome of toxicological investigations of microplastic particles (Ogonowski et al., 2016; Lambert et al., 2017; Potthoff et al., 2017).

In some studies, mussels were caged in specific areas for extended periods to investigate the microplastic pollution related to specific anthropogenic activity, such as the removal of wreck or to assess seasonal changes in plastic pollution (Avio et al., 2017; Catarino et al., 2018). To mimic environmental weathering, some studies exposed organisms to microplastics collected from beaches or deployed in a bay for a period time (Gandara e Silva et al., 2016; Nobre et al., 2015; Rochman et al., 2014; Bråte et al., 2018a). Furthermore, a photo-oxidative degradation of plastic pellets incubated in seawater, ultrapurewater and air with UV irradiation over a three-month period observed some changes in hydroxyl groups, carbonyl groups and surface textures which provides a good foundation for making environmental microplastics under laboratory conditions (Cai et al., 2018).

A recent study using weathered PE particles from toothpaste showed that following a chronic exposure (21 days) with lower dose than normally tested (~1 particle per ml), still induces tissue alterations in mussels (Bråte et al., 2018a). In contrast, long-term exposure (90 days) of *P. perna* to a less extreme concentration compared with previous studies (0.125 g/L) indicated no behavioral and physiological effects of microplastics (Santana et al., 2018). Calls for more testing on toxicological effects of long-term exposure to environmentally realistic concentrations and shapes are repeatedly made by the scientific community (Van Cauwenberghe et al., 2015; Phuong et al., 2016; Koelmans et al., 2017). Furthermore, Connors et al. (2017) and Karami (2017) provide guidance which should be considered to improve the quality and reliability of ecotoxicological studies of microplastics. This includes the characterization (physical and chemical properties) and quantification of microparticles in future laboratory exposure studies to facilitate a comprehensive understanding of the causal links between physical-chemical properties of microplastic particles and toxic effects (Connors et al., 2017).

4. Scope of mussels as global bioindicators of microplastic

4.1. Advantages of utilizing mussel

There is a consensus that mussels make good biological indicators for monitoring many anthropogenic pollutants (Beyer et al., 2017). Besides the advantages discussed above, mussels also have specific advantages as bioindicator for microplastic pollution. Feeding type affects microplastic ingestion, for example, filter-feeding makes bivalves ingest more microplastics (Setälä et al., 2016). Mussels as species susceptible for microplastic uptake have been documented widely (e.g., Browne et al., 2008; Von Moos et al., 2012; Mathalon and Hill, 2014; Santana et al., 2016; Li et al., 2016). Furthermore, potential contamination during sampling and laboratory processing is a key problem in microplastic research, mussel's hard shells and easy handling minimize contamination risk (Setälä et al., 2016; Beyer et al., 2017). Bivalves are likely the largest source of microplastics from seafood to humans because they are consumed whole (Lusher et al., 2017c). This adds to their selection as ideal indicators for microplastic pollution monitoring.

Furthermore, a vast amount of field data shows that microplastics are widespread in mussels around the world, and laboratory exposure studies have demonstrated that mussels can be good model organisms for understanding uptake, accumulation and toxicity of microplastic (Tables 1, 2, S1). This highlights the feasibility and advantages of mussels as indicator species for monitoring of microplastics from an implementation perspective.

Practically, the quantification of pollutant levels in bioaccumulator organisms and a specific response to a toxic substance by an organism provide two frequently employed pathways for

monitoring environmental quality (Reguera et al., 2018). The suitability of the first approach relies on the relationship of pollutant level between the organism and ambient environment. Based on laboratory studies, mussels show size selection for particle uptake (Ward and Shumway, 2004). Nevertheless, there are diverse ways for mussels to take microplastics (Kolandhasamy et al., 2018), and various microplastics exist in real environments. Though not all the properties of microplastics in mussles can exactly match those in their environment, quantitative correlations of abundance between microplastics in mussels and in surrounding seawaters makes it practicable to deduce environmental microplastic pollution levels from that in mussels (Qu et al., 2018). Since the concentration of pollutants including microplastics in mussels tend to remain stable after obtaining a balance between intake, assimilation in tissues and defecation/eggestion, this method can effectively mitigate or avoid error rates and misinterpretation stemming from contingency in environmental medium (Setälä et al., 2016; Beyer et al., 2017).

As for the other pathway, efforts have been taken to reveal the toxic effects resulting from microplastic intake, translocation and accumulation in mussels. Most biomarkers such as lysosomal membrane stability, inflammatory response, antioxidant enzymes are sensitive to other pollutants as well (Brooks et al., 2011; González-Fernández et al., 2016; Burgeot et al., 2017). Utilising these toxicological studies will provide evidence and scientific basis for the selection of specific biomarkers for the early warning and monitoring of microplastic pollution and related ecological risk assessment.

Recently, Fossi et al. (2018) proposed to use a threefold monitoring approach to assess the impact of ingested marine litter including microplastics on marine organisms. It combines an accurate measure of microplastic levels in target organisms, the concentrations of plastic additives and other persistent organic pollutants (POPs) in tissues and the corresponding toxicological effects. According to this new concept, mussels correspond to ideal biological models because they have been widely used as bioindicators of POPs in coastal environments (Liu et al., 2014; Martinović et al., 2016; Aznar-Alemany et al., 2017; Cunha et al., 2017; Gagné et al., 2017; Chiesa et al., 2018; Chiu et al., 2018; ; Politakis et al., 2018).

4.2. Current regional and national proposals

Recently, mussels have been proposed as suitable indicator organisms of microplastic pollution by research groups from several geographic locations (Van Cauwenberghe et al., 2015; Wesch et al., 2016; Li et al., 2016; Lusher et al., 2017b; Qu et al., 2018). Uptake and accumulation of microplastics in mussels from Belgium has been selected as a marine health status parameter, and microplastic levels in mussels have been included in European databases regarding contaminants of emerging concern in seafood (De Witte et al., 2014; Vandermeersch et al., 2015a). The possibilities of using mussels as monitoring species for microplastics in Norway and the Nordic marine environment is also supported (Bråte et al., 2017; Lusher et al., 2017b) since they have been used in other regional, national and international monitoring programmes. Lusher et al. (2017b) suggests that mussels (*Mytilus* spp.) can be a promising bioindicator of the smallest sized microplastic (<1 mm) in the water column.

In a recent workshop on "*Distribution, source, fate and impact of marine microplastics in Asia and the Pacific*" organized by the IOC Sub-Commission for the Western Pacific (WESTPAC), mussels were recommended as bioindicator species to monitor marine microplastic pollution (WESTPAC, 2017). At the European level, the MSFD has defined marine litter and microplastics as a full descriptor of the Good Environmental Status (Galgani et al., 2014). OSPAR have recommended mussels as suitable monitoring species because of their large stocks for repeated sampling and the ability to reflect the local conditions (OSPAR, 2012). Due to advantages of mussels as traditional biological indicators and mounting evidence of microplastics in mussels, ICES have advised to use mussel as a indicator of microplastic pollution (ICES, 2015; Vandermeersch et al., 2015b; Beyer et al., 2017). However, there are currently no standard monitoring procedures outlined by any of the regulatory bodies (inc. OSPAR, MSFD, NOAA, UNEP). These monitoring protocols should follow recommendations from international experts and are expected to be produced in the near future, as the GESAMP Working Group 40 is currently formulating a report to harmonise monitoring and assessment of plastics and microplastics globally.

4.3. Future developments

Based on the analysis above, we propose to use mussels as bioindicator species for monitoring microplastics in marine environments. Nevertheless, some questions require further clarification, and additional factors should be taken into consideration when it comes to building an efficient and economical approach suitable for future large-scale monitoring program with mussels.

Firstly, it is necessary to develop a global working group investigating microplastics in mussels under some international organization such as UNEP, including underlying physiological and behavioral processes and responses to microplastics. Already, mussels have been proposed to be used as bioindicators in some local or regional areas. It is time to form a working group globally so that researchers from different areas share and discuss the protocol of monitoring as well as future plans. One possible arena to advertise and promote this discussion is the Ad Hoc Open-Ended Expert Group on Marine Litter and Microplastics composed by representatives from member states to support the implementation of the United Nations Environmental Assembly resolution on marine litter and microplastics (UNEP/EA.3/L.20).

Secondly, a uniform protocol should be developed and adopted, at least on a comparable regional monitoring basis. Uniform protocols and harmonized monitoring methods are need to allow spatial and temporal comparisons and to enable assessment of the presence of microplastics and their effects in mussels at a global level (Fossi et al., 2018). Such a detailed methodology for measuring microplastics in blue mussels has also been described by Lusher et al. (2017b) which supplies a potential baseline standard to conform too. Future inter-calibration exercises will help validate and harmonize methods used across different research groups. The development and use of an internal reference sample(s), one for each matrices, might also help facilitate inter-laboratory and global validation of results.

Finally, monitoring should be practicely conducted regionally or globally. To date, comparable data of microplastic pollution characteristics in mussels from different parts of the world is scarce. Ideally, researchers should be encouraged to combine microplastic monitoring into the existing monitoring projects using mussels. A global picture of microplastic should be obtained, and the potential ecological and health risk should be assessed.

5. Conclusions

Current evidence on microplastic abundance in all parts of the marine environment including biota call for establishing a suitable indicator species for microplastic pollution, to monitor spatial and temporal trends internationally. Mussels have been widely used as bioindicators for monitoring of coastal water pollution and their susceptibility to microplastic uptake and assimilation has been well

documented. Field investigations have shown that microplastic abundance in mussels is closely related to human activity and, in some studies, there has been a positive and quantitative correlation of microplastics in mussels and their surrounding waters. Laboratory exposure studies demonstrate that mussels can be good model organisms when investigating uptake, accumulation and toxicity of microplastics. Therefore, we strongly propose the use of mussels as indicator species for monitoring of microplastics in the marine environment. We also urge the international organizations (e.g., UNEP) to facilitate the formation of an international workgroup of microplastics in mussels to develop an internationally accepted protocol to monitor and collect preliminary data comparing coastal mussels from around the world.

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Appendix A. Supplementary data

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