Paint particles in the marine environment: An overlooked component of microplastics

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1. Introduction

The recent scientific, policy and management literature contains a plethora of studies on microplastics in the environment, and in particular in the marine environment (Abbasi et al., 2018; Dauvergne, 2018; Liubartseva et al., 2018; Kane and Clare, 2019; Henderson and Green, 2020; Kor and Mehdinia, 2020). Microplastics have been operationally defined as synthetic or semisynthetic materials constructed of polymers and additives that are < 5 mm in diameter (Arthur et al., 2009), with nanoplastics defined with a 1000 nm upper size limit (Gigault et al., 2018). Microplastics may be further divided into primary particles, like pre-production pellets, clothing fibres and exfoliating beads used in cosmetic products that are < 5 mm before entering the environment, and secondary particles that are broken down from larger debris in situ to fragments, films, foams and fibres of < 5 mm in size. Microplastics may also be classified according to additional physical or chemical attributes, like polymer type, colour, roughness, transparency and shape (Gauci et al., 2019; Bikker et al., 2020).

Paint is a pigmented and usually opaque surface coating that has decorative, protective or other specific technical properties (OECD, 2009). Paint consists of polymers and additives and dried paint particles of < 5 mm in size derived from the deterioration or removal of surface applications (paint particles) should, strictly, be classified as microplastics according to the definition above. However, there is relatively
little systematic study of paint particles in the current marine scientific literature (Galafassi et al., 2019; Gaylarde et al., 2021). This may, in part, be attributed to the omission of paint in marine litter guidelines and the consequent inconsistencies regarding its classification in the microplastic literature (OSPAR Commission, 2010). Thus, some investigations include paint particles amongst the microplastic cohort (Lima et al., 2014; Cardoza et al., 2018; Haave et al., 2019) while others operationally, incidentally or deliberately exclude them or treat them differently when reporting or characterising samples (Free et al., 2014; Baini et al., 2018; Ferreira et al., 2019; Lacerda et al., 2019). In this paper, the similarities in and differences between microplastics and equivalently sized paint particles are critically evaluated based on their uses, physico-chemical characteristics, sources, behaviours and relative abundances in the environment. The focus is on the marine setting in which both types of solid clearly exist, but where the connection is often distinctly lacking or overlooked.

2. Plastics versus paints

There are fundamental similarities in and differences between plastics and paints which are conceptualised in Fig. 1. Thus, plastics are constructed of one or more synthetic polymer whose properties may be customised by functional additives and fillers, while paints consist of fine, natural or synthetic polymeric particles (the binder or resin) and additives and fillers that are held together on a surface as a “plastic-like” film when cured. Frequently used synthetic polymers in consumer and industrial plastics include acrylonitrile butadiene styrene, polyethylene, polystyrene and poly(vinyl chloride) (PVC), while most paints are based on acrylic, alkyd, polyurethane, epoxy or chlorinated rubber binders.

Some additives are common to both plastics and paints, like inorganic fillers and colourants, while others are specific to or more commonly employed in either plastics or paints because of differences in the manufacture, processing and function of the materials. For example, dryers, emulsifiers and adhesion promoters are critical to the storage and application of many kinds of paint whereas blowing agents, lubricants and impact modifiers are more important for the performance and durability of certain plastics. In general, the mass content of additives (including pigments) and fillers is significantly greater in dried paint films than in plastics and, consequently, the polymeric content is usually much greater in the latter. There is also a smaller range of additives available for plastics because the higher processing temperatures required constrain the selection of chemicals, and in particular colour pigments, based on thermal stability.

These differences generally mean that paint particles are denser, more brittle, more angular, less transparent and chemically more heterogeneous than microplastics. Moreover, the nature of paint application results in a layered structure (of identical or different formulations) that may be contaminated by residue from the underlying substrate. These characteristics may account for why paints are deliberately (through appearance under a microscope; Imhof et al. 2016; Horton et al., 2017) or operationally (via density separation or chemical treatment; Coppock et al., 2017; Rodrigues et al., 2020) excluded from the overall micro-debris pool in environmental samples. (Micro-debris is defined as all synthetic debris of < 5 mm in size, including fibres, tire-wear particles and other vehicle-derived solids, pellets, beads and paints.)

3. Sources of paint particles to the marine environment

The broad sources of paint particles to the marine environment are shown in Fig. 2, where paints are defined according to their principal users or applications; a selection of various, more specific sources is also illustrated in Fig. 3. In theory, and as with microplastics, sources can be divided into primary and secondary, with the former representing fine (< 5 mm) particles transported into the environment via runoff or the atmosphere and the latter encompassing particles that are formed in situ from, for example, the weathering of larger fragments or the deterioration or damage to coastal structures and boats. Ultimately, however, most paint particles are formed by the same mechanisms acting on the paint film and such a classification does not bear any relationship with physical or chemical properties or environmental behaviour. Mechanisms of deterioration are UV degradation of the binding polymer and intentional or unintentional mechanical disturbance of the coating (damage, wear and tear, maintenance or removal).

The poor management of municipal waste containing unused paint or painted structures (including building waste) and packaging may act as a direct source of paint particles to the coastal environment through littering and failing coastal landfills (Pope et al., 2011). More generally, however, the urban environment represents a rich and varied indirect source of paint particles that are generated when decorative, anticorrosive or safety paints on private and public buildings, road surfaces, and municipal structures and street furniture undergo natural deterioration or are deliberately disturbed during maintenance, repair or

Fig. 1. The chemical constituents of plastics and paints. Polymers, pigments, fillers and certain additives are common to both materials (with the arrow showing the direction of greater abundance) while other additives are more specific to either general or speciality plastics or paints.
removal (Jartun et al., 2009; Turner and Solman, 2016; Horton et al., 2017; Lee et al., 2018). More directly, large coastal structures like bridges and decommissioned naval establishments may act as locally significant sources of paint particles to the marine environment (Finckelstein et al., 2003; Shu et al., 2015). However, perhaps of broader concern in this respect is the uncontrolled release of paint particles from boats that are abandoned, being recycled or undergoing repair (Reddy et al., 2003; Singh and Turner, 2009; Rees et al., 2014; Soon et al., 2021). Regarding the latter, maintenance of hulls in the leisure boat sector is generally unregulated and yet produces significant quantities of toxic antifouling paints particles of a range of sizes from the scraping, sanding, stripping, sand-blasting and hydro-blasting of spent coatings. Furthermore, older formulations that are disturbed often contain metals and biocides that are now restricted or prohibited (Eklund et al., 2014; Soroldoni et al., 2018).

Paint particles from these sources are transported to the aquatic environment as airborne particulates that are subject to fallout, with road runoff or treated waste water, or via washdown from boat maintenance facilities or coastal structures undergoing repair. The significance of the local transport of airborne paint particles has been established from the levels and signatures of contamination of soils and roof top dusts of residential and municipal buildings in the vicinity of roads, dry docks, boatyards and harbours (Decelis and Vella, 2007; Jartun et al., 2009; Turner, 2013; Eklund et al., 2014; Sakata et al., 2017; Meza-Figueroa et al., 2018). Given the particle size range generated by sanding (in the range of < 50 nm to a few μm in diameter; Koponen et al., 2009), however, the potential range of airborne transport of paint particles is considerable. For example, microplastics of dimensions orders of magnitude greater (albeit less dense and usually fibrous) appear to have the propensity to be transported thousands of km with regional air masses (Bergmann et al., 2019; Brahney et al., 2020) while geosolids towards the upper size limit generated by sanding may be transported globally (Mahowald et al., 2014).

Paint particles are also generated while ships are at sea. For example, groundings and collisions are known to generate large quantities of antifouling paint around the location of impaction (Negri et al., 2002; Jones, 2007; van der Schyff et al., 2020), while vessels navigating through ice or fishing activities generating friction between painted surfaces and rope may also act as more diffuse sources (Negri and Marshall, 2009; Song et al., 2014). It is also likely that paint particles are generated more passively with the general wear and tear (erosion) of hull, waterline, topside and deck coatings (Dibbe et al., 2021).

4. Inputs of paint particles to the marine environment

Estimating the quantities of paint particles entering the marine environment or their contribution to the total marine microplastic pool is fraught with difficulties and uncertainties. One of the fundamental
considerations in this respect is whether the mass of dried paint should be converted to the mass of polymer by subtracting the weights of additives and fillers present before direct quantitative comparisons are made with microplastics. While this approach is often favoured for flux calculations and inventories, additives and fillers are integral components of both paints and microplastics that, from physical, environmental and toxicological perspectives, should not be ignored.

Input estimates are generally based on the quantities of paints manufactured for or sold in different sectors and quantitative assumptions about longevity, removal, disposal and retention by waste facilities, and are usually compared with estimates for other types of microplastic based on equivalent assumptions (Sundt et al., 2014; Lassen et al., 2015; Verschoor et al., 2016). In a recent example, Ham et al. (2018) estimated emissions of different types of microplastics to the European aquatic environment using data, information and assumptions on paints published in earlier OECD reports (OECD, 2005; OECD 2009) and supplied by the European Council of the Paint, Printing Ink, and Artist’s Colours Industry (CEPE). Neglecting inputs from poor waste management or the degradation of larger plastic waste, lower, middle, and upper estimates for each microplastic category considered were derived (Table 1). For building, marine and road marking paints, estimates were based on figures for consumption, coupled with assumptions on rates of removal during maintenance, wear and tear and polymer degradation, and rates of retention, entrapment or loss by water treatment facilities, road-side sedimentation devices, road cleaning, adjacent soils and asphalt surfaces. Not factored into the estimates for paint were emissions to water when uncur, disturbance of layers of paint below the top coat (for example, primers and base coats), weathering of marine paints while in service or at the end-of-life, and the direct input of airborne dusts. Within these assumptions and constraints, estimates for paint particle emissions to surface waters range from about 12,000 to 30,000 tonnes, or between about 10 and 17% of total microplastic inputs. Presumably, inputs to the marine environment are related to these estimates but with modification of building and road paint inputs by settlement and entrapment in rivers.

Given the many sources of marine paint particles neglected in the calculations above, it is likely that the emission figures for each paint category in Table 1 have been underestimated. In their calculations, for example, McAdams and Angelskår (2020) considered all paint applied to industrial and marine steel assets (not just marine paints, and about 42 billion litres per annum) and assumed a 20-year coating lifetime, or a 5% loss per year, and 50% efficiency of waste retrieval. A resulting 2-3 million tonnes of paint particles was predicted to enter the oceans annually, representing a highly significant fraction of the estimated 8 million tonnes of total plastic entering the marine environment each year (Jambeck et al., 2015). The calculated relative contribution of paint to the latter estimate is still subject to many uncertainties but is more consistent with measurements of the relative abundance of micro-debris constituents (that include paint) obtained by surface trawls in the ocean and as reported below.

### Table 1

| Paint Type                  | Upper  | Middle | Lower |
|----------------------------|--------|--------|-------|
| Automotive Tyres           | 136,000| 94,000 | 52,000|
| Pellets                    | 78,000 | 41,000 | 3,000 |
| Washing of Clothing        | 25,000 | 13,000 | 4,000 |
| Road Markings              | 21,000 | 15,000 | 10,000|
| Building Paint             | 8,000  | 5,000  | 2,000 |
| Fishing Gear               | 5,000  | 2,600  | 500   |
| Automotive Brakes          | 5,000  | 2,000  | 100   |
| Artificial Turf            | 3,000  | 2,000  | 300   |
| Marine Paint               | 400    | 400    | 400   |
| Total                      | 280,600| 176,300| 71,800|
| Total paint                | 29,400 | 20,400 | 12,400|

5. Presence and abundance of paint particles in the marine environment

The presence of protective and antifouling paint particles in sediment deposits in the vicinity of boatyards, marinas, harbours and abandoned boats has been well-documented from visual or microscopic inspection of sieved samples (Thomas et al., 2003; Turner, 2010; Takahashi et al., 2012; Eklund et al., 2014; Rees et al., 2014; Costa et al., 2016; Lagerström et al., 2016; Soroldoni et al., 2018; Abreu et al., 2020). Based on sediment contamination by Cu, Singh and Turner (2009) estimated antifouling paint particle abundance of up to 1% on a mass basis in a tidal inlet of southwest England, while direct counting of antifouling paint particles in sediments from a Brazilian lagoon by Soroldoni et al. (2018) revealed mass contamination of up to 4.4%.

In contrast, very little quantitative information exists on paint particles in intertidal and benthic marine sediments more remote from significant point sources. Thus, in the coastal environment, Diez-Minguito et al. (2020) refer to paint “sheets” in sediments from the Ria de Vigo, northwest Spain, at water depths of up to 40 m, while Haave et al. (2019) report paint particles in a Norwegian fjord at depths of up to 330 m. Along the strandline of a sandy beach in eastern England, Latuta et al. (2019) report that almost one half of microplastics retrieved by density separation were paint-based. In the open ocean, Fischer et al. (2015) mention paint chips being present in sediments from the northwest Pacific Kuril–Kamchatka Trench and its adjacent abyssal plain at depths of around 5 km. Unfortunately, however, none of these studies provide clear quantitative information on the paint particles observed, nor characteristics that could determine their origin.

A greater body of more quantitative data exist for paint particles captured by plankton trawls at or near the sea surface. In some cases, visible or chemical characterisation of particles has revealed that the protective or antifouling paints of the research vessel or trawl frame was a significant (and sometimes dominant) source created by shedding from the hull or collisions between the vessel and sampling equipment (Rummel et al., 2016; Bagaev et al., 2017; Eriksen et al., 2018; Lacerda et al., 2019; Suaria et al., 2020). However, in other cases, trawls deployed from unpainted vessels or paint sample characteristics sufficiently different to those of formulations applied to the research vessel or trawl frame have allowed external sources to be inferred. Lima et al. (2014) found that paint particles contributed nearly 30% of all microplastics and up to about 0.1 particle m$^{-3}$ in 300 μm nets trawled in the Goiana estuary, Brazil. The greatest concentrations were encountered in the bottom waters and during the rainy season, coincident with the period of most intensive fishing activities. Kang et al. (2015a) found alkyd-based paint particles contributed 20 to 50% of suspended microplastics in the mouth of Nakdong River, Korea, with abundances of up to about 1 m$^{-3}$ captured by a 330 μm Manta trawl but up to 230 m$^{-3}$ captured by a finer, 50 μm hand net. Dibke et al. (2012) recently estimated that up to 80% of the microplastics sampled from surface waters of the German Bight (North Sea) using an on-board deck-wash system were ship paints based on epoxy, acrylic and chlorinated rubber binders; the remaining material was dominated by polyethylene, polypropylene and PET derived from packaging and mismanaged waste. Paint particles have also been reported amongst microplastic debris retrieved from plankton trawls in the North Atlantic (Moret-Ferguson et al., 2010), Black Sea (Oztekin and Bat, 2017), tropical and equatorial western Atlantic Ocean (Ivar do Sul et al., 2014; Garcia et al., 2020), Adriatic Sea (Suaria et al., 2018), Gulf of Oman (Aliabadi et al., 2019), and over the Great Barrier Reef (Hall et al., 2015; Jensen et al., 2019). In the coastal zone of Korea, Song et al. (2014) observed paint particles in the sea surface microlayer, or the boundary layer between the atmosphere and ocean, sampled to a depth of 400 μm using a metal sieve. The mean particle abundance was found to be about 200 L$^{-1}$ or 150 m$^{-2}$ compared with an abundance of other microplastics of 16 L$^{-1}$ or 13 m$^{-2}$. Significantly, there was a distinct increase in paint particle number with decreasing size (< 1000 μm to < 50 μm) which was
attributed to the gradual breakdown of the floating stock of paint particles into smaller pieces.

Quantitatively and qualitatively, the observations and measurements above are subject to various limitations and uncertainties. Specifically, in sediment, where harsh chemical treatment (e.g., peroxodization or acidification) and flotation are employed to isolate microplastics, the abundance of paint particles is predicted to be under-reported because of their greater density and lower chemical stability (see below). In seawater, paint particle concentrations are likely underestimated because of the inherent size limits of the capturing device or detection methods and the propensity for paint particles to fragment to sizes below these limits during sample processing. Nevertheless, it is clear that paint particles are a significant, if not the dominant form of microplastic at or near the water column surface and within intertidal and benthic sediments in many marine settings. Accordingly, it would appear that the sources, processes and assumptions involved in estimates of paint inputs into the aquatic environment (and as exemplified in Table 1) require re-evaluation.

6. Environmental behaviour, transport and exposure of paint particles

One of the key physical factors that determines the transport, behaviour and fate of plastics in natural waters or engineered systems (for example, water treatment facilities) is density (de la Fuente et al., 2021). This property also forms the basis by which microplastics are extracted (through flotation) from environmental samples like soils and sediments (Miller et al., 2017). Density ranges for common plastics and paints are compared in Fig. 4. For plastics, the lowest values represent the pure polymer while maximum values denote formulations loaded with the highest concentrations of mineral or glass fibre filler, plasticiser or flame retardant according to the Omnexus material selection platform (SpecialChem, 2021). For dried paints, the overall density range and noted formulations are based on information in the scientific and commercial literature. Also annotated on Fig. 4 are the density range of seawater and the density range of solutions of ZnCl₂ that are commonly used to extract microplastics from environmental samples (Rodrigues et al., 2020).

Low density plastics like polyethylene are predicted to float in seawater while high density plastics like PET and PVC are predicted to sink. Plastics of intermediate density or with a relatively low density but broad range, like polypropylene and polystyrene, are expected to exhibit negative or positive buoyancy depending on the content and nature of additives and fillers, the precise density of ambient seawater and any biofouling present. For paints, however, all formulations are, ultimately, predicted to sink in the marine environment based on density considerations because of the greater abundance of inorganic additives and fillers in the dried formulations. This means that, in theory, paint particles of equivalent dimensions and shapes to microplastics are more readily deposited and less mobile in aquatic systems. Regarding extraction by flotation in ZnCl₂ solution, the majority of microplastics should be isolated when a relatively high concentration of the salt is employed, but many types of paint particles, including those with clear marine origins, would evade capture by this approach (Haave et al., 2019). That is, conventional extraction methods are predicted to underestimate the stock of micro-sized, synthetic polymeric particulates.

Empirical studies on the settling characteristics of antifouling paints in artificial estuarine water (salinity = 15) have shown that, above a diameter of 1 mm, settling is related to density and size, but below 180 μm most particles remain at the surface without agitation and settle only after the surface tension had been broken with stirring (Soroldini et al., 2018). Accordingly, the authors suggested that the size range of paint particles generated in local harbours and boatyards could be transported several tens of km from their point of origin in a lagoonal estuary within 24 h, depending on local hydrodynamic and meteorological conditions. More generally, surface tension, augmented in natural seawater by biogenic material (Song et al., 2014) and broken only by energy equivalent to wind speeds > 6.6 m s⁻¹ (Wurl et al., 2011), coupled with long-range airborne transport, may account for the ready detection of paint particles at or near the sea surface in locations remote from any immediate point sources and as discussed earlier.

With paint particles encountered at the sea surface and in the pelagic and benthic zones it is not surprising that they have been detected among other micro-debris in the digestive tracts (and sometimes gills) of various marine animals, including birds, fish, crustaceans, cetaceans, turtles and invertebrates, and as summarised in Table 2. In many cases, paint particles are noted without detailed or systematic quantification or characterisation and critical information on abundance, sources, selectivity, impacts and fate is lacking. Where paint particles have been counted and classified in fish guts, however, the contribution of this material to the MP cohort on a number basis may be as high as 35% (Cardoza et al., 2018), with blue often noted as the dominant colour (Herrera et al., 2019). Paint particles may be ingested passively while suspended in the water or attached to dietary material like algae.
tributyl tin, lead, arsenic and mercury phased out but still encountered many decades, with other metal-based biocides based on compounds of (I)-based compounds have played a critical role in antifouling paints for retardants have been employed in speciality paints, of more general sectors but legacy coatings remain an important source of the metal to the environment (Mielke et al., 2008; Turner and Lewis, 2018). A greater concern regarding contemporary and historical paint particles, however, is the presence and availability of additives that are hazardous. While asbestos, cadmium compounds and certain phthalate esters and brominated flame retardants have been employed in specialty paints, of more general concern are lead-based compounds that were common in historical paints with wide-ranging applications (including road markings, buildings and shipping) and biocidal additives that are fundamental to marine antifouling formulations.

A variety of lead compounds were used as drying catalysts (lead acetate, octoate and napthenate), pigments for colour, opacity and protection (e.g. lead carbonates, sulphates, oxides, chromates), and, on metal, corrosion inhibitors (e.g. lead tetroxide, calcium plumbate). Restrictions have eliminated the intentional use of lead in paints many sectors but legacy coatings remain an important source of the metal to the environment (Turner et al., 2018). Copper (I)-based compounds have played a critical role in antifouling paints for many decades, with other metal-based biocides based on compounds of tributyl tin, lead, arsenic and mercury phased out but still encountered on the hulls of older boats and abandoned vessels (Eklund and Eklund, 2014; Rees et al., 2014). Various organic- and organo-metallic “booster” compounds with herbicidal properties have often been added at smaller concentrations to antifouling paints, with many undergoing restrictions but encountered in historical boat coatings (Parks et al., 2018).

While lead compounds and certain biocides have also been added to plastics, both their concentrations and mobilities are significantly greater in paint particles. For example, the maximum lead content of beached microplastics (mainly polyolefins) retrieved from south west England was about 5000 mg kg$^{-1}$ (Massos and Turner, 2017), with PVC fragments having concentrations in the range 10,000 to 20,000 mg kg$^{-1}$ (Turner and Solman, 2016). In contrast, the lead content of paint fragments derived from the hulls of abandoned boats can regularly exceed 20% by weight (Rees et al., 2014), or an order of magnitude higher than the maximum concentration reported for PVC. Regarding biocides, the typical content in protected plastics is $< 1,000$ mg kg$^{-1}$ (Dylingowski and, 2004). This contrasts with Cu(I) concentrations in the dry films of contemporary and historical antifouling formulations of up to 50% by weight (Muller-Karanassos et al., 2019).

A specific chemical comparison between a fragment of polyethylene pigmented with PbCrO$_4$ that had been retrieved from the strandline of a beach and a copper-based antifouling paint fragment taken from an abandoned boat is shown in Fig. 5. Here, X-ray fluorescent spectra acquired under the same operating conditions are illustrated over the energy range 0 to 14 keV, with selected principal secondary X-ray peaks identified. Annotated in the boxes, and where quantification was possible by fundamental parameters, are concentrations of elements detected in each sample in mg kg$^{-1}$. Clearly, the number and magnitude of peaks and overall elemental concentrations are greater in the antifouling fragment but the percentage balance (also annotated), consisting of light elements not detected by the instrument ($Z < 11$) and indicative of the polymer concentration, is considerably lower. More generally, these observations are also consistent with the greater abundance of additives and lower polymer content of paints relative to plastics discussed earlier.

The greater mobility of hazardous chemicals in paint particles relative to plastics is evident from differences in the quantities of toxic metal ions (e.g. Cu$^2+$, Pb$^{2+}$, Cr$_2$O$_7^{2-}$) released from each particle type into physiological solutions that simulate the mammalian or avian stomach (Turner and Radford, 2016; Smith and Turner, 2020) and from the more stringent guidelines and regulations that deal with toxic metals in paint (Gooch, 1993). Factors that account for a greater mobility of metals in paint than in plastic include a higher additive to polymer/binder ratio in the dried formulation, the brittleness of weathered paint and its propensity to readily fragment into smaller particles of high surface area, and the more ready aging and degradation of the paint binder than the plastic polymer. More specifically, in hard, ablative and self-polishing antifouling paints, biocides are designed to leach out or the entire formulations dissolve at controlled rates into the surrounding medium.

The greater solubility of hazardous additives in paint than in plastic also results in a higher toxicity. Thus, in the aquatic environment, while chemicals leaching from microplastics, including Pb, have been shown to be toxic to bacteria, invertebrates and fish (Silva et al., 2016; Boyle et al., 2020; Sarkur et al., 2020), the quantities of plastic relative to

### Table 2

A compilation of reports of paint particle ingestion by marine animals. Asterisks denote reports of particles in the gills as well as the digestive tract; ns = not stated.

| Animal | Location | Size, mm | Paint type | Reference |
|--------|----------|----------|------------|-----------|
| Layson albatross, Diomedea immutabilis | Midway Atoll, North Pacific | ns | building paint | Sileo and Fefer (1987) |
| Cape petrel, Daption capense | Equatorial Pacific | ns | ns | Laist (1997) |
| Green sea turtle, Chelonia mydas | Central Pacific | ns | ns | Russell et al. (2011) |
| Various pelagic and benthic fish | Portuguese coast | < 4.8 | alkyd | Neves et al. (2015) |
| Amberstrip scad, Decaptus muroadi | Southern Pacific subtropical gyre | 1-2.5 | acrylic | Ory et al. (2018) |
| Green sea turtles, Chelonia mydas | Great Barrier Reef | < 5 | acrylic | Markic et al. (2018) |
| Various pelagic and benthic fish | Southern Pacific subtropical gyre | < 5 | acrylic | Cardozo et al. (2018) |
| Mauve stinger, Pelagia noctiluca | Tyrrenhian Sea | 3 | zinc-rich | Macalik et al. (2018) |
| Atlantic bigeye, Prionacanthus arenatus | Brazilian coast | < 10 | ns | Cardozo et al. (2018) |
| Various eels | Irish coast | ns | ns | Lusher et al. (2018) |
| Pelagic fish* | Musa estuary and Persian Gulf | 0.2 | anti fouling | Abhasi et al. (2018) |
| Ray worm, Heliate diversicolor | Plymouth Sound, SW England | 0.26 | anti fouling | Muller-Karanassos et al. (2019) |
| Atlantic chub mackerel, Scomber colias | Canary Island coast | < 5 | ns | Herrera et al. (2019) |
| Various estuarine snoeks | Goiana estuary, Brazil | < 5 | ns | Ferreira et al. (2019) |
| Gurneet fish, Genidens geniden | Laguna estuarie system, Brazil | < 5 | ns | Dantas et al. (2019) |
| Benthic jellyfish, Cassiopea xamachana | Florida estuaries | ns | ns | Duff et al. (2020) |
| Mangrove crabs* | Hong Kong | > 0.01 | ns | Not et al. (2020) |
| Harbour porpoise, Phocoena phocoena | German coast | 0.1 | acrylic/alkyd | Philipp et al. (2021) |
water volume used to generate leachates are well in excess of levels encountered in the environment. By contrast, poisoning of birds through the ingestion of leaded paint particles has been documented in situ (Molnar, 1983; Finkelstein et al., 2003) and concentrations of anti-fouling paint representative of those encountered in contaminated harbour sediments have been shown to elicit toxic responses in epibenthic copepods, crustaceans and macroinvertebrates in controlled laboratory exposures. For example, Muller-Karanassos et al. (2021) report 5-day lethal and effects concentrations (LC50 and EC50, respectively) for size-fractionated particles (100 μm to 1 mm) derived from a
modern Cu-based antifouling paint of 19.9 and 14.6 g per L of estuarine sediment, respectively, for the ragworm, *Hediste diversicolor*, and 2.3 g L$^{-1}$ and 1.4 g L$^{-1}$, respectively, for the common cockle, *Cerastoderma edule*. Soroldoni et al. (2017) report a significant decrease in fecundity for epibenthic copepods exposed to 0.01% of modern, Cu-based antifouling paint particles (< 63 μm) in estuarine sediment and an LC$_{50}$ arising from the elutriate of a preparation equivalent to 0.14% of paint. Soroldoni et al. (2020) subsequently demonstrated 10-day LC$_{50}$ values for benthic microcrustaceans exposed to estuarine sediment spiked with fractionated antifouling paint particles of 0.16 to 0.45% by mass of sediment.

The biocidal properties of antifouling formulations also impact on colonisation by marine bacteria. Thus, while microplastics are rapidly colonised by microbial communities that do not differ greatly from communities on other inert surfaces like glass, rock or wood (Wright et al., 2020), antifouling surfaces provide a habitat to select particular, but sometimes diverse bacterial populations that are resistant to the active biocides (Chen et al., 2013; Flach et al., 2017). Tagg et al. (2019) analysed and compared biofilm communities on microplastics and paint particles retrieved from sediment grabs in the coastal Baltic. Alkyd- and epoxy-based paints that likely included antifouling fragments (although this was not explicitly stated) were found to support communities that were distinct from and more consistent than biofilms on polypropylene and polyvinyl chloride microplastics and on natural particles. Significantly, an abundance of taxa from the Desulfobacteraceae family on some paint particles suggests that their presence in sediment may have impacts on the sulphur metabolism cycle.

8. Concluding remarks

The sources, pathways, behaviour and receptors of marine paint particles described above are summarised in Fig. 6. Also shown are the ions of greatest ecotoxicological concern that are, by design or otherwise, mobilised from paint additives into seawater. Given that both the abundance and mobility of harmful additives are greater in most paint particles compared with microplastics of equivalent dimensions it is concerning that the former are often overlooked or deliberately excluded from the micro-debris pool in environmental samples (Baini et al., 2018; Ferreira et al., 2019). This omission means that quantitative comparisons of plastic to paint abundance in seawater, sediments and digestive tracts are often lacking. It is also likely that the significance of the paint pool is understated from surface or near-surface water sampling because the most mobile paint particles evade capture by the mesh size of common trawling nets (100 to a few hundred μm) while larger particles have a greater propensity to sink. In sediment, paint particle abundance is also underestimated because the densities of conventional solutions used to isolate micro-debris are lower than the densities of many common paint formulations and treatments typically used to isolate microplastics may be physically or chemically destructive to paints.

Clearly, a greater awareness of the potential environmental and health impacts of paints particles and development of methods for their recovery and isolation are required amongst the microplastic community, and general measures to reduce the inputs of paints to the aquatic environment by manufacturers and users are called for. Regarding the latter, potential solutions include reducing the volume of materials that require painting, improving the wear resistance and enhancing end-of-life degradation of formulations, increasing the use and efficacy of dust extraction systems during maintenance and repair, improving waste collection systems in boatyards, and incentivising the safe and sustainable disposal or recycling of boats (Turner, 2010; Rees et al., 2014; Verschoor et al., 2016).

Declaration of Competing Interest

The author declares that he has no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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