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# The dark matter in water, air and land: from microplastic to invisible nanoplastics

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Microplastic is well known and has been the subject of many review articles. In recent years, an increasing number of reports have documented the presence of nanoplastics—plastic particles smaller than 1  $\mu\text{m}$ —in various environments, from the ocean to the human brain. In this article, I focus on nanoparticles and what we do and do not understand about their effects on our health. After an introduction to nanoplastics and their size relative to a single polymer chain, the degradation process that produces nanoplastics, similar to microplastics, is briefly summarized. Due to their high surface area, nanoplastics can behave differently in solution because they tend to aggregate. After reviewing the presence of microplastics and nanoplastics in humans, insights from the established field of nanomedicine are used to explore how nanoplastics may enter the bloodstream and reach the brain. This also includes the topic of protein corona formation, which influences the fate of nanoplastics in the body. Finally, a brief summary on the impact of plastic particles on health is provided, focusing on reports comparing nano- and microplastics. This article concludes with how materials scientists and chemists can contribute to addressing the rising plastic pollution problem.

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## New concepts

Everyone has heard of microplastics, but the term nanoplastic is only now entering the public consciousness. Recent articles have identified trigger points, showing that we have large amounts of nanoplastics in our oceans—an amount that far exceeds microplastics. Furthermore, plastic has been found in the brain, and closer inspection revealed it to be nanoplastic. We do not yet know how dangerous plastic truly is to our health, but initial reports, often using engineered micro- or nanoplastic beads, suggest it could be harmful. This feature article explores the pathways from plastic waste to nanoplastics, concluding with an examination of our current understanding of the differences between micro- and nanoplastics and their potential health implications. Although research on nanoplastic and its health effects is still limited, the mature field of nanomedicine can help us understand how nanoplastics in the environment may absorb small compounds like pollutants, form a biocorona with, for example, proteins, and transfer from the stomach, after ingestion with food and water, to the brain.

## Introduction

There is an abundance of microplastics in the ocean and in soil, mainly based on the polymers shown in Fig. 1. Pictures of coloured plastic pieces that are around the size of a grain of sand circulate widely on the internet. A recent UNESCO report paints a very dire picture of the state of our oceans. Around 8 to 10 million plastic ends up in the ocean and, alarmingly, it is estimated that by 2050, more plastic will be found in the ocean than fish.<sup>1</sup> Numerous local studies worldwide focused on quantifying the issue. The reality is that it is not as simple as collecting and counting the plastic bottles floating on water. Exposure to the elements results in the degradation of plastic

into small pieces. Some of them are visible with the naked eye, others require a light microscope. In 2004, Thompson and co-workers were the first to report on the occurrence of microplastics.<sup>2</sup> Since then, the problem is omnipresent with man-made plastics being discovered in land, air, animals, drinking water and more.<sup>3</sup> The term microplastic refers to a vast and diverse group of particles formed from the breakdown of plastics, classified by their size, which ranges from 1  $\mu\text{m}$  to 5 mm. The variety of chemical structures is huge. The most commonly produced plastics are summarised in Fig. 1, but there are many more. Just behind the term of Nylon are hidden several structural varieties that have all different properties. Even if we pick up two microplastic pieces of nylon from two different origins, they might vary in their molecular weight and in other aspects. The reality is that the plastic we use is a mixture of different ingredients. Polymers, which are the

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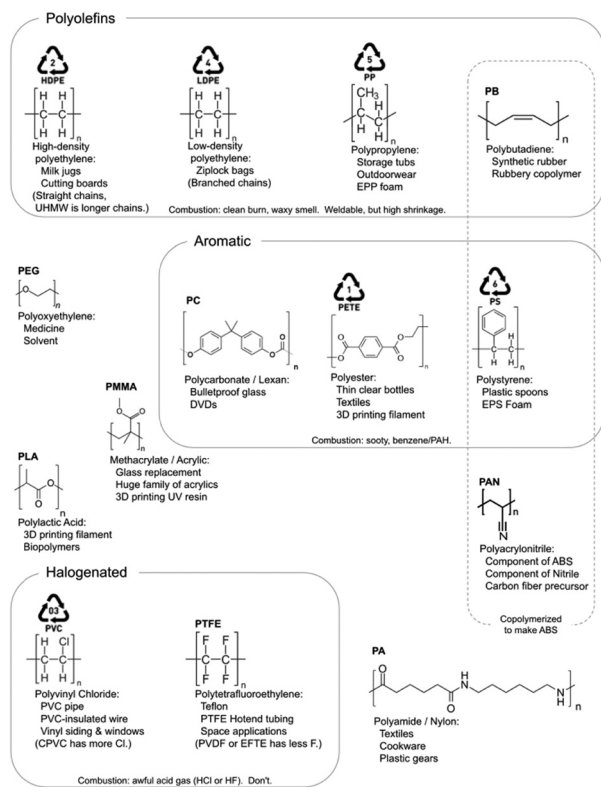


Fig. 1 Main commercial plastics. Source: Wikipedia, [https://upload.wikimedia.org/wikipedia/commons/5/5f/Plastics\\_Summary.svg](https://upload.wikimedia.org/wikipedia/commons/5/5f/Plastics_Summary.svg), CC0 1.0 DEED, CC0 1.0 Universal, Author: Orion Lawlor.

underpinning macromolecules, are processed in plastics by blending them with other polymers and with additives. These additives, which are small molecules, can serve different purposes, including enhancing the longevity of the plastic, providing colour, enhancing flexibility or rendering them with other properties.<sup>4</sup> These additives are not covalently attached to the polymer and can leach out during degradation, causing additional environmental damage.<sup>4</sup>

While there is a lot of focus on microplastic pollution, with many active community groups collecting samples and assessing the problems worldwide, recent reports have shown that nanoplastics are just as abundant. A 2023 study, published in *Nature* in 2025, highlighted that the main plastic pollution is nanoplastics, not microplastics, with around  $1.5\text{--}32.0\text{ mg m}^{-3}$  of PET, PS and PVC being found in various ocean layers. The authors estimated that the amount of nanoplastic found in the temperate to subtropical North Atlantic is around 27 million tonnes, which is exceeding or similar to the amount of large-scale plastic found in all oceans.<sup>5</sup> The authors concluded that “nanoplastics comprise the dominant fraction of marine plastic pollution”.<sup>5</sup>

Nanoplastics are defined by a size between 1–1000 nm, but I would advise against searching for nanoplastics of 1 nm. A monomer, a standard building block, typically measures about 500 pm to 1 nm. In commercial applications, many polymers consist of 500 to 1000 repeating units, but some can be significantly larger, such as Ultra-high-molecular-

weight polyethylene (UHMWPE), which can have up to 100 000 repeating units. The contour length of a PS with 1000 units is around 24 nm, but polymers do not exist as extended chains but as coils. An expression of “size” can be the random-flight end-to-end distance that increases in size with  $\sqrt{\text{Molecular weight } M}$  and equates for a polymer with 1000 repeating units to 15.4 nm.<sup>6</sup> This is only a mathematical model; in reality, environmental factors play a role. However, it should serve as an indication of how large a single polymer chain can be. Once we enter the realm of nanoplastics, other physical properties come into play. A plastic particle with a diameter of 1 mm has the same mass as  $10^{12}$  nanoparticles, each with a diameter of 100 nm, assuming they are made from the same material. At the same time, we created a 10 000 times increase in surface area, which means that now surface properties play an important role.

If we were to engineer 100 nm nanoparticles in the lab using any hydrophobic polymers such as PS, PP, PE, or others, we would observe that these nanoparticles quickly aggregate to form large visible particles in water as a means to minimise surface energy.<sup>7</sup> However, alterations to the surface properties through chemical reactions or adsorption of compounds found in the environment, facilitated by various interactions such as H-bonding, van der Waals forces or others, can create nanoparticles with colloidal stability.<sup>8</sup> Now, we have nanoparticles dissolved in aqueous solution that are invisible to the naked eye as long as the size remains below a critical scattering threshold. These nanoplastic particles sit alongside other nanoparticles found in nature.<sup>9</sup> These naturally occurring nanoparticles can be living, such as viruses, but also include non-biological ones like nano-sized soot particles, calcium ion particles in drinking water, or inorganic nanoparticles produced by bacteria.<sup>9,10</sup> Naturally occurring nanoparticles as well as man-made nanoplastic particles would behave very differently in a biological system compared to microplastics, as they can invade living organisms to an extent that microplastics cannot.<sup>9</sup>

In this focus article, the challenges related to nanoplastics will be examined. While the formation of microplastics is widely addressed, awareness of nanoplastics is only now beginning to grow. Clarivate Web of Science lists 13 766 articles mentioning microplastics, whereas only 1409 documents focus on nanoplastics while writing this article. Until 2020, there were just a few reports on nanoplastics, but the field has since expanded rapidly. An alarming report from 2025 showed the presence of plastic in the brain. The particles found were not microplastics, but the authors found particles below 200 nm in size.<sup>11</sup> The question is how nanoplastics are formed and how they can pass through the relatively impermeable blood–brain barrier (BBB). Researchers apply knowledge from nanomedicine, a field that started in the 1990s, to better understand the effects of nanoplastics on human health. However, several challenges still need to be addressed. While the impact of nanoplastics on our health is particularly interesting for medical researchers, materials scientists also play a role in helping detect nanoplastics or in developing polymer materials that are less likely to contribute to future pollution.



# How do nanoplastics end up in the environment?

Humans do not litter microplastics or nanoplastics; they litter large plastic items. In some countries, more than 2 kg of plastic is emitted into the ocean per person.<sup>12</sup> This amounts to around 100 plastic bottles, assuming a weight of about 20 g per bottle. We are all familiar with pictures of rivers that have floating layers of plastic waste. At this point, the plastic waste can be scooped up and processed safely. However, once nature exerts its full force, plastics start breaking down. We have all seen the degradation process looking at the plastic products around the house: the ones kept inside the house can last many years, but a plastic garden chair starts to yellow quickly, crumbles, and then breaks,<sup>13</sup> often without much prior warning that the end of structural integrity has been reached. Degradation of plastics can be accelerated by light (photodegradation and photo-oxidation), heat (thermal degradation), chemicals (oxidation, hydrolysis), microorganisms (biodegradation), mechanical force (physical degradation) and even electrical fields (Fig. 2). Several review articles describe plastic degradation in detail; only the main aspects are highlighted here.<sup>13–17</sup>

## Degradation by light and oxidation

Plastics in the environment are subject to light exposure that can trigger the degradation or ageing of polymers,<sup>18</sup> either in the presence or absence of oxygen.<sup>19,20</sup> UV irradiation at approximately 300 nm can directly break C–C and C–H bonds. However, the presence of chromophores, like carbonyl groups in the polymer, can accelerate this process. Certain compounds, such as metal compounds, pollutants, or oxygen, can

have catalytic effects and serve as photosensitizers.<sup>21</sup> The process catalyzed by oxygen is called photooxidation and can be divided into two mechanisms: the formation of singlet oxygen, which can lead to the formation of peroxides along the polymer chain, or the formation of a radical on the polymer that can react with oxygen, serving as an initiator for subsequent propagation across different parts of the polymer chains.<sup>19</sup> The oxygen-centred radical that is formed in the process allows polymer chain scission (Fig. 2). The rate of degradation is dependent on the intensity of light, the type of polymer, exposure time, and even the inclination angle.<sup>22–24</sup> UV-light can result in the formation of microplastics, but also nanoplastics, albeit this process is more pronounced when in combination with some mechanical agitation. UV light alone can enable the degradation into nanoparticles,<sup>25</sup> but simulated beach conditions using a standardized UV aging experiment revealed that after the initial formation of microplastics, nanoplastics less than 50 nm in size start to emerge.<sup>26</sup>

## Degradation by heat

Polymers are generally stable at moderate temperatures. They can transition from glassy to rubbery states if the glass transition temperature is reached at higher temperatures, but they do not visibly degrade until temperatures approach 200 °C or higher.<sup>27</sup> At this point, polymers degrade in a radical process encompassing initiation, propagation, and termination similar to that of UV degradation.<sup>28</sup> However, decomposition can already occur at lower temperatures, such as 30 °C, resulting in the slow release of oligomeric compounds.<sup>29</sup> It is therefore not surprising that microplastics are present when plastic is heated, such as with plastic teabags in hot water.<sup>30</sup> The Nylon and PET components of the tea bag started breaking down after only 5 min in hot water, releasing micron and nano-sized particles that affected the swimming behaviour of *Daphnia magna*, a planktonic crustacean.<sup>30</sup> Similar results were observed when hot solutions were filled into PP infant feeding bottles,<sup>31</sup> or when silicone-rubber baby teats are disinfected with steam,<sup>32</sup> both processes led to the formation of nanoplastics. Plastic containers together with their content are often placed in microwave ovens, which can locally generate very high heat. PP containers were found to release 4.22 million microplastic and 2.11 billion nanoplastic particles from only one square centimetre of plastic area within 3 min of microwave heating.<sup>33</sup>

## Degradation by mechanical abrasion

Light and heat can compromise the chemical structure through chain scission. However, it is often the movement of plastic, which includes abrasion from objects, sheer stress by tearing, or tensile stress by stretching, that causes the breakage of plastic into smaller pieces.<sup>34</sup> Under mechanical stress, polymers might initially respond with realignment of the polymer chains, but with increasing force, they can break, especially along entanglements or knots.<sup>35</sup> Most plastic is exposed to a variety of such external forces during usage, depending on the application. An example for mechanical force is washing

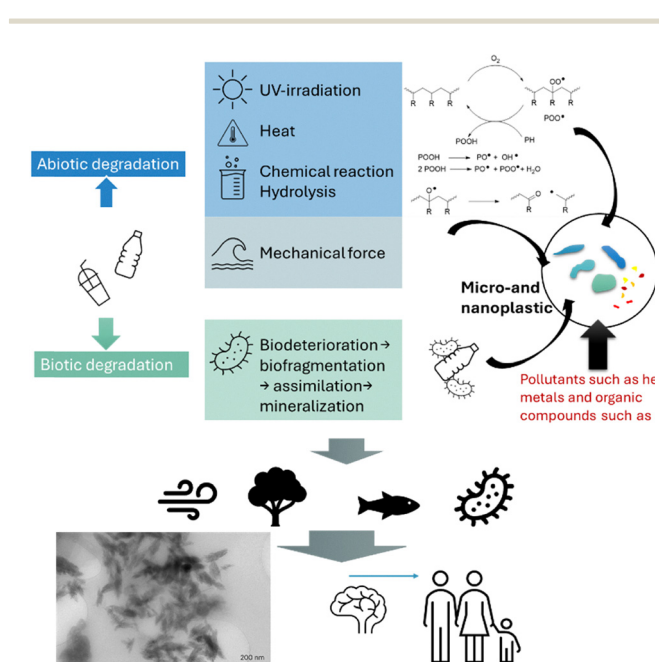


Fig. 2 Pathways of plastic degradation. The inserted TEM shows the nanoplastic pieces found in the brain by Nihart *et al.*,<sup>11</sup> reprinted from Springer Nature Medicine <https://creativecommons.org/licenses/by-nc-nd/4.0/>.

clothes. Each household produces microplastics simply by washing clothes.<sup>36</sup> A study showed that 124 to 308 mg microfibres per kg of washed polyester were produced, resulting in a set of recommendations on how to reduce microplastic pollution, such as the use of cold water or the installation of filters.<sup>37,38</sup> There is little discussion about the formation of nanoplastics, but when it comes to clothing, it is important to note that many clothing items are treated with engineered nanoparticles that might detach during mechanical stress. This can be mistaken for the formation of nanofibers generated during stress on the fabric. Other sources of microplastic in the household, that are subject to mechanical stress, include plastic chopping boards,<sup>39</sup> and a variety of food packaging, as reviewed earlier.<sup>40</sup> Microplastics have recently been found in unexpected places, like used chewing gum, which is polyvinyl acetate that breaks down through repeated mechanical stress.<sup>41</sup> A significant contributor to the microplastic pollution are tires, including bike tires, with a study finding that a mountain bike loses 3.62 g of plastic on a 100 km ride.<sup>42</sup> Even indoor climbing venues are not safe, as the constant mechanical stress on climbing shoes leads to the emission of large amounts of microplastic nanoparticles.<sup>43</sup>

While most studies have focused on microplastics, it is possible that nanoplastics might also be present. A study that accelerated the mechanical stress effect by blending polystyrene coffee cup lids immersed in water found that plastic can break down into pieces ranging from 100 to 200 nm within just 5 minutes. Microscopy analysis revealed the presence of a range of different shapes, including spindle-like structures.<sup>44</sup> This was also observed during simulations of water shear forces, where nanoplastics of approximately 50 nm in size were rapidly produced from microplastics.<sup>45</sup>

### Chemical reaction

Plastics can degrade by non-radical reactions.<sup>18</sup> This includes the reaction with ozone on double bonds, such as SEBS plastics, which is widely used in the automotive industry, for example.<sup>46</sup> Hydrolysable polymers with esters or amides in the backbone can theoretically be cleaved in an aqueous environment, but higher temperatures or pH values that are not neutral are required to accelerate the reaction.<sup>46</sup> At neutral conditions in a marine environment, it takes around 4.5 years for 50% of PET bonds to cleave when the ocean temperature is 35 °C, but 162 years if it is only 30 °C.<sup>47</sup> Earlier studies only focused on the formation of microplastics in the process of degradation,<sup>48</sup> but there is now an increased awareness that nanoplastic will also be generated.<sup>49</sup>

### Biodegradation

While polymers exhibit stability in pure aqueous environments, their degradation within biological systems is often accelerated by the synergistic effects of mechanical forces, ultraviolet (UV) radiation-induced photodegradation, enzymatic activity, and microbial colonization. The weathering of polymeric materials initially results in fragmentation into smaller particles, thereby increasing the surface area available for subsequent chemical

degradation reactions catalyzed by various environmental factors.<sup>50</sup> Many polyester polymers, such as PLA or PET, can be readily hydrolysed with the help of lipases, esterases, and others, but also polymers such as PVC and PP show signs of degradation when exposed to microorganisms such as fungi.<sup>51</sup> There is growing interest in developing genetically modified bacteria to help break down commodity polymers produced by radical polymerization, which are regarded as more difficult to degrade.<sup>52</sup> Biodegradation can be unpredictable, and labels in plastic items such as biodegradable or compostable do not mean that the plastic bag is completely depleted, as this will depend on subtle environmental differences. Placing in soil, hot compost, sea water, or simply in open air will lead to widely different degradation rates, to the point of showing little biodegradation.<sup>17,53,54</sup>

Biodegradation can also occur when plastics are ingested. Microplastics are internalised *via* drinking water, seafood, salt, honey, and many other food items.<sup>55</sup> But not all plastic can be broken down in the stomach. Large pieces of plastic often pass through the gastrointestinal (GI) system, which includes the stomach, colon, and intestines, unchanged. This is evidenced by the many LEGO pieces swallowed by children that only reappear after a few days, as shown by a study in which six paediatricians each swallowed a piece of LEGO.<sup>56</sup> Of course, large amounts of plastic can block the stomach or physically damage it and therefore result in the death of many smaller animals.<sup>57</sup> For smaller plastic pieces, the residence time in the stomach is often not long enough, and despite the high acidity in the stomach and the presence of bacteria and enzymes, medium-sized plastic pieces do not provide enough surface area to trigger substantial changes. However, there is evidence that smaller plastic pieces with higher surface area might indeed degrade in the intestines, subject to the nature of the polymer used. Understandably, esters such as PLA or PET might show quick signs of aging, but even polymers such as PS that are considered hard to degrade start breaking down, as they were observed to undergo oxidation on the surface when exposed to conditions found in the stomach.<sup>58</sup> While the low pH value in the stomach can facilitate hydrolysis, gut bacteria can assist in the degradation. Plastic-degrading bacteria were recently found in human faeces that could oxidize the surface of LDPE and PP and partly depolymerize the polymer chain.<sup>59</sup>

## Degradation to nano-sized particles and their analysis

### Effect of the plastic particle size on degradation

The occurrence of degradation reactions will also depend on the size of the plastic piece. Mechanical forces play more an effect in larger plastic pieces, where shear forces significantly contribute to the process and accelerate the breakdown into microplastic and even nanoplastic.<sup>45,60</sup> With declining size, the surface area is becoming more dominant compared to the bulk of the material. It is therefore expected that reactions that target the functionality of the surface and the underlying layer





would speed up the degradation process. Hydrolytic reactions, for example, are accelerated when the plastic particles become smaller, as these reactions rely on direct access to functional groups on the surface.<sup>61</sup> This, however, does not mean that plastic pieces quickly dissolve into monomeric building blocks once they are nano-sized. At this stage, the morphology of the plastic particle will influence the process. Since most polymers are semi-crystalline and chain scission mainly occurs in the amorphous regions, crystalline areas tend to resist further degradation.<sup>49</sup> In this study, the authors emphasized that the fact that commercial polymers are often semi-crystalline is an important aspect of understanding the degradation of micro- and nanoplastics.

### Effect of the type of environment

Plastic degrades through a combination of mechanisms, resulting in a wide variation in degradation rates depending on the location. Factors like soil or water, hot sunny locations or dry cold climates, mechanical stress or undisturbed conditions, all create unique chemical reactions that can lead to plastic breakage to microplastics and then nanoplastics.

**Water.** A noticeable difference between micro- and nanoplastics can be observed in water.<sup>62</sup> Nanoplastics have a strong tendency to aggregate as attractive forces, which are exerted by surface functionalities, might outweigh repulsive forces. They can aggregate with themselves or with other particles found in water, such as clay or natural organic matter (NOM). While these forces are also found in other media such as soil or food, the rate of aggregation, which is a function of viscosity among other parameters, can be substantial.<sup>63</sup> However, certain functional groups, as well as low salinity or favourable coating by natural matter, can ensure colloidal stability. This means that nano-sized particles will remain in solution for an extended time and Brownian motions control their movements. In contrast, aggregated nanoplastic or microplastics can settle in a few seconds or become buoyant. Therefore, the size, in combination with the physical properties, can determine whether the plastic pieces will be transported with the flow of the water or sink into the sediment.<sup>62</sup> This means that they will be located in different environments, such as varying UV intensities, which will further influence degradation.

**Air.** Microplastics and nanoplastics can form aerosol particles that are transported through the air for an extended period of time.<sup>64</sup> Here, they are subject to mainly photodegradation and oxidation and some form of wind abrasion.<sup>65</sup> The residence time in the air, also expressed by the settling velocity, depends on their size, shape, and density. While the residence time increases with size decrease, shape also plays a role although this effect is more complex. A theoretical model predicting the residence time of plastic nanofibers in the air identified the cross-sectional shape as the primary factor influencing long-range fiber transport, with flat fibers showing a 450% increase in residence time compared to cylindrical fibers.<sup>66</sup>

**Agroecosystem.** In soil, the plastic particles are in contact with soil components, which may be natural, but can also be of

synthetic origin (fertiliser or pesticides).<sup>67</sup> As the relative surface area increases when the plastic particles get smaller, this dynamic interface will determine further degradation by the various degradation pathways discussed above. Nanoplastics sit alongside black carbon and natural carbonaceous nanoparticles, which, to our current knowledge, outnumber the amount of nanoplastic. However, there is currently little known about how both interact and contribute to plastic degradation. These natural nanoparticles are internalised by plants when they are smaller than 100 nm, and there is evidence that nanoplastics will do the same.<sup>67,68</sup> Once inside the plant, nanoplastic may cause phytotoxic or genotoxic effects, but this field is still in its infancy and the researchers need to rely on knowledge obtained using engineered nanoparticles. It is safe to say, the uptake into plants by endocytosis, ion transport channels, aquaporins or other microbes such as bacteria and other pathways will again affect the surface properties and degradation.<sup>69</sup>

**Food.** There is only a short path from air, plants, and water for micro- and nanoplastics to enter animals and plants, thus food.<sup>70</sup> Micro- and nanoplastics have now been found in most food items.<sup>70</sup> In living systems, plastics can be subject to biodegradation,<sup>71</sup> but heating food and mixing it with other ingredients can potentially accelerate degradation. Extensive studies examined the formation of micro- and nanoplastics during cooking and food preparation when plastic-based kitchen utensils are used,<sup>72</sup> but our understanding of the fate of food-derived plastic particles during cooking remains limited.

Degradation in the environment is complex and often based on a feedback system. For example, the presence of microplastics was found to affect earthworms, who responded by increasing the number of burrows and thus may increase earth porosity.<sup>73</sup> These changes in soil structure will subsequently influence plastic degradation. The literature is filled with complex relationships between plastic particles and living organisms, emphasizing the challenge of fully understanding the effects of plastics.

### Identifying nanoplastics

One of the issues, however, is the difficulty in detecting nanoplastics. Current analytical techniques for analyzing plastic pollution are more tailored towards identifying the type of plastic than the particle size. Spectroscopic techniques such as Raman, Fourier Transform Infrared (FTIR) spectroscopy, or even nuclear magnetic resonance (NMR), paired with thermo-analytical techniques such as gas chromatography-mass spectrometry (GC-MS) (often coupled with other techniques) or differential scanning calorimetry (DSC), help to identify the type of plastic and they are used to analyse micro- and nanoplastics. However, the size and shape identification of nanoplastics is more challenging.

**Microscopy and light scattering techniques.** Widely used analytical techniques for determining the size and shape of the particles include microscopic techniques such as light- and fluorescent microscopy, with only scanning electron microscopy



(SEM) providing some insight into the presence of particles below 1  $\mu\text{m}$ .<sup>74,75</sup> However, direct size analysis is more complex because typical techniques, including light scattering, must be free of other natural nano-sized particles like protein (aggregates) and viruses, and the sample should also be relatively monodisperse. Additionally, smaller particles produce lower scattering intensity, making it difficult to analyze very diluted solutions. Nonetheless, techniques such as dynamic light scattering (DLS) have been successfully used to detect nanoplastics, but purification by separation and enrichment of nanoplastics is necessary.<sup>76</sup> Recent developments in the field of nanoplastic analysis have enabled the detection of nanoplastics in bottled water without the need for enrichment. The developed hyperspectral stimulated Raman scattering (SRS) imaging platform, which uses an additional Stokes laser to maximize the Raman signal to allow detection of nanoparticles of 100 nm, was able to identify high amounts of nanoplastics at concentrations previously unknown.<sup>77</sup>

**Combined techniques.** These techniques present an advancement as they facilitate analysis by enabling enrichment, separation, and identification in a single process. This includes the combination of AC electro-osmotic flows that collect nanoplastics on a hole-patterned electrode, followed by Raman identification of the nanoplastics (dielectrophoretic tweezing).<sup>78</sup> Raman tweezers, which are optical tweezers combined with Raman spectroscopy, are an alternative way to trap nanoplastics while allowing chemical identification at the same time.<sup>79</sup> Generally, combined techniques are more useful for identifying nano-sized objects in a diverse background. Transmission electron microscopy (TEM) is the most common way to extract size and shape from a nano-sized sample. However, when mixed with tissues, cells, bacteria, or other soft materials, polymers are not easily identifiable. Correlative Light and Electron Microscopy (CLEM) combines the capabilities of fluorescent microscopy with TEM to identify the location of synthetic material and determine its size and shape.<sup>80</sup> This technique could be used to locate plastic particles in tissue after staining them with Nile Red, a dye that is used to stain plastic with different plastic particles, causing slightly different Stokes shifts.<sup>81</sup>

#### Advanced enrichment techniques based on nanotechnology.

There is now a range of new solutions that aim to identify specifically nanoplastics in complex mixtures.<sup>76,82</sup> Many of these techniques are based on the idea of isolating nanoplastic based on a sieve principle, using a membrane,<sup>83</sup> a metal organic framework (MOF)<sup>84</sup> or an optical sieve.<sup>85</sup> The principle is often used to enrich nanoplastics to facilitate further analysis.<sup>76</sup>

Examples include a membrane based on nanowires that can capture nanoplastic, which are then detected with surface-enhanced Raman spectroscopy (SERS).<sup>83</sup> The cavities in MOFs can be used to extract nanoplastics from solutions as shown with a chromium-based MOF.<sup>84</sup> Researchers have developed an optical sieve that can detect nanoplastic and their size using Mie void resonances. According to the authors, this technique could be combined with  $\mu\text{-FTIR}$  or  $\mu\text{-Raman}$  spectroscopy to elucidate the structure of the nanoplastic particles.<sup>85</sup>

The reader is referred to comprehensive review articles that discuss and compare available techniques to detect micro- and nanoplastics, including the advancements in microscopy, spectroscopy, and AI-supported techniques.<sup>75,76,86</sup>

Despite the difficulty of identifying nanoplastic, they have now been widely observed in nature. In 2017, researchers found nanoplastics in the North Atlantic,<sup>87</sup> but since then, nanoplastics have been discovered everywhere, including in 2021 in soil,<sup>88</sup> in 2022 in polar ice,<sup>89</sup> in 2021 in the snow,<sup>90</sup> and in tissue in humans, as it will be discussed below. The recently published study outlined the widespread occurrence of nanoplastic in the ocean.<sup>5</sup>

## Nanoparticles may not be nanoplastics in their biological environment

However, just because nano-sized plastic pieces were detected under the microscope, it does not mean they exist as isolated pieces in the environment.<sup>91</sup> As discussed in the beginning, a reduction in size will lead to a larger surface area. As a result, the type of functional groups on the surface will influence what happens to these nanoparticles. Hydrophobic groups or hydrophilic and potentially charged groups will determine if these nanoparticles are colloidally stable or if they aggregate. In solution, this is described by the Derjaguin–Landau–Verwey–Overbeek (DLVO) theory. This theory combines attractive van der Waals forces and electrostatic repulsion, which is caused by the double layer of ions, also discussed in terms of the zeta potential of the nanoparticle. The total potential energy describes the balance between both forces and thus determines stability. While the colloidal stability of engineered nanoparticles is relatively well understood, the solution behaviour of micro- and nanoplastics is complex. The surface of these plastic particles will undergo a range of chemical changes during the weathering process.<sup>92</sup> Photooxidation usually leads to an increase in oxygen-containing functional groups, such as hydroxyl or carboxyl groups, that will help with solubility in aqueous solutions. This often coincides with changes in surface structure, such as increased roughness. This alone will already influence whether the plastic particles aggregate. However, the particles also strongly interact with other compounds present in their environment. In nature, this can include the formation of biofilm and the absorption of environmental contaminants such as metal ions and organic compounds that change surface properties.<sup>92</sup> Depending on the chemistry of these coated plastic pieces, they can either be stable in the aqueous system or start aggregating randomly.<sup>93</sup> The rate of aggregation will be determined not only by the composition of the plastic surface but also by the nature of the surroundings.<sup>93</sup> The size here plays an important role as nanoparticles aggregate faster than micron-sized particles due to their high surface energy.<sup>94</sup> Once the plastic pieces are ingested, they might be subject to further degradation as the low pH value of the stomach or the different enzymes might alter the surface properties again. Particles that



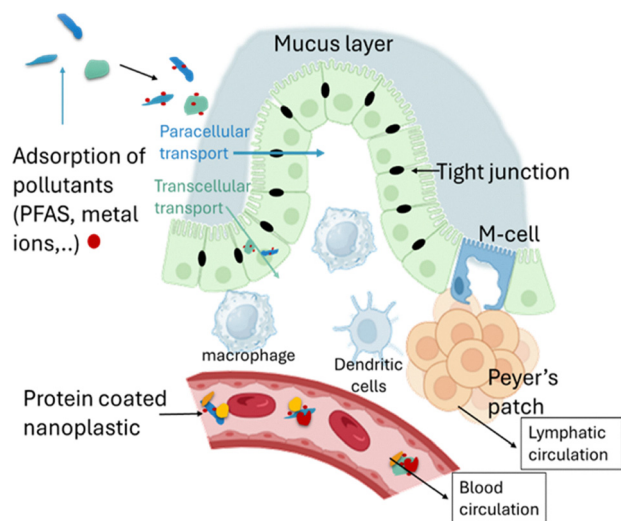


Fig. 3 Transport of micro- and nanoplastics from the stomach or intestine to the blood or lymphatic circulation (parts of this picture were created with <https://BioRender> (<https://BioRender>)).

enter the bloodstream will most likely be covered by a protein corona, which will determine the ultimate fate of the plastic pieces (Fig. 3). The types of proteins on the surface and their denaturation state will influence biodistribution, as well as whether the plastic pieces remain stable in the bloodstream.<sup>95</sup>

## How much plastic is in our body?

Humans are exposed to plastics in numerous ways.<sup>96</sup> As mentioned above, a significant amount of plastic is ingested through contaminated food or water, but plastic can also enter the body when previously uncontaminated food or water comes into contact with plastic containers that release plastic particles upon heating or other degenerative forces. Depending on the type and size of the plastic pieces, plastic is excreted through the stool. In a study with healthy volunteers, around 20 microplastic pieces between 50–500  $\mu\text{m}$  in size per 10 g of stool were detected.<sup>97</sup> This amount of particles found in faeces can be directly linked to the exposure to plastic food packaging.<sup>98</sup> This will mean that plastic will go back into the ecosystem.<sup>99</sup> Other options include the inhalation of plastic particles from the air.<sup>100</sup> Moreover, the skin is frequently in contact with plastic, whether it is clothing or other plastic items, such as phones or personal care products. Healthy skin usually serves as a barrier,<sup>101</sup> although it was proposed that nanoplastics might display some toxicological effects on the skin.<sup>102</sup>

The main question on everyone's mind is whether micro- and nanoplastics are dangerous to humans. First of all, we need to explore where plastic can be found in the body. This is not an easy undertaking, as many organs can only be analysed after the person is deceased. It is therefore not surprising that we know the most about micro- and nanoplastic by analysing the placenta (Table 1). In the first study, published in 2021, microplastics were detected in 4 out of the 12 placenta samples, with particles measuring approximately 10  $\mu\text{m}$  in size, primarily

composed of PP.<sup>103</sup> Another study found microplastic pieces larger than 50  $\mu\text{m}$  in the placenta,<sup>104</sup> but also in the baby's first stool, the meconium, which is clear evidence that microplastic is transferred from the mother to the baby. Microscopy techniques, such as FT-IR or Raman, were typically the method of choice for counting micron-sized microplastic particles. Occasionally, fluorescence microscopy was also used after visualising the microplastic with dyes such as Nile Red (Table 1). While seeing these large plastic pieces in organs can be confronting, a real game changer was the introduction of pyrolysis-gas chromatography and mass spectrometry (Py-GC-MS), which is capable of quantifying plastic in tissue, independent of the particle size.<sup>105</sup> However, a very recent study raised concerns about its suitability in complex media like blood, indicating that the concentration values, particularly for PVS and PE, are not reliable.<sup>106</sup> In a 2024 study by Garcia *et al.*, between 6.5 and 685  $\mu\text{g}$  of plastic was found in one gram of placental tissue (mean  $126.8 \pm 147.5 \mu\text{g g}^{-1}$ ).<sup>107</sup> The amount varied widely, with a specific measurement of 685  $\mu\text{g}$  found in the placenta of one mother, equating to more than 0.06 wt%. Let's compare microscopy samples with these results: microplastic pieces vary widely in size and shape, but let's assume one piece with a prolate shape with a length of 10  $\mu\text{m}$  and a thickness of 2  $\mu\text{m}$ , which equates to a volume of  $V = 20.9 \mu\text{m}^3$ . A piece like that would weigh approximately  $2 \times 10^{-5} \mu\text{g}$ , which is only a fraction of the 126.8  $\mu\text{g}$  of polymer per g tissue found in the study by Garcia *et al.* (Table 1). This calculation is only a rough example, but it highlights that microscopy might reveal less than 1% of the plastics present in the tissue. The invisible part, nanoplastics, makes up more than 99%, and can only be visualized by techniques that are suitable for nanoscopic analysis. Table 1 lists how microscopy studies uncovered a few microplastic pieces in a gram of tissue in various organs, while Py-GC-MS exposed large plastic quantities that can only be assigned to nanoplastic pieces. We are only at the beginning of understanding how much plastic in the form of nanoplastic has accumulated in the body, but the initial studies provide an insight that is concerning.

## How does plastic get into organs?

The fate of these plastic particles in the lungs (after inhalation) or the stomach (after ingestion) depends on the properties of the particles. Size will play a major role, as will shape and surface chemistry. Plastic that will find its way into the stomach will be exposed to very acidic environments and a gastrointestinal epithelium that does not allow the transport of larger molecules from the stomach to the blood. In general, the gastrointestinal (GI) tract, encompassing the oesophagus, stomach, and intestine, is relatively impermeable to many compounds.<sup>120</sup> The transport across these barriers has been the focus of attention in the field of nanomedicine. Researchers have developed particles of different sizes, shapes, and properties and investigated how they can cross these impermeable barriers. Nanoparticles can cross through the transcellular





**Table 1** Micro-and nanoplastic found in humans (Py-GC-MS: pyrolysis-gas chromatography and mass spectrometry; LDIR: laser direct infrared imaging; IRMS: isotope ratio mass spectrometry)

| System              | Organ                                     | Analytical technique   | Concentration: wt plastic per wt biological sample  | Amount of pieces  | Size of plastic particles  | Dominant type of plastic     | Ref.  |
|---------------------|---|--|---|---|--|------------------------------|---|
| Nervous system      | Brain (2024)                              |  | 4917 $\mu\text{g g}^{-1}$   |   | Below 200 nm, spherical and anisotropic  | PE                           | Nihart <i>et al.</i> <sup>11</sup>          |
|                     | Digestive system and urinary system       |  | 433 $\mu\text{g g}^{-1}$  |   | Rod-shape 1–5 $\mu\text{m}$ (potentially aggregates) and smaller shard-like below 400 nm   | PE, PP                       | Nihart <i>et al.</i> <sup>11</sup>          |
| Circulatory system  | Liver (2022)                              | Fluorescence microscopy<br>$\mu\text{Raman}$<br>spectroscopy |   | 0 particles per g (kidney)<br>1.1 particles per g (spleen)<br>0.7 particles per g (healthy liver)<br>8.4 particles per g (liver with cirrosis)  | 3.3 to 20.1 $\mu\text{m}$  | PS, PVC, PET                 | Horvatis <i>et al.</i> <sup>108</sup>       |
|                     | Kidney (2024)                             |  | 404 $\mu\text{g g}^{-1}$  |   | Rod-shape 1–5 $\mu\text{m}$ (potentially aggregates) and smaller shard-like below 400 nm   | PE, PP                       | Nihart <i>et al.</i> <sup>11</sup>          |
|                     | Blood                                     | Py-GC/MS   | 1.6 $\mu\text{g ml}^{-1}$   |   | Size not reported  | PET, PS, PP                  | Lesli <i>et al.</i> <sup>109</sup>          |
|                     | Carotid plaque (2024)                     | Py-GC/MS<br><br>TEM<br>SEM<br>IRMS                           | 21.7 $\pm$ 24.5 $\mu\text{g g}^{-1}$ (PE)<br>5.2 $\pm$ 2.4 $\mu\text{g g}^{-1}$ (PVC)                           |   | Jagged-edged nanoparticles below 1 $\mu\text{m}$   | PE, PVC                      | Marfella <i>et al.</i> <sup>110</sup>       |
| Reproductive system | Corona                                    | Py-GC/MS   | 118.66 $\pm$ 53.87 $\mu\text{g g}^{-1}$   |   | Size not reported  | PET                          | Liu <i>et al.</i> <sup>111</sup>            |
|                     | Coronary artery                           |  |   | 60–75 504 particles per g   | 20 to 469 $\mu\text{m}$  | PET, PE                      | Yang <i>et al.</i> <sup>112</sup>           |
|                     | Carotid artery                            | LDIR   |   | 12 pieces in the placentas of four women of samples sizes of 23.3 $\pm$ 5.7 g   | Fibre-like 5–10 $\mu\text{m}$  | PP                           | Ragusa <i>et al.</i> <sup>103</sup>         |
|                     | Heart, different parts                    | SEM  |   | All the analyzed MPs were pigmented.  |  |                              |   |
|                     | Placenta (2021)                           | Visible light<br>Raman microscopy                            |   | A particle found in 1 $\times$ 1 $\times$ 1 cm Placenta sample and in some Meconium samples   | > 50 $\mu\text{m}$   | PS, PP                       | Braun <i>et al.</i> <sup>104</sup>          |
|                     | Peripheral and central placenta, meconium | FT-IR microscopy   |   | 2.70 $\pm$ 2.65 particles per g   | 20.34 to 307.29 $\mu\text{m}$  | PVC, PP, PBS, PET            | Zhu <i>et al.</i> <sup>113</sup>            |
| Respiratory system  | Placenta                                  | LDIR   |   | 153.07 $\pm$ 84.45 per 0.4 g  | Size not reported  | PE dominant                  | Garcia <i>et al.</i> <sup>107</sup>         |
|                     | Placenta                                  | Fluorescence microscopy                                      | 126.8 $\pm$ 147.5 $\mu\text{g g}^{-1}$  |   | Size not reported  | PE dominant                  | Jochum <i>et al.</i> <sup>114</sup>         |
|                     | Preterm placenta                          | Py-GC-MS   | 224.7 $\pm$ 180.7 $\mu\text{g g}^{-1}$<br>175.5 $\pm$ 137.9 $\mu\text{g g}^{-1}$<br>328.44 $\mu\text{g g}^{-1}$ |   | Size not reported  | PE dominant                  | Hu <i>et al.</i> <sup>115</sup>             |
|                     | Placenta                                  | Py-GC-MS   |   | 0.23 $\pm$ 0.45 particles per mL (Semen)  | 96.19 $\pm$ 74.17 $\mu\text{m}$ (fibres, films, flakes and others)   | PVC and PE (semen)           | Zhao <i>et al.</i> <sup>116</sup>           |
|                     | Testes                                    | LDIR   | 15.34 $\pm$ 23.31 $\mu\text{g g}^{-1}$ (semen)  | 11.60 $\pm$ 15.52 particles per g (testis)<br>0.56 particles per g  | 83.15 $\pm$ 56.25 $\mu\text{m}$<br>5.5 $\mu\text{m}$ particles, and fibres of 8.12 to 16.8 $\mu\text{m}$                                       | PS (testis)<br>PP and PE     | Amato-Lourenco <i>et al.</i> <sup>117</sup> |
|                     | Semen                                     |  |   | upper (0.80 $\pm$ 0.96 MP g <sup>-1</sup> ), middle/lingular (0.41 $\pm$ 0.37 MP g <sup>-1</sup> ), and the lower (3.12 $\pm$ 1.30 MP g <sup>-1</sup> )<br>24 microplastics (> 20 $\mu\text{m}$ ) in 100 human lung tissues | 223.10 $\pm$ 436.16 $\mu\text{m}$ (length) and 22.21 $\pm$ 20.32 $\mu\text{m}$ fibre, fragment, films<br>1000–5000 $\mu\text{m}$ length fibres | PP and PET                   | Jenner <i>et al.</i> <sup>118</sup>         |
| Respiratory system  | Lung                                      | Raman spectroscopy   |   |   |  | Polyester and natural fibres | Chen <i>et al.</i> <sup>119</sup>           |
|                     | Lung                                      | Light microscopy<br>$\mu\text{FTIR}$                         |   |   |  |                              |   |
| Respiratory system  | Lung                                      | $\mu\text{FTIR}$   |   |   |  |                              |   |
|                     | Lung                                      | SEM  |   |   |  |                              |   |



route through epithelial cells, paracellular pathways between cells along the tight junctions, using the M-cells of the Peyer's patches or *via* receptor and transcytosis-mediated endocytosis (Fig. 3). While many of these pathways are irrelevant for larger micron-sized particles, the M cells in Peyer's patches can internalise microplastics of several micrometers. Similar to all cell internalization pathways, the uptake by M cells depends on the types of proteins that can be found on the surface.<sup>121</sup> It appears, though, that there is an upper size limit as plastic particles that are larger than 150  $\mu\text{m}$  are predominantly excreted.<sup>122</sup> Translocation into the bloodstream increases as the size of the plastic particle decreases because additional transport pathways open up (Fig. 3). Particles in the low micron size can now use phagocytosis or endocytosis to move into the bloodstream, albeit the paracellular route can be ruled out when the particles are still in the microrange.<sup>123</sup> Uptake is further facilitated when these plastic pieces are below 1  $\mu\text{m}$ . A study on engineered nanoparticles of various sizes showed that nanoparticles below 100 nm have significantly higher permeation across intestinal cells than spheres of 1000 nm.<sup>124</sup> The authors also prepared disk- and rod-shaped nanoparticles and observed significant uptake. Understanding the unique behavior of anisotropic nanoparticles is important because TEM analysis of plastics in organs revealed that most pieces are non-spherical (Table 1 and TEM insert in Fig. 2).

The field of nanomedicine also provides us with plenty of studies on the effect of size and surface properties on bio-distribution.<sup>125</sup> Particles with sizes between 50 and around 100 nm usually have a long circulation time in the bloodstream as they are only slowly detected by the mononuclear phagocytic system (MPS).<sup>125</sup> A landmark paper also showed the significantly extended circulation time when non-spherical, worm-like nanoparticles were used.<sup>126</sup> The differences in circulation time between spherical nanoparticles and other shapes are crucial when dealing with nanoplastics. TEM analysis revealed a myriad of shapes, which are the result of various erosion processes, but the shapes are rarely spherical. Moreover, the surface chemistry will play a role. In nanomedicine, blood clearance is often evaded by coating nanoparticles with polymers such as poly(ethylene glycol) PEG and zwitterionic structures that can introduce stealth properties to the nanoparticles.<sup>127</sup> The surface chemistry of nanoplastic is given by the degradation mechanism. While commodity plastics are hydrophobic, introducing functional groups will not only enhance the colloidal stability, but it will also influence the interaction with biological entities such as proteins, as discussed earlier. A protein corona that contains predominantly opsonin proteins will have a shorter circulation time than one with dysopsonins.<sup>128</sup> After circulating in the bloodstream for a few hours, or in some cases several days, the nanoparticles begin accumulating in various organs, primarily in the liver and spleen, unless they have been excreted by the kidneys. It is therefore not surprising that large amounts of micro- and nanoplastics have been found in these two organs (Table 1). The final destination of nanoparticles is, however, hard to predict, although attempts have been made to link the properties of nanoparticles with *in vivo* outcomes.<sup>129</sup> Even very small changes

to the nanoparticle properties can result in different protein coronas and biodistributions, which is explored in nanomedicine to target a range of organs.<sup>129</sup>

While the reader probably quietly accepts that plastic was found in the liver, an organ whose role is to collect and excrete unwanted products, discovering that plastic was found in the brain is concerning.<sup>11</sup> Nanoparticles are widely developed as a way to treat brain diseases such as brain cancer, neurodegenerative diseases, or Alzheimer's disease. However, the primary challenge that still exists is effectively delivering nanoparticles across the BBB.<sup>130</sup> That engineered plastic nanoparticles, which are spherical in shape and made from commodity polymers, can penetrate this layer was indeed shown in animal studies.<sup>131</sup> Key to brain entry was not only the size of the particle,<sup>132</sup> as only nanoplastics, but not microplastics, could cross the BBB, but also the composition of the protein corona.<sup>132</sup> Indeed, from the nanomedicine field, we know that the smaller nanoplastic pieces are, the more dangerous they will be in crossing the BBB.<sup>133</sup>

## Does more plastic exposure lead to higher accumulation in the organs?

The studies listed in Table 1 only had a handful of participants in each study, and it is difficult at this point to draw clear conclusions. In general, there seems to be some correlation between what we eat and drink and the amount of plastic found in our bodies.<sup>98,134</sup> However, the authors listed in Table 1 all exercise caution about potential relationships, as the number of samples is often limited. There seems to be, for example, a moderate correlation between the amount of bottled water a volunteer drank and the microplastic pieces in faeces.<sup>135</sup> However, in this study, the authors only counted microplastics, while the amount of nanoplastics may be huge but invisible. In another study, this time involving infants, a PET content of 36  $\mu\text{g}$  per gram of stool was measured using solid-phase extraction (SPE). In contrast, only 2.6  $\mu\text{g}$  was found in adults. The excessive use of plastic products like bottles during infant feeding could explain the difference.<sup>136,137</sup> In a study that tried to compare the consumption of bottled water with the occurrence of microplastics in semen, no such correlation was found.<sup>116</sup> A direct correlation between plastic consumption and enhanced accumulation can, therefore, not confidently be drawn at this stage, as there are probably too many factors that play a role.

One would also expect plastic accumulation to increase with age, but the absence of a direct correlation highlights the complexity of accumulation and clearance. For example, the concentration of microplastics in human testes was found to be independent of age and more likely related to the process of sperm production.<sup>115</sup> Similar observations were made when linking the amount of plastic in the brain with age, where there was a lack of correlation.<sup>11</sup> Yet, when comparing samples collected eight years ago with recent samples, there was a noticeable increase in the amount of plastic deposited in the



organs, which is in line with the increase in plastic pollution in the environment.<sup>11</sup> It appears that plastic accumulation in the organ plateaus as incoming plastic is also slowly cleared again, as observed in a zebrafish study.<sup>138</sup> In contrast, when analysing the presence of microfibrils in lung tissue, the authors found a clear increase with age.<sup>119</sup>

The concentration in various organs seems to be a function of the unique physiology of each living being. An enhanced concentration of microplastics was found in patients with liver cirrhosis. It was proposed that the “leaky gut” in these patients allows additional migration of microplastics into the liver.<sup>108</sup> A huge amount of nanoplastic was measured in the brains of dementia patients. It was highlighted that it is likely the inability of these brains to purge nanoplastic, although it might be tempting to link nanoplastics with dementia.<sup>11</sup> However, the authors state clearly that this is not an indication that nanoplastic will cause dementia, but they also warn that more needs to be done to understand the effect on human health. At this point we do not have enough knowledge to separate cause and effect. Does the enhanced accumulation trigger a disease, or does the disease lead to enhanced accumulation? An example of the former might be the higher amount of microplastics found in lung cancer,<sup>119</sup> where it appears that plastic fibres in ground-glass nodules (GGNs) in the lung might have contributed to the progression of the disease.

## Is nanoplastic dangerous?

The complex relationship between the plastic accumulation of microplastics in various organs and potential health effects has been subject to a range of reviews,<sup>96,123,139–143</sup> therefore, only a brief overview is given here.

Ultimately, the question will be whether micro- or nanoplastic will make me sick? Unfortunately, we do not have sufficient human data yet to answer this, but there are enough *in vitro* and *in vivo* data using animal studies to show that we should be concerned. There is a large body of work on the toxicity of micro- and nanoplastics that proposes that plastic in our body might cause adverse health effects. However, setting up suitable studies is complex. In many cases, the researchers used engineered microplastics, which are often spherical particles made from commodity polymers, with a well-defined surface chemistry. Apart from micro- and nanoplastics being irregular in shape, we also know that plastic composition from various sources can vary substantially, as each plastic item produced has a different mixture of additives. However, these engineered nanoparticles allow us to gain some insight into potential toxic effects.

The initial step in toxicity testing is to study micro- and nanoplastics on cell lines similar to what researchers in the nanomedicine sector would do. From there, we know that nanoplastics can be readily taken up by a large variety of different cells while microplastics struggle to gain entry.<sup>96</sup> Macrophages, which play an important role in the immune system, are known to phagocytose even micron-sized particles.<sup>80</sup>

The more particles enter the cells, the more likely they are to cause cell stress or even cell death; therefore, size-dependent effects are observed in some reports.<sup>144,145</sup> Plastic particles often do not directly display cytotoxicity, only at relatively high concentrations.<sup>146,147</sup> Again, this is often a function of size as nanoplastics in the low nanometer range were found to be toxic already at low concentrations, while larger particles require higher concentrations to be cytotoxic towards various phagocytes.<sup>145</sup> Cytotoxicity is often not directly observed but manifests as a range of more subtle interferences with cells since nanoplastics can interfere with the mitochondria,<sup>148</sup> inhibit membrane transporters,<sup>148</sup> trigger the production of reactive oxygen species (ROS),<sup>144,146</sup> or cytokines<sup>144</sup> or lead to an increase in a range of other cell markers that might indicate cell stress.<sup>145,149</sup> The effects of micro- and nanoplastics on cytotoxicity, immune response, oxidative stress, barrier properties, and genotoxicity were compared in a *meta-analysis*, and the authors found that the shape of the plastic particle is a much stronger predictor of toxicity than its size or polymer type.<sup>150</sup> Nanoplastic pieces with rough edges and irregular shapes showed noticeably more toxicity than spherical ones, which can be concerning, considering that most naturally occurring nanoplastic pieces are non-spherical.<sup>151</sup> It also highlights that engineered spherical nanoparticles may not be the best models to test plastic toxicity.

Organoids and spheroids have recently emerged as an alternative tool to study the effects of plastic particles.<sup>152–156</sup> Next to toxicology information, organoids and spheroids can provide information on the ability of plastic pieces to penetrate into tissue.<sup>157</sup> Cortical or cerebral, kidney, cardiac, lung or liver organoids could reveal significant damage after exposure to predominantly nanoplastic particles as small as 100 nm.<sup>156</sup> Organoids were able to uncover various toxic effects, but according to the author of a recent review paper, organoids need to become more complex and include features such as vasculature and immune cells. Additionally, the authors recommended that the experiments should better simulate long-term exposure to nanoplastics.<sup>156</sup>

What is, however, on people's minds is how nanoplastic might affect their health or their ability to have healthy children. As highlighted earlier, we don't have enough data on human studies, but *in vivo* work using animals can shed some light on this.

### Digestive system and urinary system

The first line of contact of micro and nano plastics is with the digestive system and urinary system. Particles that are ingested will potentially damage the stomach or intestine, and after entering the bloodstream, will be transported to the kidneys, liver, and bladder. It is evident that damages already occur in the stomach and intestine, with the various effects being summarized in review articles.<sup>158,159</sup> Feeding mice with 5  $\mu$ m PS particles over 28 days led to oxidative damage, inflammation, and disturbances of the intestinal microbiota.<sup>160</sup> This led further to an increase in undesirable gut bacteria and increased levels of interleukin-1 $\alpha$  in blood serum, a hallmark of inflammation as observed after feeding mice with micron-sized PE.<sup>161</sup>



Similar results were observed after feeding micron-sized PE to zebrafish, including intestinal inflammation and increased infection probability.<sup>162</sup> While a pro-inflammatory response has been observed with nanoplastics *in vitro*,<sup>163</sup> there is only a limited number of studies that look at the effects of nanoplastics in animal models. A direct comparison of micro- and nanoplastic on zebrafish showed that while all particles affected the microbiome and the intestinal immune cells, only nanoplastic led to genetic changes that led to increased ROS production and mucus secretion.<sup>164</sup>

The observed disturbance of the microbiome will affect the gut-liver axis and can potentially result in liver diseases such as liver fibrosis and cirrhosis, as reviewed here.<sup>165</sup> Accumulation of microplastics in the liver is widely observed, which is not surprising considering that the liver and spleen filter the blood to remove harmful substances. Once lodged there, microplastics can cause oxidative stress, alter metabolism, increase cell death, and inflammation.<sup>165</sup> Most studies used microplastics, but there are a few studies using fish that show that nanoplastics are just as dangerous.<sup>166</sup> Zebrafish that were exposed to 50 nm PS particles for 28 days showed not only signs of liver damage, but the authors found evidence for the depolymerization of PS in the liver by the cytochrome P450 enzymes, which may lead to the formation of cytotoxic styrene and styrene oxide.<sup>167</sup> A review that directly compared studies on microplastics and nanoplastics and their effect on the liver found that microplastics are more likely to accumulate in the liver and cause hepatotoxicity.<sup>158</sup> According to these initial reports, it appears that microplastics could be more damaging than nanoplastics to the liver.<sup>158</sup>

### Nervous system

Microplastics can cause significant neurotoxicity in mammals and fish,<sup>168,169</sup> and an increasing number of studies now include nanoplastics. Micro- and nanoplastics were directly compared when mice were given drinking water containing PS particles with diameters of 500 nm, 4, and 10  $\mu\text{m}$ , respectively, over 180 days. The smallest nanoparticles were able to cross the BBB to a significant extent, but nevertheless, the authors found no size effect on subsequently studied cognitive dysfunction, only a concentration effect.<sup>170</sup> This does not mean that nanoplastic is not potentially more dangerous, as shown in studies on fish using 70 nm and 5  $\mu\text{m}$  PS particles. While both particles led to significant toxic effects, nanoplastic reduced the activity of acetylcholinesterase (AChE), an enzyme that controls nerve transmission.<sup>171</sup> Several studies showed that nanoplastics, when fed to mice and rats, result in significant behaviour changes, cellular dysfunction, as reviewed in detail here.<sup>172</sup> For example, PS particles measuring 30–50 nm in size, and administered orally to mice, were capable of reaching the brain, where they induced cognitive impairment, most likely by modulating gene expression in microglia.<sup>173</sup>

### Blood, circulatory and lymphatic system

Once micro- and nanoplastics cross into the bloodstream, they will immediately be exposed to the innate immune system,

most likely to neutrophils.<sup>174</sup> Plastic particles have been widely found in the blood of animals and humans.<sup>175</sup> Several studies on fish exposed the changes in neutrophils and the subsequent effect on the fish population, which includes reduced disease resistance,<sup>174</sup> but also chromosomal harm to fish erythrocytes.<sup>166</sup> In mice, an increase in eosinophils and basophilic leucocytes<sup>176</sup> and altered lipid metabolism, as evidenced by elevated serum markers, were found.<sup>177</sup> While the effects of plastic particles on blood cells and cardiovascular parameters are evident as reviewed here,<sup>175</sup> there are few studies comparing micro- and nanoplastics *in vivo*. One study that investigated the effect of plastic particles of 500 nm and 5  $\mu\text{m}$  in size reported an increase in body weight after plastic ingestion. The mice fed with the 500 nm nanoplastic particles had significantly higher glucose levels than the control group.<sup>176</sup>

Once in the blood, the particle can be shuttled to the lymph nodes and various organs<sup>178</sup> with nano-sized plastic pieces reaching the lymph nodes more rapidly than microplastics.<sup>179</sup> Plastic was detected in humans in the aorta, coronary arteries, and carotid arteries, and the concentration was correlated to the disease history of the individual humans.<sup>110,111</sup> The authors suggested a potential connection between the presence of large amounts of nanoplastics in carotid plaques and the occurrence of stroke, myocardial infarction, and other deaths in patients.<sup>110</sup> Moreover, it is also believed that plastic particles might be linked to atherosclerosis.<sup>103</sup> However, the authors measured the content using Py-GC/MS and did not distinguish between micro- and nanoplastic, which might have shed light on the effect of size differences. Finally, the plastic pieces might reach the spleen, where they can cause adverse effects.<sup>159</sup>

### Reproductive system

The effects of micro- and nanoplastics on the reproductive system are widely studied and the subject of many review articles,<sup>180–183</sup> and the main effects are summarised in Fig. 4. In this field, there is a substantially greater volume of research focusing on human subjects relative to studies examining health effects on other organs. This disparity can be primarily attributed to the relative ease of obtaining biological samples such as placentas or semen, whereas obtaining other organs is generally limited to specimens from diseased individuals or those who have undergone partial organ resection. Consequently, studies involving the uterus, cervix, or testis tend to be scarce and often characterized by limited sample sizes. For example, uterine fibroids, which are benign tumors of the uterus, are usually removed together with the neighboring myometrial tissue during surgery.<sup>134</sup> It was observed that there was more microplastic in tissue that contained fibroids than in healthy tissue. What was particularly interesting about this study was the link between these findings and the lifestyle choices of the tissue donors. It was discovered that a higher consumption of takeaway meals and bottled water led to increased plastic accumulation, highlighting that most plastic found in the body is ingested directly through contaminated food.<sup>134</sup> Microplastics are also found in testis (on average 328.44  $\mu\text{g}$  per gram tissue)<sup>115</sup> and uterus (amount not determined).<sup>184</sup>



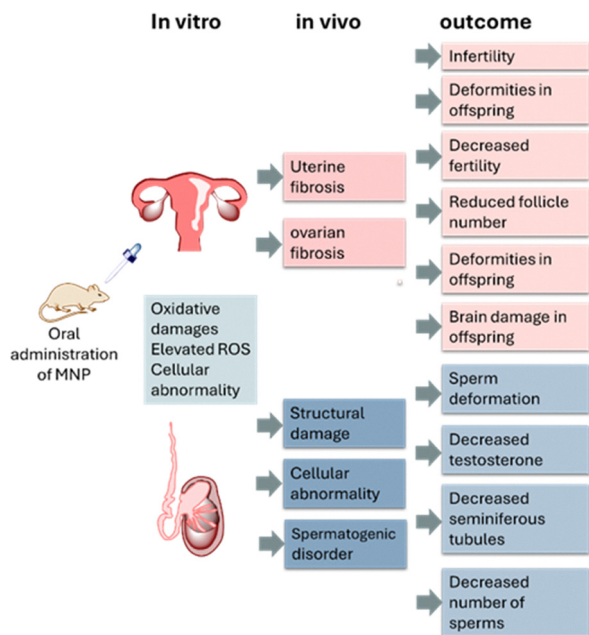


Fig. 4 Visual representation highlighting the impacts of micro- and nanoplastics (MNP) on the male and female reproductive system. Inspired and adapted from Ye *et al.*<sup>190</sup>

Analysis of tissue samples from cervical cancer patients showed that when microplastics were found, metabolic changes in cervical invasive cancer tissues were measured, but the researchers did not draw any conclusions about what this means for our health.<sup>185</sup> However, some concerning correlations have been reported in recent years. For example, reduced foetal growth is associated with an abundance of plastic in the placenta of women,<sup>186</sup> while plastic in the women's villous tissues is thought to lead to an increased occurrence of miscarriages.<sup>187</sup>

As the number of studies on humans is limited, researchers have used animal models to learn more about this topic. The extent to which microplastics may influence fertility rates or adversely affect offspring remains to be fully explored.<sup>180</sup> In mice, administration of plastic particles led to reduced fertility, an abnormal sex ratio in their offspring,<sup>184</sup> increased embryo resorption rate,<sup>188</sup> and embryonic growth retardation.<sup>189</sup> Several animal studies suggest that disruptions in the neuro-endocrine system *via* the hypothalamic-pituitary-gonadal (HPG) axis may affect the synthesis and secretion of sex hormones, in addition to endometrial dysfunction, atrophy, fibrosis, and other adverse effects, as reviewed here.<sup>180</sup> Comparison of micro- and nanoplastics in zebrafish led to the conclusion that nanoplastics might be able to reach areas such as the yolk sac, which microplastics cannot reach,<sup>180</sup> but in general, the number of studies comparing the effects of different sizes of plastic particles on reproductive health is limited.<sup>190</sup> A study that uses PS beads in the size of 100 nm, 3  $\mu\text{m}$ , and 10  $\mu\text{m}$  found a larger accumulation of nanoplastic in the testis compared to the microplastic particles, but not in the epididymis.<sup>191</sup> Nanoplastics are also more likely to cross the placental barrier to reach the foetal brain in mice.<sup>192</sup> Depending on the surface charge of

the nanoparticles, damages to the foetal brain by ROS were observed.<sup>192</sup> Next to damages to the brain of the unborn mouse during gestation, nanoplastic can also be more easily delivered from the female to the offspring through breast milk.<sup>193</sup> Both particles, micron- and nano-sized, reduced fetal growth, but this study on mice did not find significant differences between 50 nm and 5  $\mu\text{m}$  particles.<sup>194</sup> A study that followed the health of mice from the gestational stage to puberty after their mothers were fed plastic particles of 500 nm and 5  $\mu\text{m}$ , respectively.<sup>195</sup> The offspring showed metabolic changes although they were never exposed to plastic directly, which could either mean that the maternal metabolism was affected or that plastic particles passed through the placenta. Interestingly, this effect was more pronounced with microplastics.

### Respiratory system

It is not unexpected that plastic particles should have a profound impact on the respiratory system, considering that inhalation is a common pathway of entry. As reviewed elsewhere,<sup>100,196</sup> the deposition of plastic particles on the lungs can potentially cause respiratory diseases, as shown in animal models. However, Vasse and Melgert also concluded that more work is needed to understand the effect of actual environmental plastic pollution on human health.<sup>196</sup> However, it is reasonable to expect that nanoplastics might cause more damage than microplastics, as they are more likely to reach the respiratory bronchioles, alveolar ducts, and alveoli, whereas larger particles tend to remain in the upper respiratory system.<sup>197</sup>

In summary, there is evidence indicating that plastic particles may have detrimental health effects. However, it remains unclear at this point whether nanoplastics pose a greater or lesser risk compared to larger plastic particles, as studies that compare sizes directly are limited. Moreover, many studies employ engineered spherical PS nanoparticles while actual micro- and nanoplastic particles are often fibrous, rough with many edges, or film-like in appearance. Furthermore, certain studies employ exceedingly high concentrations of plastic particles in animal experiments (several milligrams of plastic per kilogram per day), which may not be representative of typical human consumption. While it was proposed that we eat the equivalent of one credit card per week (5 grams), Pletz revisited these numbers and proposed that 4.1  $\mu\text{g}$  per week is a more likely number.<sup>198</sup> Even with a 5 g intake by a 70 kg adult, the daily dose is 10 mg per kg, which is lower than the concentrations used in some studies.

### Is nanoplastic more dangerous than microplastic?

If we reduce the size of a plastic particle from 5  $\mu\text{m}$  to 50 nm, we increase the surface area relative to the volume by a factor of 100. Therefore, the types of functional groups on the surface will determine the fate of the plastic pieces. It is the degradation pathways, the degradation environment such as soil, air or water, the season, the type of polymer, and even subtle





differences in the constitution or stereochemistry of a polymer, such as molecular weight, branching and tacticity, that will influence the rate of degradation, but also the type of functional groups that will dominate the surface.<sup>199–201</sup> This interface will control if the nanoplastic pieces are stable in water or if and how they interact with other compounds.<sup>202</sup> Both processes are interrelated. Surface functionalities such as charged groups will ensure water-solubility, colloidal stability, but they will also decide with what organic molecule or biomaterial they interact. This corona of absorbed molecules will then decide the stability in water, which is either long-term colloidal stability or aggregation into larger micron-sized aggregates. Nano-sized particles are more likely to circulate around the blood stream for an extended period of time, they are more likely to enter cells in the body or even cross the BBB. In contrast, the surface of microplastics is relatively small. There is less room to absorb any molecules, such as pollutants, and they are likely to be cleared *via* the liver. Although more work needs to be done, nanoplastic is likely more dangerous.

## Responsible plastic use

It is evident that there are many unknowns about the effects of plastic particles on our health, but we need to err on the side of caution and reconsider how we use plastics. Plastic is a unique material that is irreplaceable in our daily lives for certain applications, but in other areas, we need to consider how to reduce plastic consumption, reuse items, or recycle them. An informed population is a prerequisite for this step. For example, it is not widely known that washing of clothes made from synthetic fibres such as Nylon is a huge contributor to microplastic pollution.<sup>203</sup> A recent citizen science project in the UK showed that households grossly underestimate how many plastic items they dispose of (plastic blindness),<sup>204</sup> but awareness of it can help with behaviour change. The “One Planet Network”, a global community of practitioners, policymakers and experts, governments, businesses, civil societies, academia and international organisations, has assembled strategies that help with behavior change.<sup>205</sup> While there is a focus on personal responsibility, the responsibility also lies with big corporations and lawmakers.

Plastic pollution can be reduced by better remediation techniques, such as the removal of plastic particles from storm water, better waste water sludge management, the development of a circular economy, better waste management and many other approaches<sup>203,206</sup> More importantly, policies need to be in place that aid the process. Tackling plastic pollution is a major aim of the UN environmental program,<sup>207</sup> and law makers around the world already act upon. Examples are the “National Plastic Plan” in Australia,<sup>208</sup> the “Plastic strategy” by the European Commission,<sup>209</sup> but also international agreements are being developed that seek to tackle the plastic problem (UNEA resolution (5/14)).<sup>210</sup> Policies include the phasing out of single-use plastic, improving waste management, reducing plastic leakage, and appropriately financing the

circular economy, with some responsibility given to the private sector.<sup>211</sup>

Until then, it is recommended to limit personal exposure to plastics. This includes the protection from airborne plastic particles, which is particularly crucial for workers in recycling plants. This aspect is often overlooked: recycling plants, where plastic is shredded, can be a source of micro- and nanoplastics formation.<sup>203</sup> On a personal level, households can reduce the amount of plastic particles in the air by using air purifiers equipped with HEPA filters, which can help alleviate exposure,<sup>212</sup> consider the use of plastic items in the kitchen and household,<sup>72</sup> and other plastic items, such as water in cheap plastic bottles.<sup>40</sup>

## The role of chemists and materials scientists

Understanding the health effects of micro- and nanoplastics will need to be the subject of many studies for biomedical researchers in the coming years, as we still do not understand the magnitude of the problem. This problem can only be tackled by combining the effects of interdisciplinary teams comprising analytical chemists, environmental chemists, food, soil, water scientists, toxicologists, and medical researchers. Materials scientists can play a central role in the discussion around plastic pollution as they can contribute to our understanding of the behaviour of plastic particles, but they can also be instrumental in helping to reduce plastic waste. The following outlines areas where polymer scientists can potentially contribute to the discussion:

### Making plastic more durable and repairable

This may be a strange demand, but it needs to be considered that plastic materials have properties that cannot be found with any other materials. Plastic materials are known to save lives and can help reduce energy consumption, among other favourable properties. There are plastic items we want to be durable and to last for a long time, yet many have been found to break down during usage.<sup>213</sup> Some plastic pollution is the product of abrasion, such as tires, or premature breakage. As a result, plastic is released in the environment or contributes to landfills. To extend the lifetime of a plastic consumer item, it is desirable to be able to repair it or reshape it into an alternative product. The development of self-healing plastic<sup>214</sup> or dynamic materials that can be upcycled<sup>215,216</sup> Even broken plastic might be a valuable raw material in such a case.

### Creating better micro-and nanoplastic reference materials

Many animal studies have used spherical PS particles that are commercially available as models to test the effects on health. However, we know from the plastic pieces found in various tissues that plastic particles are neither spherical nor made from PS, as the dominant plastic material found in tissue was often PE (Table 1). The impact of the shape on biodistribution and cell uptake is significant. Additionally, PS cannot be directly compared to PE because one is an amorphous polymer



and the other is highly crystalline, which influences degradation. Micro- and nanoplastics have diverse shapes and a broad size range,<sup>217</sup> making it challenging to predict their health effects. While it is feasible to use actual micro- and nanoplastic samples from nature, the issue would be that the size, composition and surface chemistry represent a very unique fingerprint of the geographic area and time point that created these specific plastic pieces. Materials scientists could potentially help to create models using accelerated aging conditions while providing in-depth structural characterization, such as size, shape, and surface functionalities. A variety of approaches have been summarised here,<sup>75</sup> but it is evident that some reliable techniques to generate nano-sized materials are still missing. A notable challenge involves generating nanoplastics with surface chemistries that closely emulate the characteristics of natural nanoplastics. While accelerated procedures are never on par with conditions found in nature,<sup>218</sup> they can still serve as better models than the ones currently used. Materials scientists can, moreover, think about introducing labels in the process that facilitate subsequent biological analysis. Introducing stable or radioisotopes is one option, but it is not readily available to all labs. Alternatives could be metal-doped nanoparticles.<sup>219,220</sup> However, whatever functionality is introduced, care should be taken not to significantly alter the surface chemistry, as this may lead to the adsorption of different proteins, thus different biodistributions.

### Developing techniques to identify nanoplastics

As Ivleva emphasized,<sup>75</sup> the analysis of nanoplastics within environmental matrices remains a significant analytical challenge. Conventional methods for microplastic detection, primarily based on diffraction techniques, lack the requisite resolution to identify nanoparticles at the nanometer scale. Moreover, visualizing low-abundance nanoplastics against complex backgrounds containing naturally occurring nanoparticles—including viruses, organic matter, and inorganic particles—poses additional difficulties. Preconcentration, enrichment, and fractionation procedures can enhance detection sensitivity; however, the advancement and integration of hyphenated analytical techniques, wherein two complementary methods are combined, are arguably the most promising strategies for the effective visualization and characterization of nanoplastics amidst diverse environmental constituents. Because nanoplastics are usually found in low concentrations in most environments, developing reliable analytical techniques that prevent false positives poses extra challenges. Once such samples are transferred to the lab for analysis, the samples are introduced to an environment that is filled with plastic products such as gloves, stoppers and syringes. There is an increasing awareness that analysis of samples that contain plastic nanoparticles requires additional care to avoid further contamination<sup>221</sup> but also more thought needs to be devoted to the development of control experiments that help verify detection thresholds and the absence of false positives.<sup>75</sup>

### Contributing to the discussion on standards

While materials scientists are not necessarily policymakers, their involvement in the discussion on standardized protocols

is crucial. This would include the discussion on the best way of separating micro- and nanoplastics from the surrounding media, such as natural inorganic nanoparticles and organic matter, *etc.*, and the establishment of protocols and procedures that have limited false readings and can be universally applied. The International Standardization Organisation has now established the first internationally recognized microplastic testing standards, ISO 24187, “Principles for the analysis of microplastics present in the environment.” This is a first step forward, but this standard focuses only on microplastics. Therefore, informed discussion towards clear guidelines on nanoplastics is needed.

### Analysing the interaction between nanoplastic with pollutants

While current research primarily focuses on assessing the toxicity of micro- and nanoplastics, predominantly utilizing engineered polystyrene (PS) beads, the potential risks may be more alarming. Traditionally, polymer scientists exploit the ability of plastics to interact with small molecules, such as additives, to enhance desired material properties. In natural environments, this mechanism inadvertently occurs as plastic particles either absorb pollutants into their bulk or adsorb contaminants onto their surfaces, potentially facilitating the transfer of harmful substances. Depending on the polymer and the pollutant, accumulation is not limited to the surface but these small molecules will penetrate deeper into the nanoplastic material. This can lead to unexpected toxicity. While some pollutants may not be toxic because they lack a pathway to enter cells in the body, nanoplastics can now transport them directly into cells, where they are released. It was discovered that a mixture of microplastics and perfluoroalkyl substances (PFAS) exhibits synergistic toxicity compared to PFAS and plastic particles alone.<sup>222</sup> Similar effects were observed with heavy metals.<sup>223</sup> This interaction depends not only on the compatibility between the polymer and the compound but also on the presence of functional groups, which typically result from the degradation process. Additionally, amorphous polymers can absorb significant amounts of low molecular weight compounds if the Flory–Huggings interaction parameter  $\chi$  is favorable. In contrast, crystalline polymers generally have limited space to accommodate potential pollutants. It is important to remember that loading pollutants in bulk or on the surface will alter the properties of nanoplastic particles, thus affecting colloidal stability, protein adsorption, and ultimately determining the fate of these nanoplastics in the body. This relationship between the size of the plastic particles, the degradation mechanism that determines functionality of the surface, the type of pollutants, and the resulting potentially toxic warheads is complex, and further work is required to fully understand how plastic will increase the toxicity of environmental pollutants.

### Developing strategies to remove plastic from complex media

There are several proposed strategies for detoxifying the body from plastic, including the use of melatonin, probiotics, and other compounds that appear to mitigate the effects of plastic



in animals.<sup>212</sup> Very recently, extracorporeal apheresis, which is the filtration of the blood, was tested as a way to remove microplastics from the blood,<sup>224</sup> although approaches like this need to be followed up with more studies, including detailed quantification and size analysis. Most of these studies are preliminary and require more attention from medical scientists.

Materials scientists can, however, help to create better techniques to remove micro- and nanoplastics from the environment as part of a remediation process. Currently, a range of techniques are already applied to remove micro- and nanoplastics from wastewater, such as filtration, centrifugation, and flocculation, but not all of them are suitable for nanoplastics.<sup>225</sup> New ideas are currently being explored to remove nanoplastics from the environment, such as the electrophoretic removal of nanoplastics from industrial wastewater,<sup>226,227</sup> the photolytic decomposition of plastic,<sup>225</sup> the use of bioreactors,<sup>225</sup> or the adsorption of nanoplastics onto other materials.<sup>225</sup> Most of these techniques appear to be successful in removing nanoplastics from water, but it is evident that other matrices, such as soil, are more challenging. Isolating nanoplastics from complex media, however, is an important step in enabling robust quantitative analysis.

### Developing an understanding that polyethylene is not polyethylene

Just because two plastic items are made from polyethylene, it does not mean they are the same; this statement applies to any plastic items. There are differences in additive, molecular weight, crystallinity, branching, and so on that result in different behaviours in the environment, such as different degrees of degradation.<sup>228</sup> This can lead to a large amount of data as each polymer has its unique degradation rate in a specific environment. Machine Learning could be applied more widely in the future to help identify the link between the type of polymer and its degradation.<sup>229</sup> The subtle differences between polymer structure and outcome are not only evident when analysing the behaviour of the nanoplastic in the environment. Very recently, it was shown that even Py-GC-MS, a standard technique to identify plastics, could not provide reliable results when PS of different tacticities (syndiotactic, isotactic and atactic) were measured.<sup>230</sup> This means that we need to adjust some analytical techniques and rethink its robustness.

### Participation in networks

This complex topic can only be understood through collaboration among researchers from different fields. Chemists and materials scientists contribute their expertise on analyzing micro- and nanoplastics and designing reference materials, but their role extends further. Large-scale comparison studies across multiple labs help validate methods and ensure their accuracy. These interlaboratory studies are already underway or have been completed, such as a study involving 84 labs that compared microplastic detection methods.<sup>231</sup> Other networks explored better reporting options and open-access tools.<sup>232</sup> A network can help establish key principles for managing plastics, which can become government-supported standards

such as the Australian Standard AS ISO 24187 “Principles for the analysis of microplastics present in the environment”.<sup>233</sup> This entails academics dedicating some of their time and resources, but it is a crucial step toward addressing this issue.

### Creating alternative polymers for single-use plastic

While the mantra should be to avoid single-use plastic at all costs and for everyone to take on the responsibility to limit daily plastic usage, there are still fields where plastic is necessary. An example would be the need for sterile catheters, blister packs for medicine, packaging that can enhance the shelf-life of food, or personal protective equipment. Biodegradable plastics could be an alternative as they are designed to degrade fully. However, this approach has caveats. Comparison of biodegradable microplastics with that of commodity polymers shows that degradable materials pose similar challenges such as the adsorption of contaminants, but they can also display negative effects towards marine life<sup>234</sup> or modulate the carbon and nitrogen cycle in marine sediment.<sup>235</sup> A recent review warned that we should not replace one problem with another, as there is evidence that incompletely degraded plastics might affect the ecosystem.<sup>236</sup> While polymer scientists have many innovative ideas regarding the design of new degradable polymers, their concepts need to be discussed with ecotoxicologists in a timely manner.

## Conclusions

There is no denying that excessive plastic consumption, especially single-use plastics, has created a significant environmental problem. Now, increasing evidence shows that tiny plastic particles enter humans and are detected in blood and organs, including the brain. The plastic particles found in the brain measure 200 nm in size; therefore, we need to expand our discussion from microplastics to nanoplastics. Nanoplastics are produced in similar ways to microplastics. However, when discussing nanoplastics, new aspects need to be considered, such as the crystallinity of the polymer, which affects degradation. The results of degradation are often fiber-like pieces, irregular with rugged edges. Sometimes, they resemble film pieces, but they are rarely spherical particles. When transitioning from microplastic to nanoplastic, other effects come into play, such as the large surface area. The type of functional groups on the surface, usually resulting from the degradation pathway, will influence colloidal stability and the adsorption of biomolecules. This determines whether the nanoplastic particles are nano-sized entities or large micron-sized aggregates in an aqueous environment. How these nanoplastic particles reach the brain can be understood through the field of nanomedicine. From there, it becomes clear that size, shape, and surface functionality— which influence the protein corona composition— are highly important parameters that shape the behavior of nanoparticles. This also highlights that published toxicity studies with round engineered nanoparticles may not indicate actual toxicity effects, as we know that



non-spherical structures can behave quite differently. Materials scientists can help create better reference materials, but they can also develop reference materials that can be easily detected in tissue and complex biological solutions, as these small nanoparticles are harder to identify than microplastics. While we often discuss microplastics, it appears that we need to shine a spotlight on nanoplastics. This complex problem can only be tackled when materials scientists, analytical chemists, environmental scientists, medical researchers, and researchers from related fields work together.

## Author contributions

The article has been written by MS with no further support.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

This is a review article and does not contain any original data.

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