



12 **Abstract**

13 Nanoplastics have attracted increasing attention in recent years due to their  
14 widespread existence in the environment and the potential adverse effects on living  
15 organisms. In this paper, the toxic effects of nanoplastics on organisms were  
16 systematically reviewed. The translocation and absorption of nanoplastics, as well as  
17 the release of additives and contaminants adsorbed on nanoplastics in the organism  
18 body were discussed, and the potential adverse effects of nanoplastics on human  
19 health were evaluated. **Nanoplastics can be ingested by organisms, be accumulated in  
20 their body and be transferred along the food chains.** Nanoplastics showed effects on  
21 the growth, development and reproduction of organisms, and disturbing the normal  
22 metabolism. The toxic effects on living organisms mainly depended on the surface  
23 chemical properties and the particle size of nanoplastics. Positively charged  
24 nanoplastics showed more significant effects on the normal physiological activity of  
25 cells than negatively charged nanoplastics, and smaller particle sized nanoplastics  
26 could more easily penetrate the cell membranes, hence, accumulated in tissues and  
27 cells. Additionally, the release of additives and contaminants adsorbed on nanoplastics  
28 in organism body poses more significant threats to organisms than nanoplastics  
29 themselves. However, there are still knowledge gaps in the determination and  
30 quantification of nanoplastics, as well as their contaminant release mechanisms,  
31 degradation rates and process from large plastics to nanoplastics, and the  
32 transportation of nanoplastics along food chains. These challenges would hinder the  
33 risk assessment of nanoplastics in the environment. It is necessary to further develop

34 the risk assessment of nanoplastics and deeply investigate its toxicological effects.

35 **Keywords:** Nanoplastics; Organisms; Humans; Toxic effects; Toxicity assessment

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## 37 1. Introduction

38 Nowadays, the production and application of plastics are increasing. In 2015, the  
39 global amount of plastic production has reached 322 million tons (PlasticsEurope,  
40 2016). Due to the chemical stability, persistence, bioaccumulation of plastics, the  
41 problem of plastic pollution in the environment is becoming increasingly prominent  
42 (Barnes et al., 2009; Fossi et al., 2016; Hu et al., 2019; Shen et al., 2019). Up to now,  
43 plastics wastes have been detected in soils (Zhang and Liu, 2018), lakes (Xiong et al.,  
44 2018b), sediments (Abidli et al., 2018; Rodrigues et al., 2018; Wen et al., 2018),  
45 oceans (Keswani et al., 2016; Mendoza et al., 2018), and even in places with rare  
46 human activities, such as Antarctic (Munari et al., 2017) and Arctic (Cózar et al., 2017;  
47 Peeken et al., 2018). Plastic wastes can be degraded into small fragments or particles  
48 due to weathering, sunlight radiation and biodegradation. The plastic species with the  
49 particle sizes less than 5 mm were defined as microplastics (Thompson et al., 2004).  
50 Cosmetics and detergents that contains small plastic particles are the major sources of  
51 microplastics (Dris and Cours, 2016; Lei et al., 2017). The characteristics of small  
52 particle size and low density enable microplastics to escape the traditional wastewater  
53 treatment process and enter into the marine systems (Carr et al., 2016; Murphy et al.,  
54 2016). Microplastics can be further degraded to nanoplastics, which have the particle  
55 size between 1 nm and 100 nm (Jambeck et al., 2015). The commercial application of  
56 nanoplastics is also an important source of nanoplastics in the environment.

57 Compared with other nanomaterials, such as carbon nanomaterials (Gong et al.,  
58 2009; He et al., 2018b; Xiong et al., 2018a; Yi et al., 2018) and metal nanomaterials

59 (He et al., 2018a; Qin et al., 2018; Xu et al., 2012; Yang et al., 2018), the research of  
60 nanoplastics is still in its infancy. Due to the limitation of technology, the degradation  
61 processes and degradation rates of microplastics to nanoplastics are not yet clear. But  
62 it is certain that the decomposition processes will not stop and the microplastics will  
63 continue to form nanoplastics (Mattsson et al., 2015). It is a long-term and lasting  
64 process in the environment. The amount of nanoplastics in the environment is not yet  
65 known, because there are no effective methods for the determination, quantification  
66 and assessment. Due to the considerably small particle size, nanoplastics are widely  
67 distributed in the aquatic environment and can be easily ingested by organisms. Lu et  
68 al showed that polystyrene (PS) nanoplastics (200 nm) could be ingested and  
69 accumulated in the body of zebrafish (*Danio rerio*) (Lu et al., 2016). The authors  
70 further found that PS nanoplastics with a concentration of 20  $\mu\text{g L}^{-1}$  ( $1.1 \times 10^8$   
71 particles/mL) could cause local necrosis and lipid aggregation in the liver, resulting  
72 in metabolic disorders and energy cycling disruption. Kashiwada et al studied the  
73 uptake and accumulation of PS nanoplastics in medaka (*Oryzias latipes*) (Kashiwada,  
74 2006), and found that PS nanoplastics (39.4 nm, 10  $\text{mg L}^{-1}$ ) were mainly distributed  
75 in gills and viscera, but also in testis, liver and blood. More seriously, PS nanoplastics  
76 could pass through the blood-brain barrier, a highly selective barrier that can prevent  
77 the entry of potential neurotoxins via an active transport mechanism mediated by  
78 P-glycoprotein, and enter into the brain tissues, posing profound effects on organisms.  
79 On the other hand, nanoplastics could be transported along the food chain, eventually  
80 accumulated in the higher trophic levels, such as fish, birds, and even human beings.

81 Cedervall et al showed that **PS nanoplastics (28 nm) at a concentration of 0.01% (w/v)**  
82 could transfer along the food chain from algae to zooplankton and fish, and the  
83 feeding behavior and lipid metabolism of fish were noticeably affected (Cedervall et  
84 al., 2012). Nanoplastic particles can adsorb chemical substances from water (**heavy**  
85 **metals**, polycyclic aromatic hydrocarbons, polychlorinated biphenyls, etc.), which  
86 may aggravate the adverse effects of nanoplastics on living organisms (Koelmans et  
87 al., 2016). Once these nanoplastic-contaminant complexes were ingested,  
88 contaminants can be simultaneously absorbed by organisms. Plastic additives and  
89 polymer monomers, such as bisphenol A, anti-ultraviolet radiation stabilizers,  
90 polybrominated diphenyl ethers and phthalates, can be released during degradation  
91 process. Many of these additives and polymer monomers are toxic to organisms, and  
92 can bring about acute poisoning symptoms, endocrine disorders, and reproduction  
93 toxicity (Bejgarn et al., 2015). Consequently, nanoplastic contamination should be  
94 seriously considered, and the ecotoxicity and environmental risks of nanoplastics need  
95 to be well studied.

96 Although some studies have been carried out on the toxicity of nanoplastics,  
97 further efforts are still needed. For example, to date, toxicological studies on  
98 nanoplastics mostly focused on model organisms, such as zebrafish and earthworm,  
99 but few experimental studies on toxicity of nanoplastics in mammals were carried out.  
100 Although nanoplastics could enter the **cell membranes** and blood circulation of  
101 **organisms**, the metabolic system of **lower trophic levels is different from that of**  
102 human beings and the ability of clearance and resistance to nanoplastics is also

103 different. The release of toxic pollutants from nanoplastics was affected by many  
104 factors, so it was not certain whether or not toxic substances would be released in the  
105 human body. In this paper, in order to provide valuable reference for the biological  
106 safety assessment and potential toxicity studies of nanoplastics, sources and  
107 environment behaviors of nanoplastics, toxic effects of nanoplastics on organisms,  
108 toxic effects of pollutants brought about by nanoplastics, and potential human health  
109 risks are comprehensively reviewed, and some future research needs are proposed.

## 110 **2. Sources and environmental behaviors of nanoplastics**

111 The sources of microplastics in marine environment were mainly from  
112 land-based input, aquatic aquaculture and fishing, and coastal tourism (Browne et al.,  
113 2011). It has been reported that about 800 million tons of plastic wastes in the ocean  
114 from the land (Jambeck et al., 2015). Due to small particle size, traditional wastewater  
115 treatment processes could not thoroughly remove such plastic particles, which cause  
116 huge amount of **microplastic particles entering the marine environment** (Vance et al.,  
117 2015). Loss or erosion of soil polluted by microplastics is another way of the  
118 land-based source of microplastics to the marine environment (Horton et al., 2017). In  
119 addition, shipping industry and coastal tourism cause the release of a large number of  
120 plastic products and wastes into the beach and ocean, which is an important reason for  
121 the increase of marine plastic pollution. According to the estimation issued by the  
122 United Nations Environment Program, the amount of waste plastics was about 275  
123 million tons in 2010, and approximately 4.8 – 12.7 million tons entered the marine  
124 system (Mattsson et al., 2015).

125 Compared with terrestrial ecosystem, plastic wastes are more easily decomposed  
126 into smaller plastic fragments because of the effects of high salinity and  
127 microorganisms in the marine environment (Sudhakar et al., 2007; Watters et al.,  
128 2010). The mechanism of degradation can be divided into two pathways:  
129 non-biodegradation and biodegradation (Figure 1). Non-biodegradation of plastics  
130 mainly includes thermal degradation, physical degradation, photodegradation,  
131 thermos-oxidative degradation and hydrolysis (Andrady, 2011). Thermal degradation  
132 of plastics is a commercial degradation process and do not occur in the environment.  
133 Physical degradation is an important process of large plastic wastes into fragments,  
134 and this process is mainly controlled by weathering and sea waves. Hydrolysis, a  
135 bond breaking reaction, is an efficient process contributing to the degradation of  
136 plastics in aquatic environment, changing the high polymer to low polymer.  
137 Photodegradation is a very efficient non-biodegradation method of plastic degradation  
138 in the environment. These types of plastic degradation would decompose the structure,  
139 change the mechanical properties of high polymers, and increase the effective surface  
140 area of plastics during their physical-chemical reactions and their interactions with  
141 microorganisms (Nathalie et al., 2008). Biodegradation is another important pathway  
142 that further degrades plastic fragments into nanoplastics in the environment (Trishul  
143 and Mukesh, 2010). During the biodegradation, plastics are often degraded outside the  
144 bacteria. Extracellular enzymes excreted by living microorganisms can breakdown the  
145 polymer chains. This process produces smaller plastic particles of different structure,  
146 eventually forming nanoplastics. Once reaching to the nanoscale level, the specific

147 surface area will extensively increase. For example, the specific surface area of a  
148 common plastic bag (0.2 m<sup>2</sup>) would become of 2600 m<sup>2</sup> after being totally  
149 decomposed into nanoplastic particles with the average diameters of 40 nm (Mattsson  
150 et al., 2015). With the extensive plastic wastes entering the marine environment, large  
151 amount of nanoplastics would be generated.

152 Furthermore, nanoplastics can act as carriers of chemical and biological  
153 contaminants (Hodson et al., 2017; Koelmans et al., 2016; Lagana et al., 2019).

154 Nanoplastics can adsorb chemical contaminants due to their low polarity and high  
155 roughness surface characters. The accumulation of chemical contaminants on the  
156 surface of nanoplastics increased their potential of transport and uptake by organisms,  
157 because of the ubiquitous existence of nanoplastics (Pittura et al., 2018).

158 Contaminants can enter the food chain via ingestion of these nanoplastics by  
159 organisms, thereby increasing the bioavailability of chemical contaminants to the  
160 organisms. Nanoplastics can also adsorb harmful algae, bacteria and viruses. The  
161 aggregation of these harmful species may induce gene exchange among different  
162 species and promote the diffusion ability of drug resistant bacteria and pathogenic

163 bacteria, which may aggravate toxicological and pathological responses to organisms.

164 New bacteria may be produced during the gene exchange. In particular, pathogenic  
165 and antibiotic resistance bacteria contain abundant pathogenic and antibiotic  
166 resistance genes, which may be transferred by multi pathways between communities  
167 on biofilms (Arias-Andres et al., 2018; Parthasarathy et al., 2019; Schmidt et al.,  
168 2014). The presence of nanoplastics may enhance the migration of antibiotic

169 resistance genes because of their strong floatability and mobility. Antibiotic resistance  
170 genes may be transferred to different areas. The exchange of antibiotic resistance  
171 genes between communities or the surrounding environment and the presence of  
172 antibiotic resistance bacteria may result in huge uncontrollable disasters.

### 173 3. Toxic effects of nanoplastics on organisms

174 Research on the impacts of nanoplastics to organisms showed that nanoplastics  
175 could be ingested and accumulated by various organisms. They can enter the  
176 circulatory system through enteric tissues, causing toxic effects at cellular and  
177 molecular levels. The adverse effects of nanoplastics on organisms are closely  
178 associated with the particle size, composition, morphology, aging time and surface  
179 properties. Some examples of toxicity evaluations of nanoplastics to organisms are  
180 shown in **Table 1**.

181 Bhattacharya et al showed that charged PS nanoplastics (20 nm) can be easily  
182 adsorbed on the surface of green algae (2 – 10  $\mu\text{m}$ ) and affect its photosynthesis  
183 process (Bhattacharya et al., 2014). The authors further showed that the absorption  
184 capacity of nanoplastics on the surface of these algae by scallops was greatly  
185 enhanced. Another research performed by Della et al reported that charged PS  
186 nanoplastic particles (40 nm) were easily accumulated in the digestive tract of sea  
187 urchin; as compared with negatively charged particles (PS-COOH), positively  
188 charged particles (PS-NH<sub>2</sub>) showed more obvious toxicity to sea urchin (Della et al.,  
189 2014). PS-NH<sub>2</sub> nanoplastics could bind to lipid bilayers on the cell membrane with  
190 high affinity, which promotes the cell absorption through endocytosis. Canesi et al

191 suggested that PS-NH<sub>2</sub> could decrease lysosomal membrane stabilization,  
192 significantly increase oxyradical production in hemolymph serum, and induce rapid  
193 cellular damage such as membrane blebbing and loss of filopodia (Canesi et al., 2016).  
194 The generation of PS-NH<sub>2</sub> protein corona in hemolymph serum was observed with the  
195 existence of PS-NH<sub>2</sub>. Micro-scale aggregates of PS-COOH nanoplastics occurred in  
196 the media, which may efficiently reduce the bioavailability of PS-COOH nanoplastics  
197 and decrease its toxicity. Actually, PS-NH<sub>2</sub> nanoplastics could exist as nanoparticles  
198 in the environment for a long time, which greatly increased their bioavailability,  
199 exposure risk, and the ability of penetrating cells and tissues.

200 Composition and particle size of nanoplastic polymers would also have  
201 significant effects on the toxicity of nanoplastics to organisms. Smaller nanoplastics  
202 may more easily enter the body tissues and cells. The remove of these nanoplastic  
203 particles is even more difficult, thereby increasing their exposure period and risks to  
204 organisms. The nanoplastic toxic effects on various test subjects, including different  
205 polymers used in experiments and different toxic endpoints, were summarized based  
206 on previously published literature (Figure 2). Compared with other nanoplastics, PS  
207 and polymethylmethacrylate (PMMA) nanoplastics were more commonly used. This  
208 may be due to the difficulty in the synthesis of other polymers. Ward et al reported  
209 that mussels (*Mytilus edulis*) could directly absorb PS nanoplastic particles (30 and  
210 100 nm) through the intestine (Ward and Kach, 2009). Evidence showed that  
211 bioaccumulation of PMMA nanoplastics occurred in barnacles at even low  
212 concentrations (1 ppm) in chronic exposure tests (Bhargava et al., 2018). PMMA

213 nanoplastics could persist in the body throughout the whole stage of growth and  
214 development from Nauplius to juvenile barnacle, posing a potential long-term threat  
215 on invertebrate communities. The abundance of mRNA transcript was obviously  
216 increased and the immune system might be impaired after exposure to PMMA  
217 nanoplastics. Nanoplastics could change molecular signaling pathway and potentially  
218 interfere with the metabolism of lipids.

219 The coexistence of nanoplastics and other nanoparticles or chemical substances  
220 would significantly affect their toxic effects on organisms. Kong et al studied the  
221 combined toxicity of PS nanoplastics and titanium dioxide (TiO<sub>2</sub>) nanoparticles on  
222 nematode (*Caenorhabditis elegans*) (Dong et al., 2018). The authors found that  
223 co-exposure to PS and TiO<sub>2</sub> nanoparticles changed the molecular basis of oxidative  
224 stress. The presence of PS nanoplastics further enhanced the toxicity of TiO<sub>2</sub>  
225 nanoparticle via inducing intestinal reactive oxygen species production and  
226 decreasing locomotion behaviors in *sod-3* mutant nematodes. Another study showed  
227 that mussel (*Mytilus galloprovincialis*) was more sensitive to PS nanoplastics  
228 exposure even at low concentrations (0.05 mg L<sup>-1</sup>) (Brandts et al., 2018). PS  
229 nanoplastics changed the expression of gene, decreased the enzymatic activity,  
230 induced effects on neurotransmission, increased the oxidative status and finally  
231 resulted in peroxidative damage (Brandts et al., 2018). Chen et al reported that both  
232 bisphenol A and PS nanoplastics (50 nm) could result in myelin basic protein or gene  
233 up-regulation in the central nervous system, and significantly inhibit  
234 acetylcholinesterase activity of zebrafish (*Danio rerio*) (Chen et al., 2017). The

235 existence of **PS nanoplastics** obviously enhanced neurotoxic effects in central nervous  
236 system and dopaminergic system. Besides, **PS nanoplastics could be transferred** along  
237 the food chain (algae–zooplankton–fish), the lipid metabolism and behavior of  
238 crucian were significantly affected (Cedervall et al., 2012).

239 At present, due to the small particle size of nanoplastics, the detection methods  
240 of nanoplastics in biological tissues are few, and there are relatively few studies on the  
241 composition, distribution of nanoplastics and their impact on the environment and  
242 organisms. **Although the growth, development and reproductive toxicity of living**  
243 **organisms by nanoplastics has been revealed, huge concentrations of nanoplastics**  
244 **were used to test the effects within short periods (Table 1), and these concentrations**  
245 **cannot occur in the environment. More tests are needed to examine the impacts of**  
246 **lower dosage or environmental dosage and long-term exposure to nanoplastics.**  
247 **Moreover, to date, most researches on toxicity and risk assessments of nanoplastics**  
248 **were performed using PS materials. We expect that more studies are needed using**  
249 **other nanoplastic materials such as polypropylene (PP), polyethylene (PE), polyvinyl**  
250 **chloride (PVC), polyethylene terephthalate (PET), etc. Many conclusions and**  
251 phenomena of nanoplastics in organisms need to be further explored.

#### 252 **4. Toxic effects of pollutants brought about by nanoplastics**

##### 253 *4.1 Additives*

254 Some chemical compounds are often used in the production of plastics, such as  
255 anti-ultraviolet radiation stabilizers, phthalates and bisphenol A (Hirai et al., 2011). It  
256 is worth mentioning that plastic polymer monomers, such as styrene monomers, and

257 phthalates can be easily released from **plastic matrices** during decomposition and  
258 degradation. Evidence showed that these additives and polymer monomers have  
259 estrogenic or anti-androgenic (Fries et al., 2013), reduction of reproductive rate and  
260 endocrine disrupting effects on organisms (Iguchi et al., 2006). The release of  
261 additives from microplastics or nanoplastics was affected by many factors, such as  
262 ionic strength of leaching solution, **aging time**, surface roughness, polymer type, etc.  
263 The mortality, growth inhibition and abnormal embryo of organisms caused by the  
264 exposure to micro(nano)plastic leaching solution have been revealed. We summarized  
265 recent toxicity tests of plastic leachates of different polymer types. According to these  
266 studies (as seen in **Table 2**), manufacturing process, polymer type, aging time and  
267 particle size had obvious influences on the toxicity of leaching solution. However,  
268 there are few data on the toxicity of leaching solution to nanoplastics. **Further**  
269 **researches on release mechanism of polymer monomer and additives from**  
270 **nanoplastics and toxicity exposure of leaching solution are needed.**

#### 271 4.2 Attached contaminants

272 Due to large specific surface area and inherent hydrophobicity, persistent organic  
273 pollutants (such as polychlorinated biphenyls (PCBs) (Ren et al., 2018b; Velzeboer et  
274 al., 2014) and polybrominated diphenyl ethers (PBDEs) (Ren et al., 2018a; Ye et al.,  
275 2017a; Ye et al., 2017b)) and heavy metals (Tang et al., 2018) can be easily adsorbed  
276 on the surface of plastic particles. Mato et al reported that **concentrations of PCBs and**  
277 **dichloro-diphenyl-trichloroethane (DDT)** in plastic particles were significant higher  
278 than in surrounding sea waters, demonstrating strong affinity of plastics for these

279 chemicals (Mato et al., 2001). Polymer type, particle size and surface structure of  
280 nanoplastics are important factors affecting the surface bound contaminants (Guo et  
281 al., 2012). Nowadays, few studies are conducted on the adsorption of pollutants by  
282 nanoplastics, however, researches on the adsorption of pollutants by microplastics  
283 have been carried out.

284 Laboratory studies generally suggested that pollutants adsorbed on the surface of  
285 micro(nano)plastics can be transported and accumulated into organisms via feeding  
286 behaviors. Nevertheless, there is no clear conclusion of whether a linear relationship  
287 exists between the concentrations of micro(nano)plastics and the residual  
288 concentrations of pollutants. For instance, Browne et al showed that the contents of  
289 pollutants in the abdominal wall and intestine of *Arenicola marina* increased  
290 significantly after being fed with microplastics and gravel (Browne et al., 2013). The  
291 authors further showed that there was no clear judgment whether pollutants were  
292 adsorbed on gravel or onto microplastics. Besseling et al reported that the relationship  
293 between concentrations of microplastics and content of ingested pollutants was  
294 complex in *Arenicola marina* after being exposed to soils containing microplastics  
295 and PCBs (Besseling et al., 2013). The study found that when the concentration of  
296 microplastics was low, the accumulation of PCBs in the organisms increased  
297 obviously; however, when the concentration of microplastics rose to a higher level,  
298 the aggregation of PCBs slightly decreased. Another study also reported that no  
299 significant influence on the concentration of PBDEs in earthworms (*Eisenia fetida*) by  
300 the addition of microplastics was observed (Gaylor et al., 2013). Chua et al showed

301 that microplastics and nanoplastics could serve as vectors for pollutants when entering  
302 marine organisms, while the transport and absorption efficiency of pollutants was  
303 related to their inherent properties (Chua et al., 2014).

304 On the other hand, some simulation experiments showed that the migration  
305 ability of pollutants on the surface of microplastics was limited (Gouin et al., 2011;  
306 Koelmans et al., 2013). Gouin et al studied the effects of different temperature, pH  
307 and intestinal surfactants on the accumulation of pollutants on the surface of  
308 microplastics using a single storehouse model (Gouin et al., 2011). The results showed  
309 that microplastics had little effect on the intake of pollutants via the organism  
310 intestinal. Meanwhile, according to the research on thermodynamic food web model,  
311 pollutants on the surface of microplastics also had less contribution to the total  
312 amount of transfer and accumulation of pollutants in organisms. Another simulation  
313 study on North Sea cod (*Gadus morhua*) also suggested that microplastics had less  
314 effect on the aggregation of pollutants in organisms (Koelmans et al., 2013). However,  
315 some complicated factors were not taken into account in these models, such as the  
316 accumulation of pollutants in lipids and the existence of absorption and distribution  
317 dynamics. These factors would lead to the gradual dissociation of pollutants from  
318 microplastics and consequently the absorption and accumulation of pollutants in the  
319 intestinal tract of organisms. Therefore, more reasonable and explicit models should  
320 be established in further researches to simulate the release and accumulation of  
321 pollutants attached on microplastics in organisms.

## 322 **5. Potential human health risks of nanoplastics**

323 *5.1 Translocation and absorption in the body*

324 Evidences showed that these tiny particles can be transferred along the food  
325 chain to higher trophic level organisms, or into the human food chain through other  
326 pathways (such as via sea salt or animal feed) (Yang et al., 2015). It is of great value  
327 to investigate and evaluate the transportation, absorption and toxic effects of  
328 nanoplastics in the human body. More recently, the studies on the toxic effects of  
329 nanoplastics were mainly focused on their transportation and absorption efficiency in  
330 the intestine, and their accumulation in tissues of various model animals. We here  
331 reviewed the endocytosis mechanism and toxicity evaluation of micro(nano)plastics  
332 in various animals and in vitro models (Table 3).

333 Whether nanoplastics can break the intestinal barrier and enter other parts after  
334 being ingested is an important basis for studying whether nanoplastics can be  
335 accumulated in organisms. This would be an important starting point for analyzing  
336 and assessing the toxic effects of nanoplastics. For example, in vivo and in vitro tests  
337 are important methods to investigate the endocytosis and absorption mechanisms of  
338 nanoplastics. Studies on a series of different types of nanoplastics suggested that they  
339 can cross the intestinal barrier into the circulatory system and eventually lead to  
340 systemic exposure (Bouwmeester et al., 2015). Magri et al simulated that the  
341 interaction between PET nanoplastics and cells using in vitro Transwell model of the  
342 intestinal epithelium (Magri et al., 2018). The authors reported that PET nanoplastics  
343 (26.7 nm) showed small size and long-term stability in various biological media,  
344 which increase the possibility of living organism exposure. No obvious toxic effects

345 were observed during the evaluation of the biological impact of PET nanoplastics on  
346 ingestion on human intestinal cells, but they can efficiently pass through an in vitro  
347 Transwell model of the intestinal epithelium. The absorption and transportation in  
348 vivo of nanoplastics was depended on their own structure and properties, such as  
349 chemical composition and surface modification. The oral bioavailability of 50 nm PS  
350 nanoplastics was in the range of 0.2% – 7% (Jani et al., 2011). However, some studies  
351 reported that the bioavailability of 60 nm PS nanoplastics was relatively high, ranging  
352 from 1.5 % to 10% (Hillery et al., 2008; Mishra et al., 2018). The possible reasons for  
353 such variation may be related to the aging time and surface modification of  
354 nanoplastics. Due to the difference of surface modification, the bioavailability of  
355 different sized nanoplastics (50 – 500 nm) showed a significant difference (0.2% –  
356 10%) (Kulkarni and Feng, 2013; Walczak et al., 2015a; Walczak et al., 2015b). Due to  
357 their large surface area and complex surface structure, nanoparticles can interact with  
358 various molecules, such as lipid, protein, water, ions etc. For example, the interaction  
359 of proteins and nanoparticles can lead to the generation of coronal protein rings  
360 (Huang et al., 2018; Lundqvist et al., 2008). These protein rings show a significant  
361 effect on the endocytosis of nanoparticles of cells. Coronal protein rings on the  
362 surface of nanoplastics would have similar chemical effects on the endocytosis and  
363 can promote the translocation efficiency of 50 nm nanoplastics during digestion. Also,  
364 the interaction of nanoplastics and iron ions could promote the uptake of irons  
365 (Mahler et al., 2012), the presence of nanoplastics would affect the transporting  
366 function of cell membrane to induce toxic effects at cellular and molecular level.

367 5.2 Potential toxic effects on human health

368 Generally, smaller nanoplastics are easier to enter and accumulate into tissues  
369 and cell organisms, the positively charged nanoplastics showed a significant impact  
370 on cell physiological activities. Xia et al found 60 nm cationic PS nanoplastics had  
371 being significantly toxic to macrophages and epithelial cells (Xia et al., 2008). Forte  
372 et al studied the role of PS nanoplastic particle sizes on its toxicity effects (Forte et al.,  
373 2016). The authors suggested that, as compared with 100 nm nanoplastics, 44 nm  
374 nanoplastics could enter gastric cancer cells more quickly and efficiently, and further  
375 affect the cell morphology, the expression of genes and cell proliferation by inducing  
376 up-regulation of some gene expression levels. Bhattacharjee et al showed that  
377 different modification of nanoplastics' surface had different effects on cell membrane  
378 and cell oxidative stress (Bhattacharjee et al., 2014). Compared with anionic PS  
379 nanoplastics, cationic PS nanoplastics significantly increased the concentration of free  
380  $Ca^{2+}$  and the content of intracellular reactive oxygen species. Meanwhile,  
381 mitochondrial membrane potential and the content of adenosine triphosphate  
382 decreased. Another research done by Liu et al reported that 50 nm  $NH_2$ -PS  
383 nanoplastics could obviously destroy cell integrity, as well as the proliferation of  
384 cervical cancer cells and mouse embryonic fibroblasts (Liu et al., 2011). The  
385 reorganization of cytoskeleton and chromosomes during cell mitosis was also directly  
386 affected, including prolongation of division cycle and decrease in expression of  
387 cyclin.

388 Due to the stable properties and difficulty of degradation, nanoplastics can be

389 accumulated in tissues and cells easily, causing metabolic disorders and local  
390 inflammation. Especially in patients with intestinal diseases, changes of tissue  
391 permeability caused by inflammatory infection would significantly increase the  
392 transportation and absorption of nanoplastics, thereby further increasing the exposure  
393 risk. More recently, it is noteworthy that researchers and scientists have begun to be  
394 conscious of the potential impact of nanoplastics on human health. Researches on the  
395 direct and indirect effects of nanoplastics on organisms have been gradually  
396 preformed. However, test subjects of these studies are limited to model cells and  
397 organisms, the shape and composition of nanoplastics investigated are comparatively  
398 single. Consequently, further studies should focus on nanoplastic contamination in  
399 ordinary organisms or in the food chain to accurately and comprehensively assess the  
400 impact of nanoplastics on ecosystem and human health.

## 401 **6. Perspectives and challenges**

402 As plastic products are widely used and difficult to be degraded, plastic pollution  
403 will continue to affect the health of the ecological environment. The large-scale  
404 growth of plastic production causes the continuous increase in the number of plastic  
405 fragments in the environments, leading to the accumulation of nanoplastics in  
406 different media. Although researches on the effects of nanoplastics on wildlife are  
407 limited, the adverse effects of nanoplastics on organisms will surely damage the  
408 ecosystem, sometimes even affecting human health. In order to obtain more  
409 information on the toxicity consequence of nanoplastics, further studies should focus  
410 on:

411 a. Efficient methods for determination, quantification and assessment of  
412 nanoplastics in the environment.

413 b. Degradation rates from large plastic particles to nanoplastics and mechanisms  
414 of degradation process.

415 c. Transmission and toxicological effects of nanoplastics on different trophic  
416 layers in marine representative ecosystems.

417 d. Using environmental concentrations of nanoplastics to test.

418 *a. Determination of nanoplastics in the environment*

419 Detection and quantification techniques of nanoplastics in the environment are  
420 limited. This analytical challenge has bogged the obtainment of data on the  
421 occurrence, fate, assessment and ecotoxicology of nanoplastics. There are few  
422 methods for detection of nanoplastics in the environment. Due to their small particle  
423 size, most existing sampling methods for microplastics have artificially neglected the  
424 detection of nanoplastics, and these methods were not applicable to nanoplastics.  
425 Consequently, establishment of new technologies and approaches for nanoplastic  
426 detection and quantification in the environment are urgent. Separation of organic  
427 materials and nanoplastics in samples is necessary. Based on our knowledge, chemical  
428 solvents, such as strong acids, would affect organic materials and probably also the  
429 nanoplastics. Thus, enzyme separation could be a better option.

430 *b. The process from plastics to nanoplastics and degradation rates*

431 Due to technologies and methods limits, it is difficult to obtain nanoplastics from  
432 the environmental samples. Exploration on the degradation processes and rates of

433 large plastic particles being decomposed to nanoplastics is necessary and urgent.  
434 Laboratory experiments of **stable isotope labeling** to study the process are feasible.  
435 **For example, large plastic particles are exposed to natural conditions under**  
436 **experimental simulation where the temperature, pH, sunlight and other forces can be**  
437 **easily controlled.** Under controlled conditions, the separation of degraded plastics  
438 based on particle sized can be achieved, as well as the determination of the fragments  
439 content at nanoscale. This procedure can probably reflect the existence of plastics  
440 with different particle sizes in the natural environment, and estimate the formation  
441 amount of nanoscale size after degradation, thereby approximately assessing its  
442 ecological effects. Moreover, the degradation processes not only reduce particle size,  
443 but also change the chemical properties of the surface of nanoplastic particles.  
444 **However, the characterization techniques of nanoplastics have not been reported yet,**  
445 **new methods and technologies need to be developed and applied to the**  
446 **characterization of nanoplastics.**

447 *c. Transmission via food chain/web and toxicological effects*

448 More recently, evidences showed that nanoplastics not only have adverse effects  
449 on organisms, but also accumulate in organism bodies. However, most of researches  
450 on the accumulation and toxicity of nanoplastics were only at individual level of  
451 organisms. Studies on the mechanisms of transmission of nanoplastics via the food  
452 chain/web should be strengthened. Before that, two important factors need to be  
453 considered: origin and particle size. Nanoplastics in the environment can originate  
454 from the degradation of large waste plastics and the direct release of commercial

455 applications. The properties of nanoplastics from these two sources are quite different.  
456 In order to accurately evaluate the environment effects, nanoplastics used in tests need  
457 to be originated from the accordingly sources. The size of nanoplastics adopted in  
458 experiments should be set according to the size range of nanoplastics in real  
459 environment. Additionally, biological experiments should not be limited to single  
460 organisms or individual level, but along the food chains or webs. For instance,  
461 phytoplankton, zooplankton, protozoa, metazoan, fish or higher trophic levels need to  
462 be exposed to the same nanoplastics containing environments. Survival, development,  
463 reproduction, behavior changes and toxicity need to be characterized. It is important  
464 to describe the effect of exposure time, composition and size of nanoplastics on  
465 organisms. Although laboratory experiments can obtain data on the toxicity of  
466 nanoplastics to organisms by food chain, its extension to real environment conditions  
467 can be very complicated. Therefore, a reasonable way forward still needs to be explored.  
468 The low trophic level bears different metabolic system with human beings, their  
469 ability of clearance and resistance to nanoplastics is also varied. Data on exposure  
470 experiments of nanoplastics may not be suitable for human. Whether nanoplastics can  
471 enter the human body via the food chain and the trigger of adverse effects on human  
472 also needs further investigation.

473 *d. Using environmental concentrations of nanoplastics to test.*

474 **Facts proved that everyone was using huge concentrations of nanoplastics to test**  
475 **the effects within short periods (Table 1). These concentrations cannot occur in the**  
476 **environment at all. More tests are needed to examine the impacts of lower dosage or**

477 environmental dosage and long-term exposure to nanoplastics. Most researches on  
478 toxicity and risk assessments of nanoplastics were often performed using PS materials.  
479 This will lead to misunderstand of the toxicity of nanoplastics to the ecosystems.  
480 Therefore, the establishment of new technologies and approaches for nanoplastic  
481 detection in the environment is needed. The impact on organisms and human health  
482 should be further developed in conjunction with the current status of nanoplastic  
483 pollution in the environment.

484

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490

#### 491 **Declaration of interest**

492 The authors report no conflict of interest.

493 **Reference**

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