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**Perspectives on Palladium-Based Nanomaterials: Green
Synthesis, Ecotoxicity, and Risk Assessment**

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As nanomaterials develop, it is important to deliberate about environmental protection and sustainable development. Therefore, the comprehensively assess the ecological risk of rapidly developing nanomaterials are needed. This perspective analyzes and discuss the full life circle, ecotoxicity, and assessment methods to provide recommendation for the sustainable development of palladium-based nanomaterials and other similar or related nanomaterials.

**Perspectives on Palladium-Based Nanomaterials: Green Synthesis, Ecotoxicity,
and Risk Assessment**

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10 **Abstract:**

11 Palladium-based nanomaterials (Pd-NMs) have been mass produced and applied due
12 to their remarkable properties and higher earth abundance. This makes Pd-NMs come
13 into frequent contact with the environment, and enter ecological environment. The
14 comparative analysis of toxicological data has presented that Pd-NMs showed acute
15 or chronic toxicity of Pd-NMs in both in vitro and in vivo biological receptors model,
16 but only limited information has been provided on the possible environmental
17 migration and transformation or concentration distribution in environmental media of
18 Pd-NMs. Therefore, a perspective is needed to propose the existing data to provide
19 more professional and comprehensive assessments for their ecotoxicity and
20 sustainable development. This perspective describes the critical knowledge needed to
21 assess their ecological risks. We recommend focusing on the current and future
22 concentration and distribution of Pd-NMs in the environment, guiding the assessment
23 of the full cycle ecotoxicity of Pd-NMs, and strongly encouraging the quantitative
24 measurement of the concentration level of Pd-NMs in the real environment.

25 Introduction

26 In recent decades, nanoscience has made major breakthroughs and greatly
27 improved human's life.¹⁻³ As an important achievement of nanoscience, nanomaterials
28 are produced and used in many technologies and consumer products owing to their
29 inimitable properties.⁴⁻⁶ Among them, metal-based nanomaterials (M-NMs) have
30 attracted widespread attention owing to their widespread application prospects.⁷ In
31 particular, noble M-NMs are widely used in the fields of catalysis,⁸ environmental
32 remediation,^{9,10} sensors,^{11,12} nanomedicine,^{13,14} and so on.¹⁵⁻¹⁷ However, it is
33 challenging to meet sustainability from the synthesis of products to their applications
34 and disposal.¹⁸⁻²⁰

35 Palladium-based nanomaterials (Pd-NMs) are one of the most widely used noble
36 M-NMs. Pd-NMs offer opportunities for efficient catalyst materials due to their high
37 specific surface area, abundant active sites, and high catalytic activity.²¹⁻²³ Trend of
38 publishing activities highlights the continuous development of Pd-NMs and related
39 materials. Pd-NMs have valuable catalytic and optical properties, which provide
40 extensive opportunities for their chemical, medical, and environmental applications in
41 human activities.²³⁻²⁵ Growth in the use of Pd-NMs has been accompanied by
42 increasing exposure to the ecological environment. In contrast, toxicology research of
43 Pd-NMs and the related materials is far behind their applications, while their

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4 44 toxicological study has positive significance for sustainable development and
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9 46 With the rapid development of Pd-NMs and related and/or similar materials, a
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12 47 variety of different voices have emerged in the public. On the one hand, Pd-NMs are
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15 48 favored by engineers in the fields of material, chemical, and energy science because
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18 49 of their excellent characteristics (Fig. 1).^{24,26,27} On the other hand, with the extensive
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21 50 application of Pd-NMs, the increasing interaction between Pd-NMs and the ecological
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24 51 environment makes individuals in the field of environmental ecology and biomedicine
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27 52 worry that Pd-NMs may violate the principle of sustainable development and cause
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30 53 ecological harm (Fig. 1).^{28,29} To better protect the ecological environment, we need to
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33 54 control the usage amount of Pd-NMs and improve the techniques and methods for
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36 55 assessing their toxicity.

37 56 In 2015, Chen and Ostrom introduced in detail various synthesis methods of
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40 57 Pd-NMs, as well as their outstanding properties and wide application range.²⁴
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43 58 Pd-NMs have unique chemical and catalytic properties, and their synthesis and
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46 59 application are expected to see remarkable growth over the next decade. Especially in
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49 60 the biomedical field, the need for real-time care and monitoring will accelerate the
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52 61 emergence of them.³⁰⁻³² Pd-NMs are released into the environment or in contact with
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55 62 organisms, initially exist as nanoparticles (NPs), which may subsequently migrate,
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58 63 transform, and accumulate in the environment or in organisms, ultimately causing
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61 64 harm to organisms. Existing data lack the assessment of the biological activity of

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4 65 Pd-NMs, and toxicity is often neglected by folks in the design of new nanomaterials,
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7 66 while outstanding properties are regarded as the only selling point of a new material
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10 67 system. Therefore, Egorova and Ananikov called for specific measurements of special
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12 68 metal catalysts, including Pd NMs, and for the nature, toxicity, bioavailability, and
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15 69 possible exposure pathways to be taken into account in the development and
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18 70 application of these chemicals.³³

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20 71 Meanwhile, Leso and Iavicoli critically analyzed data from the existing literature
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23 72 on toxicological and occupational risk assessment of Pd-NMs, and pointed out the
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26 73 negative effects of these chemicals on ecosystem function for use in determining
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29 74 appropriate strategies to assess and manage occupational risk of them.³⁴ For gain a
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32 75 more comprehensive understanding of the toxicology of Pd-NMs and provide
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34 76 guidance for their sustainable development, the current perspective summarizes and
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37 77 analyzes the risks of Pd-NMs on the environment, and discuss in vitro and in vivo
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40 78 research problems related to Pd-NMs. This perspective aim to describe the
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43 79 ecotoxicological effects of Pd-NMs and propose that individuals should consider the
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46 80 potential environmental hazards of these nanomaterials while promoting them. The
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49 81 insights of this perspective apply not only to Pd-NMs but also to the other similar and
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52 82 related M-NMs. Excellent properties or toxicity cannot be used as the sole reason for
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55 83 promoting new nanomaterials or hindering their development without comprehensive
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58 84 and dedicated evaluation.
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85 Environmental Impact of Palladium-Based Nanomaterials

86 Exploration of Scalable Production Routes

87 Are there multiple synthesis routes and are some more sustainable than the others?

88 Multiple routes to prepare Pd-NMs (such as nanofilms, nanowires, nanocrystals,
89 nanotubes, nanospheres, nanorods, etc.) including physical synthesis methods (such as
90 sputtering, ion or electron beam deposition, and laser ablation),³⁵⁻³⁷ hydrothermal
91 methods (one-step synthesis),³⁸ electrochemical deposition,³⁹ chemical deposition,⁴⁰
92 and other methods (such as microemulsion⁴¹ and photochemically⁴² assisted
93 synthesis). The important environmental impact of these synthetic methods are the
94 need to use toxic reaction reagents and produce toxic by-products.

95 Compared to other reaction reagents, polyols have unparalleled advantages.⁴³ First,
96 alcohols have multifunctional properties, such as being a reducing agent, solvent, and
97 stabilizer for metal precursors. Secondly, polyols have inherent hydrogen bond
98 interaction and adjustable number of -OH groups, which give them elastic structure.
99 Meanwhile, due to the existence of hydrogen bond supramolecular structure, polyols
100 can prevent the agglomeration of NPs.⁴⁴ Third, the physicochemical properties of
101 different polyols are different, which can meet the different synthetic conditions for
102 the synthesis of Pd-NMs with different properties. To illustrate, Pd-NMs synthesized
103 by rapid reduction of ethylene glycol (EG) at relatively high temperature usually have
104 a cubic or rod-like single crystal structure.⁴⁵ However, the use of other polyols and/or

changing reaction conditions can affect the growth rates of different surfaces of Pd-NMs, thus affecting the shape of them.⁴⁶ Finally, noteworthy, polyols derived from environmentally friendly plant extracts are also outstanding substrates for the synthesis of Pd-NMs.⁴⁷⁻⁴⁹

To minimize secondary contamination in the preparation of Pd-NMs, the synthesis of Pd-NMs or other related novel nanomaterials in green media (such as plant extracts)⁵⁰⁻⁵² is beneficial to the sustainable development of nanomaterials. Green chemical substance in plant extract can be used as a reducing agent or stabilizer in the preparation process of Pd-NMs, which can simply and quickly reduce Pd metal ions to zero-valent Pd metal without agglomeration.⁵³⁻⁵⁵ Noteworthy, these phytochemicals not only greatly reduce the adverse environmental effects of the synthesis process, but also ensure high reaction rates and controllable yields.^{52,56} The method is based on the ability of phytochemical molecules to absorb, accumulate, transform and recycle metal ions, which has the characteristics of economy, sustainability and environmental protection.

In 2008, varma et al., one of the pioneers, reported the green synthesis of a large number of Pd-NMs at room temperature using coffee and tea extracts.⁵⁷ The main components of these extracts are caffeine and polyphenols, which can form complexes with metal ions in solution and act as reducing agents to reduce metal precursors to metals. Meanwhile, they can also be used as dispersants in the synthesis process to avoid the accumulation of metal NPs (MNPs). The synthesized Pd-NMs

were mostly spherical in shape, but varied in size, depending on the quality and source of the extract (Fig. 2). Subsequently, Li et al. and Philip et al. extracted polyols from *Cinnamomum camphora* leaf⁵⁸ and dried leaf powder of *Anacardium occidentale*⁵⁹ respectively for the synthesis of Pd-NMs. They found the polyol component binds to the metal complex and is used to reduce metal precursors, while the heterocyclic component stabilizes the reduced MNPs. Furthermore, they found the size of Pd NPs could be controlled by changing the concentration of Pd ions in the solution without the need for additional templates. Besides, *Cacumen Platycladi* leaf extract,⁶⁰ *Pulicaria Glutinosa* Extract,⁶¹ Fruit and *Aloe Vera* Juices,⁶² artichoke leaf extract,⁶³ *Catharanthus roseus* leaf extract,⁶⁴ and *Terminalia chebula* aqueous extract⁶⁵ have also been reported as a reaction medium for the simple and green synthesis of Pd-NMs.

Release and Transformation during Use

Pd-NMs are characterized by invaluable catalytic, mechanical and optical properties which may offer the opportunity for their application in human activities (Fig. 3), including electrochemical reactions,^{66,67} nanomedicine,^{68,69} fine chemistry,^{70,71} sensors,^{72,73} and especially in the treatment of environmental pollutants such as control harmful exhaust emissions of automobiles, eliminate indoor air pollutants and treat wastewater.⁷⁴⁻⁷⁸ Environmental concentration of Pd-NMs has prominently increased from these human activities. To illustrate, for catalytic applications, all types of Pd-based catalytic systems, even immobilized catalysts

(including homogeneous, heterogeneous, metal complexes, and supported Pd-based catalysts), have been demonstrated the inevitable leaching.^{23,79,80} Leaching can cause metallic substances and NPs to be released directly or indirectly into the environment. When the diffused Pd-NMs come into contact with organic or inorganic ligands and water components, chemical reactions occurring on the surface of Pd-NMs, resulting in morphological changes and the formation of core-shell Pd-NMs,⁸¹ which further promotes environmental migration of the Pd-NMs.

Especially from the transportation sources, with the increase of car use in densely populated areas, the increase of environmental concentration of Pd-NMs is documented.⁸²⁻⁸⁴ Pd-NMs used in automotive catalytic converters are discharged as particles in the exhaust gas and accumulate in the local soil.⁸⁵⁻⁸⁷ Subsequently, through various chemical processes (such as redox reactions and complexation reactions), their environmental mobility is enhanced, eventually leading to their interactions with a variety of organisms.^{88,89} The Pd-NMs that enter the organism diffuse across cell membranes, and may end up in various organs or throughout the body through lymph and blood circulation.^{90,91}

Accumulation and deposition of the nanomaterials cause damage to aquatic organisms in a water environment.^{92,93} Changes in the surface chemistry of NPs and the chemical properties of the aquatic environment can change the degree of aggregation and deposition of nanomaterials.^{94,95} Furthermore, combined effects of the environment, such as acid rain, could increase the solubility of Pd-NMs and

168 spread them across the ecosystem through runoff or atmospheric transport.⁹⁶ In
169 summary, Pd-NMs in the environment exhibited toxicity to plants, animals, and
170 microorganisms (Fig. 4), and environmental behavior of the Pd-NMs could aggravate
171 they ecological risk.⁹⁷

172 **Bioavailability and Toxicity in Organisms**

173 **Acute and Chronic Toxicity to Animals**

174 Pd-NMs have the highest bioavailability among platinum group metals and greater
175 fluidity than other platinum group metals.^{98,99} Pd-NMs have been proved to be
176 enriched in living organisms and their retention in the animal depends on how it is
177 administered.^{28,100} Pd-NMs enters animals mainly by inhalation and injection, and
178 exert cytotoxic and proinflammatory effects in vitro while affecting different target
179 organs in an animal model test. The mechanisms of toxicity of Pd-NMs in animal
180 models are mainly toxicity of released Pd ion and oxygen stress response, which
181 disrupts the energy metabolism balance, inhibiting the transcription of RNA,
182 damaging DNA, and cell inflammation (Fig. 4). For example, Pd²⁺ ions may be
183 slowly absorbed by animal cells and distributed in the nucleus and mitochondria and
184 can affect and inhibit enzyme systems in animal cells.¹⁰¹

185 Pd-NMs in contact with animals in the environment could enter the organism and
186 produce form transformation. Pd initially is released as metal and oxide particles, but
187 they can then be transformed in the environment, in the digestive tract or in the cell

188 compartment to produce more harmful soluble substances.¹⁰² The main factors that
189 determine the toxicity of Pd NPs in the air are related to particle size and chemical
190 composition, and the inhalation routes are considered to pose a greater risk for health
191 effects.⁹⁸ The metal forms of Pd-NMs that are drawn into the respiratory tract from
192 the air are usually biologically inert, but some of the metal salts can become
193 sensitized chlorine compounds that are strong allergens and sensitizers.¹⁰³

194 Pd-NMs also displayed specific cross-sensitization with nickel and can penetrate
195 through the skin.²⁹ As the surface-to-mass ratio of Pd-NMs increases, their biological
196 activity increases, which in turn releases more active metal ions, ultimately leading to
197 an increased likelihood of penetrating the skin.²⁹ Noteworthy, compared with intact
198 skin, damaged skin significantly increases the absorption of Pd-NMs (Fig. 5).²⁹ Pd
199 content in the whole skin layer from the epidermis to dermis was significantly
200 decreased. Within the skin, these nanomaterials may have a long-term role and may
201 be involved in sensitization or spread throughout the body.

202 Significant cytotoxic effects of Pd-NMs have been observed in several human cell
203 models, including respiratory cells,^{87,104} cervical,¹⁰⁵ liver.¹⁰⁶ Pd-NMs released into the
204 environment can be inhaled by humans and accumulate in the respiratory tract, and
205 the absorption of Pd through the digestive tract was shown to be insignificant.¹⁰⁷
206 Meanwhile, human exposure to Pd-NMs may cause strong sensitization
207 reactions.^{108,109} Furthermore, Pd-NMs that enter the animal may accumulate in the
208 liver, kidneys, lungs, and bones.^{30,110} However, serum biochemical evaluation¹¹⁰

209 showed no significant hepatotoxicity in the mice after 7 days of intravenous injection
210 of various Pd-NMs (Fig. 6a). Moreover, no significant damage of Pd-NMs to the
211 organs of mice within 28 days was observed in the H&E stained images (Fig. 6b). But
212 meanwhile, noteworthy, the toxicity of Pd-NMs is affected by multiple factors, of
213 which dose and time are two main factors. Therefore, systematic study of the
214 ecotoxicity of Pd-NMs at different doses within different time periods (focusing on
215 the long-term potential toxicity) is still required.

216 Soluble Pd-NMs have been found to have serious health effects on higher
217 vertebrates in acute and chronic studies.¹¹¹ Soluble Pd-NMs enter the mouse, rat, and
218 rabbit body, which can cause mitochondrial membrane potential disorder, arrhythmia,
219 organ dysfunction, and even death.^{112,113} Furthermore, the time of retention of Pd in
220 the animals could affect the level of toxicity. Intratracheal or intravenous
221 administration can lead to the prolonged retention time of Pd in the animals than oral
222 administration of PdCl₂, resulting in higher biological toxicity. Pd-NMs accumulated
223 in living organisms may also be expelled from living organisms, and the main way for
224 these MNPs to be eliminated is through urine and feces.¹⁰¹ Several studies have
225 shown that soluble Pd-NMs also may cause patient poisonings, such as accumulate in
226 organ tissues, especially in the kidneys, lungs, liver, spleen, bones, and heart.¹¹⁴
227 Meanwhile, the mitochondria are important subcellular organelles for Pd-NMs
228 toxicity.¹¹² The works have shown that Pd-NMs toxicity is caused by the breakdown
229 of mitochondrial membrane potential and the depletion of cellular glutathione (GSH)

230 levels,¹¹⁵ which proposes Pd-NMs are more susceptible to toxicity to kidney tissue
231 than liver tissue. Histopathological findings in the kidney indicate that they change
232 renal tubular epithelial that further affects the glomerular filtration function.¹¹⁶

233 Furthermore, Pd-NMs can inhibit DNA and protein synthesis, and damage different
234 types of DNA in mouse lymphoma cell lines.¹¹² Pd-NMs can also inhibit the gene
235 expression of multiple metal markers and induced the conformational changes and
236 cleavage of DNA.¹¹⁷ Meanwhile, Pd-NMs interfere with the inflammatory process by
237 increasing the adhesion of eosinophils on endothelial cells at low concentrations.¹¹⁸
238 More pieces of research are needed to further study cellular and molecular
239 mechanisms to help understand the process by which Pd-NMs induce allergies and
240 inflammation.

241 To further understand the toxicity of Pd-NMs, the toxicity of Pd-NMs during
242 embryonic development are investigated. Zebrafish have a high genetic similarity
243 with humans. Zebrafish embryos have high optical clarity and can be screened on a
244 large scale, and have a low operating cost, which are powerful tools for environmental
245 toxicity detection.¹¹⁹ The molecular mechanism of Pd-NMs inducing embryonic
246 morphological changes by using zebrafish as a model. The results propose that Pd
247 treatment resulted in zebrafish embryo pericardial edema, inhibiting embryo survival
248 and hatchability, resulting in embryonic pericardial edema and cardiac malformation
249 (Fig. 7), affected the expression levels of several cardiac-related genes and

antioxidant enzymes, as well as revealing the underlying molecular mechanism of Pd-NMs inducing zebrafish embryonic heart malformations.¹⁰⁰

In many cases, technical problems prevented zebrafish model platforms from being fully established. First, zebrafish farming techniques are highly demanding, and changes in simple parameters such as temperature, pH, and symbiotic microbes may confuse test results.¹¹⁹ Second, so far, zebrafish models have focused on a limited range of environmental chemicals that are toxic at an early life stage.^{120,121} Therefore, powerful new phenotypic techniques and systematic approaches to identify a wider range of chemicals are needed to investigate the toxic effects of a large number of environmental factors to zebrafish models throughout their life stages. Finally, if we want to maximize the use of zebrafish models to describe the toxicological reactions of higher-level organisms, a more detailed understanding of the similarities between zebrafish and higher-level organisms is needed, including the combination of multichannel organ-specific studies, functional genomics, and automated image analysis techniques.¹¹⁹ In summary, the zebrafish model as an explanatory tool in the field of toxicology is promising for the systematic exploration of nanomaterial-environment interactions. And conceivably, with the establishment of a fully and robust zebrafish model platform, toxicity prediction and the development of new materials will eventually be synchronized.

269 Evaluation of Phytotoxicity

270 What are the hazards of Pd-NMs entering the environment to plants? Rare Pd-NMs
271 have been extensively redistributed in the biosphere as a result of human activity.
272 Lethal toxicity and cytotoxicity of Pd-NMs in terrestrial plants are developing rapidly.
273 There is a serious lack of public awareness of the environmental hazards caused by
274 the widespread use of trace nano-metal pollutants. In a natural state, plants have less
275 contact with rare metals and fewer resistance mechanisms than those found.¹²²
276 Therefore, to promote the application of these materials, it is necessary to investigate
277 the harm they caused in plants.

278 For the ecological toxicological effects of Pd-NMs on plants, Pd-NMs can quickly
279 enter the kiwifruit plant in large quantities, change the shape of pollen, and cause a
280 rapid loss of endogenous calcium in pollen grains which would result in pollen
281 plasma membrane damage.^{123,124} Moreover, Pd-NMs affect plant growth,¹²⁵ and seeds
282 germination,¹²⁶ but no effect of Pd NPs on plant growth was observed within 15 days
283 of plant culture, suggesting that NPs may not directly affect plant growth but
284 indirectly.¹²⁶ Further research is needed to understand whether environmental soil
285 exposure to rare Pd NPs has a short-term or long-term impact on crop production, as
286 well as specific toxicity mechanisms. Pollens are highly sensitive to environmental
287 pollutants, which is helpful for the accurate detection of the effects of Pd-NMs on
288 biological systems.

289 Taking kiwi pollen as an example, the studies have found that Pd-NMs enter the
290 kiwi pollen grains faster and enters more than soluble Pd(II) (Fig. 8a and b).¹²³
291 Compared with the effect of soluble Pd(II), under the granular action of low
292 concentration of Pd, the endogenous calcium of kiwifruit pollen is rapidly lost (Fig.
293 8c and d), thereby causing the damage of pollen plasma membrane, which seriously
294 inhibit the growth of pollen tube. Toxicity of Pd-NMs to plants is largely caused by
295 the reactive oxygen species (ROS) production,¹²² which can result in plant
296 photosynthesis disorders, membrane integrity destruction, and mitochondrial
297 dysfunction by attacking the interactions of cell membranes, proteins, lipids, and
298 DNA in cells (Fig. 4).

299 **Damage to Microorganisms**

300 Microorganisms are completing the cycle of various trace elements and are often
301 the common target of environmental toxicology research.¹²⁷⁻¹²⁹ The toxicology of Pd
302 catalysts was explored using the marine bacteria *V. fischeri* as a model.¹³⁰ *V. fischeri*
303 are cultured by designing a microenvironment that simulates the actual sampling
304 location. During the culture process, the effects of different concentrations of Pd-NMs
305 on respiratory metabolism are reported, and no side effects are found (Fig. 9).¹³⁰
306 Meanwhile, the effect of Pd-NMs on the structure of Polychlorinated biphenyl
307 (PCB)-dechlorinated microbial community, which enriched from marine sediment, is
308 also reported.¹³⁰ The investigation finds that Pd-NMs have no permanent impact on
309 their community organization, and even, on the contrary, increase the biodiversity of

microbial communities. All in all, the results of this work counter the hypothesis that Pd-NMs affect marine microbial communities. This is the first comprehensive study of the effects of Pd-NMs on marine microbial communities and provides significant information for the assessment of the toxicity of Pd-NMs.

For the other example, results have shown that Pd-NMs cause genomic alterations in the freshwater green algae *Pseudokirchneriella subcapitata*,¹³¹ which in turn causes great damage to the *Pseudokirchneriella subcapitata* at the growth process and morphology. Compared with animals and plants, the mechanism of toxicity of Pd-NMs to microorganisms mainly affect microbial community structure, growth, and diversity. Main mechanism of Pd-NMs to microorganisms is still oxidative stress. Exposure to Pd-NMs in the environment produces ROS directly or indirectly that interact with membrane proteins and bacterial cell walls, which may cause cell lysis, inhibiting DNA- RNA- and protein synthesis (Fig. 4).

Toxicity Assessment Methods and Challenges

As described in the previous sections of this article, the toxicity of Pd-NMs to the ecosystem (cytotoxicity, genotoxicity, inflammatory response, oxidative stress response, and so on) has been demonstrated. Parallel studies and toxicity assessments of these nanomaterials are needed to promote their production and application. However, the toxicity of Pd-NMs and/or other related nanomaterials and their interaction with ecosystems have not yet been systematically studied.¹³²⁻¹³⁴ Currently, most methods of toxicity assessment have been developed based on chemical toxicity.

331 Due to the unique properties of Pd-NMs, the results of these assessment methods may
332 be disturbed.^{106,135} Compared with traditional materials, the size, shape, specific
333 surface area, doping degree, solubility, agglomeration state, crystallinity, and other
334 characteristics of Pd-NMs may affect the characterization results of biological
335 effects.¹³⁶⁻¹³⁸ Incomplete and inaccurate characterization could lead to incorrect
336 assessment results. Besides, the non-development of unified and standardized
337 assessment methods is not conducive to the comparison of toxicity assessment results
338 among different research groups.

339 Detection of Pd dispersed in the environment requires a sophisticated analytical
340 method. Many techniques for detecting Pd in environmental samples have been
341 developed.^{139,140} To improve sensitivity, it is feasible to combine the existing
342 detection technology such as gas chromatography-mass spectrometry (GC-MS)
343 analysis.¹⁴¹ Compared to conventional solid-phase extraction methods, this newly
344 developed strategy could achieve higher sensitivity by increasing the concentration of
345 the maximum allowable coexisting heavy metal ions.

346 From the perspective of the toxicity test, in vitro toxicity test is the primary method
347 to study the ecotoxicity of Pd-NMs, because it is faster, easier and does not pose
348 ethical problems. However, toxicity studies of cultured cell systems in vitro do not
349 clearly explain cell-cell and cell-stromal interactions, ignoring cell diversity, and
350 lacking consideration for hormonal effects in vivo. Moreover, in vitro experiments
351 could be difficult to reflect the actual effect of the Pd-NMs in vivo.¹⁴² Therefore,

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4 352 toxicity assessment also requires in vivo experiments, and in vivo experiments can
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7 353 also study the long-term toxicity of Pd-NMs. Furthermore, pharmacokinetic studies
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10 354 help to maximize the interpretation of the correspondence between compounds in
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12 355 vivo and in vitro investigations.¹⁴³⁻¹⁴⁵ The pharmacokinetics of the Pd-NMs are
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15 356 helpful for the comprehensive quantitative analysis of the target tissues or cells acted
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18 357 on by the materials, the residence time in vivo, the toxicity time and dose, but before
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20 358 starting the pharmacokinetic study, should be the simulation the Pd-NMs into the way
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23 359 of organisms.

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25 360 In terms of microbiological inhibition, the toxicological evaluation results of
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28 361 Pd-NMs are different under different experimental conditions and in different
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31 362 microbial communities. For example, Pd-NMs had a strong growth inhibition effect
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34 363 on bacteria in the culture of single microorganism.^{146,147} However, in the simulated
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37 364 native microbial environment, as in the soil microbial system and the Marine
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40 365 microbial system, the toxicity inhibition of Pd-NMs on the microbial community can
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43 366 be ignored.^{126,130} This indicates that toxicity evaluation is multi-directional and
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46 367 environmental media can protect microbial community and prevent the toxicity of
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49 368 Pd-NMs to microbial community. Therefore, in addition to a single toxicity test,
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52 369 microbial metabolism and microbial community composition should also be assessed
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55 370 in the toxicological investigation of Pd-NMs on the actual microbial community, so as
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58 371 to understand the toxic effects of Pd-NMs on the microbial community from the
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60 372 perspective of molecular mechanisms.

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Conclusions and Perspectives

The widespread applications of Pd-NMs lead to their close contact with the ecological environment and diffused into the atmosphere, soil, water, and sediments. Subsequently, they migrate and transform in the environment and/or in the organism, and eventually cause harm to the organism. Many progresses have been made in the investigation of Pd-NMs release, absorption, transport, and risk assessment in animal models, contaminated soil-plant systems, and microbial communities (Table 1).^{21,28,29,87,100,103,110,116,123,125,130,148-158} However, the research progress in the toxicology of Pd-NMs lags far behind their production rate. The lack of uniform criteria and guidelines for evaluating research projects and outcomes in this field. As well as the unique nature of Pd-NMs as new materials have led to many problems in evaluating the toxicology of these Pd-NMs. All these problems could lead to conflicts and hinder the development of emerging nanomaterials.

These Pd-NMs may affect ecosystem function, exert cytotoxic and pro-inflammatory effects in vitro, and induce early changes in different target organs in vivo model tests. Further studies should more comprehensively and deeply characterize the physicochemical properties of Pd-NMs to explain in detail the complex interaction between their intrinsic characteristics and their toxic effects. As well as the nano-size of Pd-NMs makes it difficult to track them in the environment, so it is necessary to exploit a more accurate method to detect trace amounts of

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4 393 Pd-NMs in the environment. The problems should be analyzed by a systematic
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7 394 approach that involves the use of new high-precision equipment or new technologies
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10 395 and the characterization of Pd-NMs by multi-technique joint analysis.

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12 396 Most in vitro studies have reported that Pd-NMs induce severe cytotoxic effects
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15 397 and dysfunction in different animal and human cells (Table 1). In vitro studies are
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18 398 helpful to understand the toxic molecular mechanism of Pd-NMs. However, current
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21 399 data on toxicity of Pd-NMs in vitro are insufficient to reach a unified conclusion on
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24 400 mechanisms such as oxidative stress response, apoptotic pathways, cell cycle
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27 401 disorders, and DNA damage. In addition, the physicochemical properties, organic
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30 402 ligands, and surfactant interference with the toxicological properties of the Pd-NMs
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33 403 should also be considered. Therefore, it is necessary to further investigate the
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36 404 properties of Pd-NMs and the effects of complex interactions between cell growth
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39 405 mediators on different toxic patterns. As an example, differences in particle intake and
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42 406 culture conditions in cell models simulating upper and lower respiratory tract
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45 407 environments can interfere with the toxicological characteristics of Pd-NMs.⁸⁷ Faced
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48 408 with this situation, it is necessary to establish more complex in vitro models to obtain
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51 409 more comprehensive data and to help infer actual in vivo pathological characteristics.

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53 410 In vivo experiments show that exposure to palladium-based nanomaterials affects
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56 411 multiple organ systems, including endocrine system and kidney system (Table 1).
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59 412 However, the toxicokinetic behavior, and changes in short-term biological indicators
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413 (cytokines, hormone serum concentrations, and urine protein content, etc.) reported in

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4 414 these reports, remains to be explained whether they are stable at low doses and under
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7 415 long-term exposure conditions. From this perspective, Pd-NMs have a long-term and
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10 416 complex interaction with biological systems in the in vivo environment. Meanwhile,
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12 417 the toxicological characteristics of Pd-NMs may be related to exposure patterns,
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14 418 differences in individual immune systems, and concentrations in biological media.
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17 419 Future investigations should focus on the toxicological effects of Pd-NMs on animals
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20 420 under low-dose, long-term exposure conditions (in practice, some workers may be
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23 421 exposed to low-dose Pd-NMs for a long period of time).

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26 422 Many studies have focused on the biological hazards of nanoscale risk assessment,
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28 423 but few have studied the current and future concentrations and distribution of Pd-NMs
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31 424 in the environment. This requires a multidisciplinary team effort involving experts in
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34 425 materials, molecular biology, environmental toxicology, and physical chemistry. As
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37 426 well as the innovative experimental methods and protocols are needed to guide the
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40 427 assessment of the full cycle ecotoxicity of Pd-NMs. Furthermore, the quantitative
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42 428 measurement of the concentration level of Pd-NMs in the real environment is strongly
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45 429 encouraged. Considering the lack of standardized testing procedures, the diversity of
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48 430 exposure pathways, and the differences in toxicity tolerance among individuals, there
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51 431 are still many difficulties in environmental monitoring of Pd-NMs in the actual
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53 432 environment. Therefore, biological monitoring can be carried out at the same time,
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56 433 focusing on the investigation of biomarkers that mark physiological toxicity exposed
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59 434 in the real environment.
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4 435 In summary, effectively define the risk from the release of Pd-NMs, and use
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6 436 multidisciplinary methods to provide the guidance are beneficial. Both help to protect
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9 437 the health and safety of the ecological environment, conduce to clarify the toxicity of
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12 438 Pd-NMs, and contribute to paving the way for sustainable nanomaterials.
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16 439 Declaration of Interest

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21 440 The authors declare no conflicts of interest.
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Figure Captions

Fig. 1. Diagram summary of the applications, environmental behavior, and ecological risk of Pd-NMs.

Fig. 2. TEM image of Pd NPs synthesized using (a) Sanka coffee, (b) Bigelow tea, (c) Luzianne tea, (d) Starbucks coffee, (e) Folgers coffee and (f) Lipton tea extract at room temperature in one step without using any hazardous reducing chemicals or non-degradable capping agents. Reprinted with permission from ref. 57. Copyright 2008 the Royal Society of Chemistry.

Fig. 3. The source composition of Pd-NMs including electrochemical applications, catalytic organic transformation, the treatment of environmental contaminants, sensors, and nanomedicine.

Fig. 4. The toxicity effects of Pd-NMs to plants, animals and microorganisms.

Fig. 5. (a) Permeation profile of Pd after skin application of Pd NPs solution. (b) Mean values and standard deviations of Pd amounts (mg cm⁻²) in intact and damaged skin. Reprinted with permission from ref. 29. Copyright 2016 Elsevier Science Ltd.

Fig. 6. (a) Serum biochemistry analysis of mice treated with different coated Pd NSs at 7 days post injection. (b) Photos of H&E stained diaphragm slices from the mice treated with different coated Pd NSs at 28 days post injection. Reprinted with permission from ref. 110. Copyright 2015 American Chemical Society.

Fig. 7. Pd caused pericardial edema and cardiac malformation in zebrafish embryos. Reprinted with permission from ref. 100. Copyright 2014 Elsevier Science Ltd.

Fig. 8. Pd (a, b) and Ca (c, d) content in controls and in pollen treated for 30 and 90 min with increasing Pd concentrations administered as either Pd-NPs (a, c) or PdCl₂ (b, d). Reprinted with permission from ref. 123. Copyright 2009 Elsevier Science Ltd.

Fig. 9. Concentrations of spiked PCB congeners and their dechlorination products constituting more than 1% w/w of total PCBs at the end of incubation in the biologically active (white bars) and sterile (black bars) sets of spiked microcosms. (a) unamended microcosms; (b) hydrogen-amended microcosms; (c) microcosms amended with hydrogen + bioPd 5mg/kgdw; (d) microcosms amended with hydrogen + bio-Pd 50 mg/kgdw. Values are means of triplicate microcosms with error bars

representing standard deviation. Reprinted with permission from ref. 130. Copyright 2016 Elsevier Science Ltd.

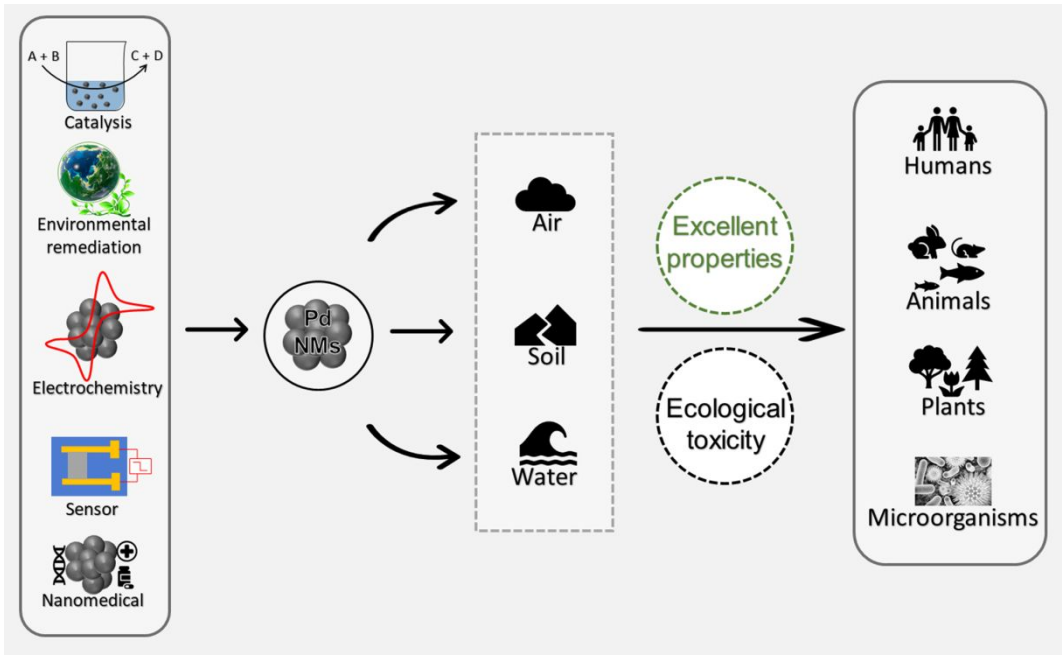


Fig. 1

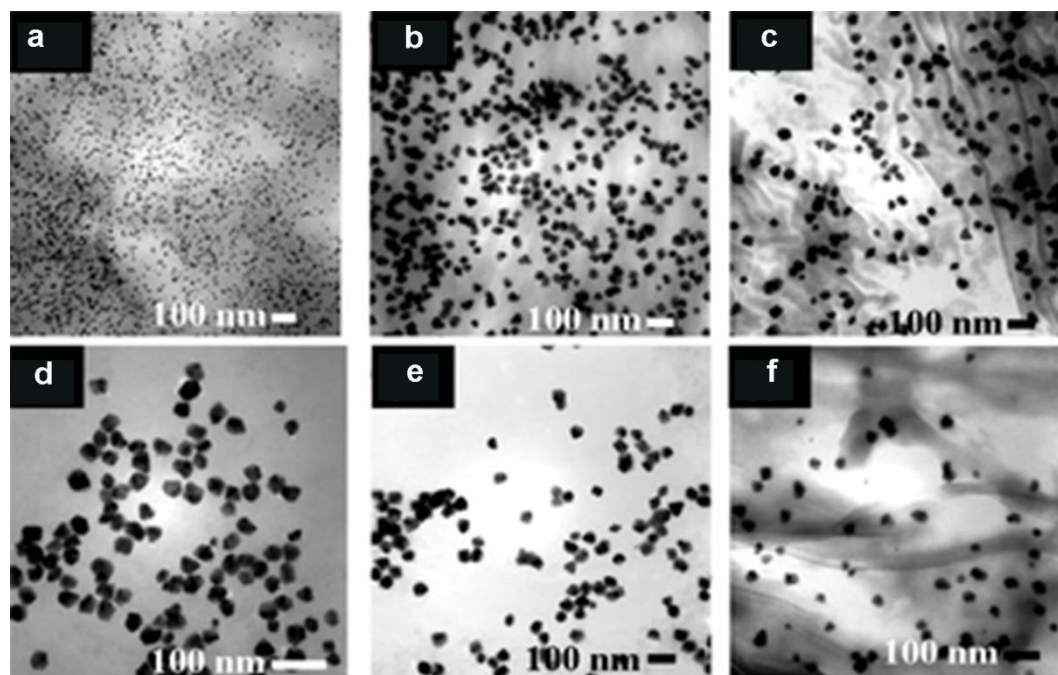


Fig. 2

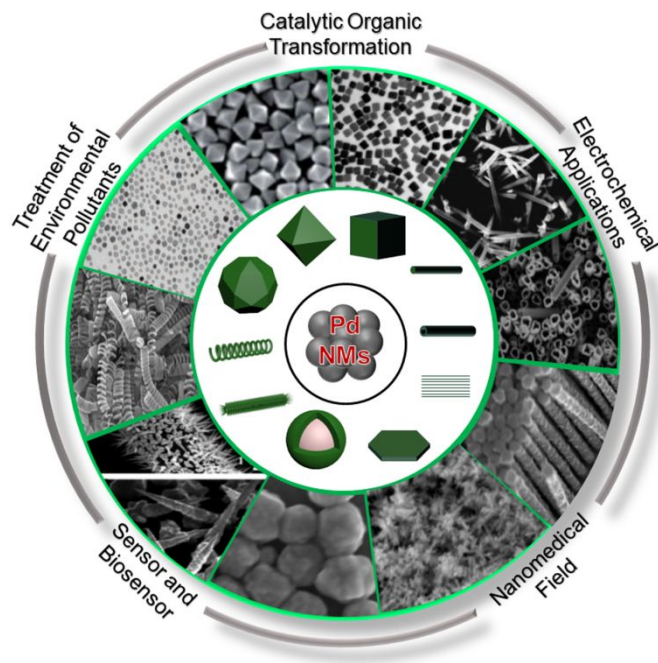


Fig. 3

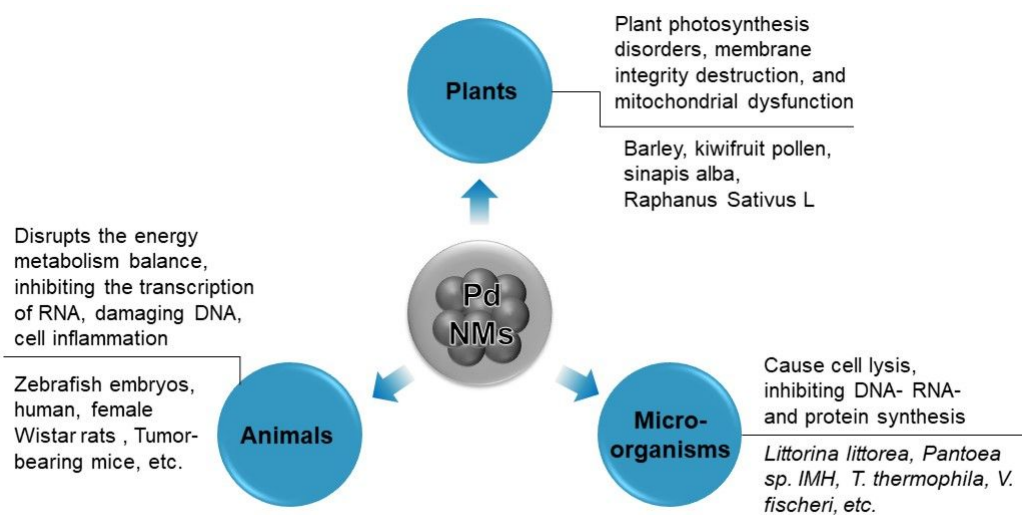


Fig. 4

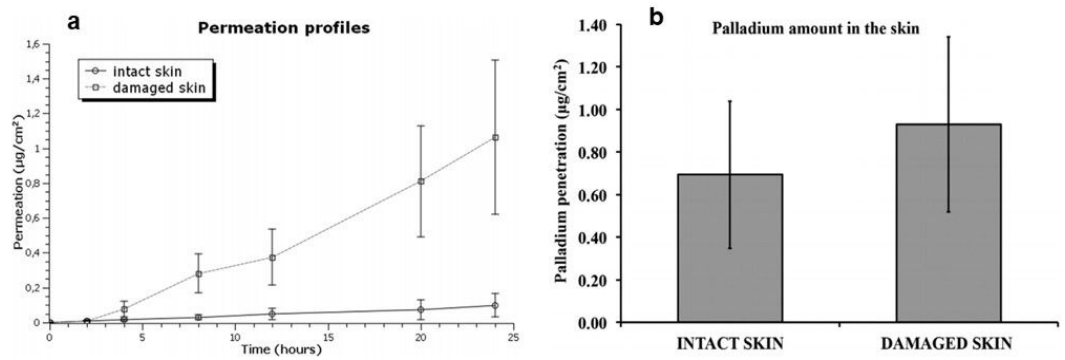


Fig. 5

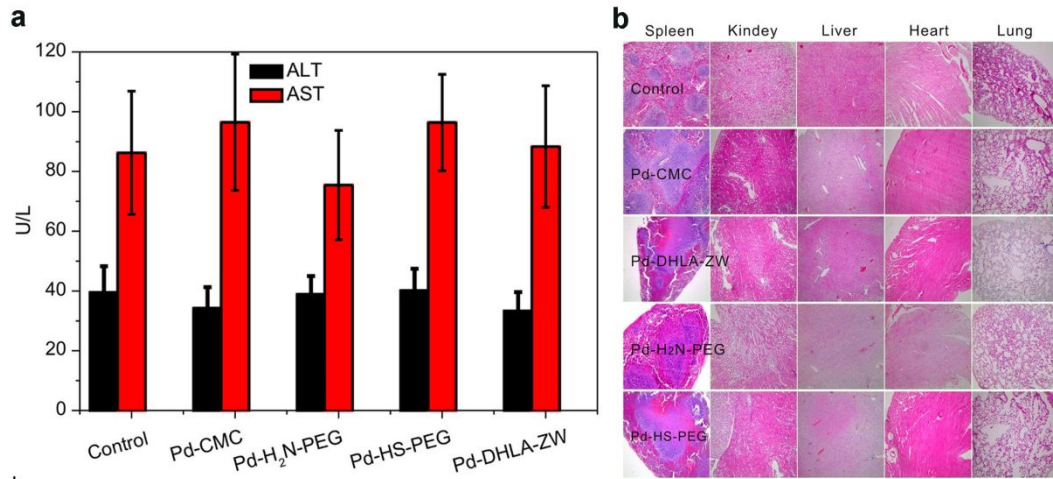


Fig. 6

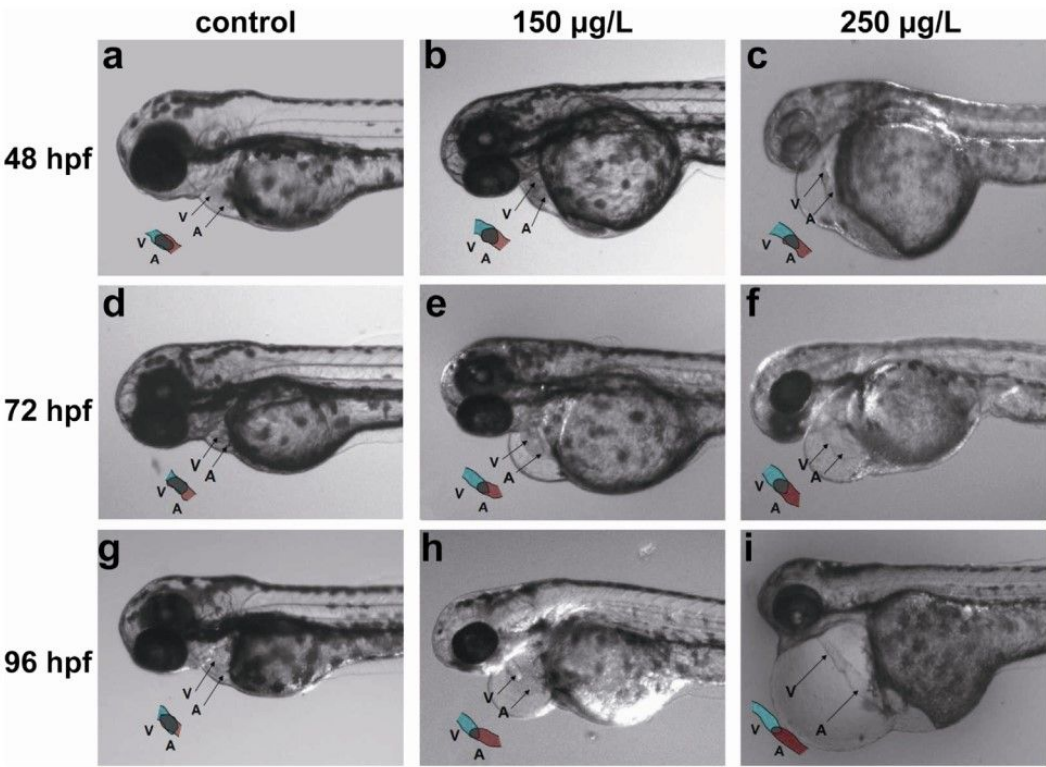


Fig. 7

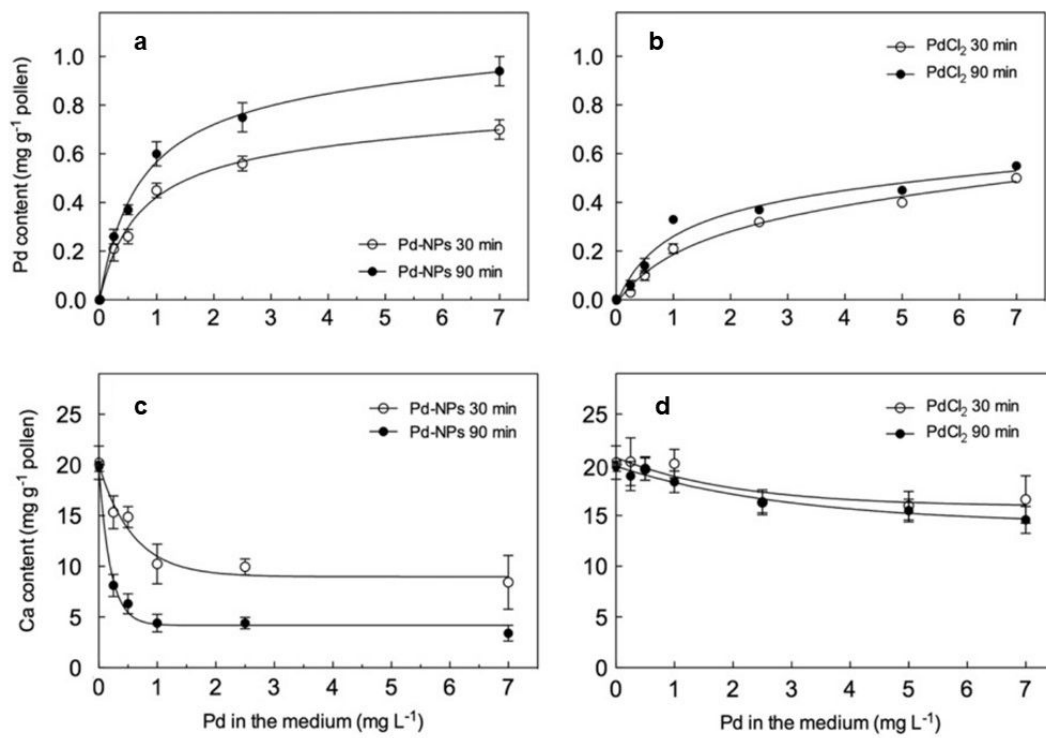


Fig. 8

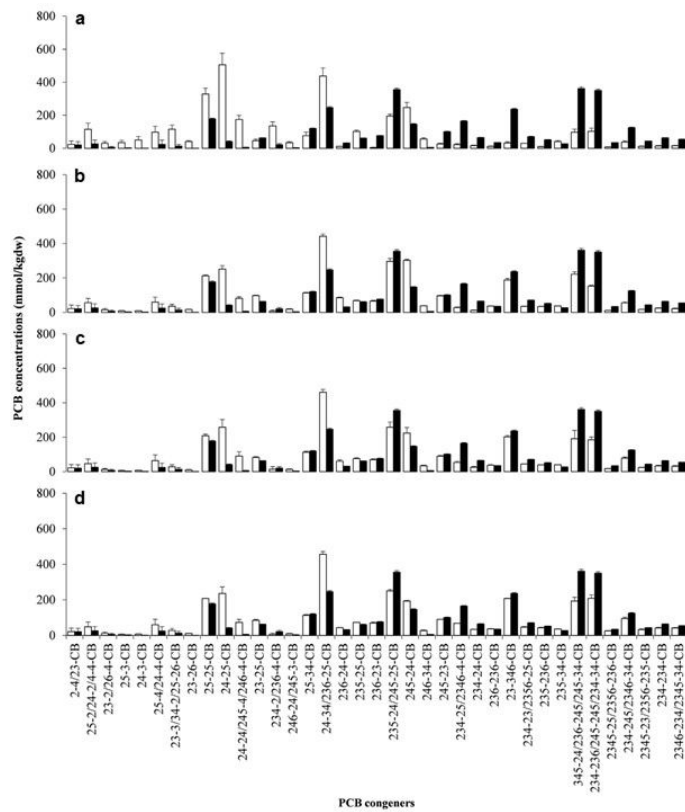


Fig. 9

Table 1: The toxicity of Pd-NMs to organisms.			
Pd-NMs	Investigated model	Findings	Ref.
Toxicity to animals			
Pd(OH) ₂	Simulated lung fluids	Form in the respiratory tract, and are toxic and allergic to humans and other organisms.	103
PdCl ₂ solution	Zebrafish embryos	Inhibits the survival rate and hatchability; Leading to pericardial edema and cardiac malformation; Inhibits the heartbeat rate; Induces the maladjustment of stress-related genes.	100
Pd/magnetite HNMs	Human skin (HaCaT) cell lines; Human colon (CaCo-2) cell lines; Rainbow trout gills (RTgill-W1) cell line.	Not trigger the production of ROS; Not affect the viability of selected mammalian and fish cell lines.	28
ZnO/Pd HNMs	Human skin cells	ZnO/Pd NPs were more photocytotoxic than ZnO NPs on the viability of human skin fibroblasts.	148
PdO/Co ₃ O ₄ HNMs	BEAS-2B cells; RAW 264.7 cells; Mouse lung	Superoxide production, glutathione depletion, cytokine production, and hierarchical cellular responses involving cytotoxicity in epithelial and macrophage lines; acute pro-inflammatory effects for mouse lung.	149
Pd NPs	Human primary bronchial epithelial cells (PBEC); Human alveolar carcinoma cell line (A549)	Absorbed by cells in PBEC; Reduce the response of PBEC to the pro-inflammatory cytokine TNF-R.	87
Pd NPs	Female Wistar rats	Cause significant tubular dysfunction of female Wistar rats; Significantly altered the epithelial cells of proximal and distal renal tubules with varying degrees of severity.	116
Pd NPs	Intact and damaged human skin in Franz cell	Permeate the skin in an in-vitro system; A potential long-term effect inside the skin.	29
Pd NPs	Human Ovarian Cancer Cells (SKOV3)	Cause a decrease in cell activity and proliferation ability; Cause the increase of cytotoxicity when the concentration increases; Induce SKOV3 cell apoptosis by inducing mitochondrial dysfunction.	150
Pd NSs	Tumor-bearing mice	No obvious hepatic toxicity by blood biochemistry assay; No detectable organ (spleen, kidney, liver, heart, and lung) damage by hematoxylin and eosin-stained imaging.	110
Pd NSs	Female Balb/c mice	Slight lipid accumulation in the liver; Led to spleen inflammation.	158
Pd NSs	ICR mice	Accumulate in the liver, spleen, tumor, and kidney; Cleared faster in the oral administration group.	151
Phytotoxicity			
Pd element	Barley	Causes stress to the leaves at a low nutrient concentration	125

Pd NPs	Kiwifruit pollen	Changed the morphology of kiwifruit pollen; Lead to the rapid loss of endogenous calcium in pollen, resulting in the damage of the pollen plasma membrane.	123
Pd NPs	<i>Sinapis alba</i>	The largest amount was found in leaves, followed by stems and roots.	21
HNT-Pd	<i>Raphanus Sativus L</i>	Increased the number of aberrations in low-vigor seeds.	152
Damage to Microorganisms			
Pd solution	<i>Littorina littorea</i>	Diet is the most important carrier of Pd accumulation for <i>Littorina littorea</i> .	153
Pd(II)	<i>Pantoea</i> sp. IMH	Induced expression of anti-stress protein; Induced detoxification of glutathione.	154
Pd NPs	<i>T. thermophila</i>	Accumulated extensively in the food vacuoles of <i>T. thermophila</i> .	155
Pd NPs	<i>V. fischeri</i>	Increased the biodiversity of <i>V. fischeri</i> ; No adverse effect on the overall structure of <i>V. fischeri</i> .	130
Pd NPs	<i>Candida albicans</i> ATCC10231; <i>Aspergillus niger</i>	Cell wall damage and oxidative stress.	156
Pd element	Caenorhabditis Elegans	Affect the growth ability of nematodes and effect on their reproductive ability.	157
NPs: nanoparticles; HNT: Halloysite nanotube; SP-ICP-MS: Single particle inductively coupled plasma mass spectrometry; HNMs: hybrid NMs; NPs: nanoparticles; NSs: nanosheets; IL-8: interleukin-8; PGE2: prostaglandin E2.; ISO 11348-3: International Organization for Standardization; ROS: reactive oxygen species; DCFH2-DA: 2',7' dichlorodihydro-fluoresceindiacetate.			

Table of Contents Entry

