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Abstract	Remediation using nanoparticles depends on proper documentation of safety aspects, one of which is their ecotoxicology. Ecotoxicology of nanoparticles has some special features; while traditional ecotoxicology aims at measuring possible negative effects of more or less soluble chemicals or dissolved elements, nanoecotoxicology aims at measuring the toxicity of particles, and its main focus is on effects that are unique to nano-sized particles, as compared to larger particles or solutes. One of the main challenges when testing the ecotoxicity of nanoparticles lies in maintaining stable and reproducible exposure conditions, and adapting these to selected test organisms and endpoints. Another challenge is to use test media that are relevant to the matrices to be treated. Testing of nanoparticles used for remediation, particularly redox-active Fe-based nanoparticles, should also make sure to exclude confounding effects of	

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## Chapter 28 Ecotoxicity of Nanomaterials Used for Remediation

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Abstract Remediation using nanoparticles depends on proper documentation of 5 safety aspects, one of which is their ecotoxicology. Ecotoxicology of nanoparticles 6 has some special features; while traditional ecotoxicology aims at measuring possi-7 ble negative effects of more or less soluble chemicals or dissolved elements, 8 nanoecotoxicology aims at measuring the toxicity of particles, and its main focus 9 is on effects that are unique to nano-sized particles, as compared to larger particles or 10 solutes. One of the main challenges when testing the ecotoxicity of nanoparticles lies 11 in maintaining stable and reproducible exposure conditions, and adapting these to 12 selected test organisms and endpoints. Another challenge is to use test media that are 13 relevant to the matrices to be treated. Testing of nanoparticles used for remediation, 14 particularly redox-active Fe-based nanoparticles, should also make sure to exclude 15 confounding effects of altered redox potential that are not nanoparticle-specific. Yet 16 another unique aspect of nanoparticles used for remediation is considerations of 17 ageing of nanoparticles in soil or water, leading to reduced toxicity over field- 18 relevant time scales. This review discusses these and other aspects of how to design 19 and interpret appropriate tests and use these in hazard descriptions for subsequent 20 risk assessments. 21

 Keywords
 Environment · Nanoparticles · Organic pollutants · Polluted soil ·
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 Toxicity
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#### 28.1 Introduction 24

25 Ecotoxicology for evaluating possible negative effects of nanomaterials (nanoecotoxicology) has some special features; while traditional ecotoxicology 26 aims at measuring possible negative effects of more or less soluble chemicals or 27 dissolved elements, nanoecotoxicology aims at measuring the toxicity of particles. In 28 addition, nanoecotoxicology has its main focus on those effects that are unique to 29 30 nano-sized particles, as compared to larger particles or solutes. For this reason, experiments in nanoecotoxicology usually compare the results with effects caused 31 by larger particles with similar composition. Another common comparison is that to 32 the effects of dissolved ions of the same elements constituting the nanoparticles 33 tested, since many metal-based nanoparticles may partly dissolve, and the toxic 34 35 effects can be due to their soluble ionic component. These so-called control treatments are not always easy to establish, or they may result in imperfect comparisons, 36 as larger-scale particles (often referred to as "bulk material") may behave quite 37 differently due to their larger size, and soluble salts of elements found in many 38 nanomaterials may not exist, or may precipitate during the tests (Kahru and 39 Dubourguier 2010; Handy et al. 2012a; Sørensen et al. 2015). 40

Exposing organisms to nanomaterials requires stable suspensions of these 41 nanomaterials. This is typically obtained through the use of surface-active agents 42 reducing the attractive forces between particles (Labille and Brant 2010), causing 43 them to remain suspended in water or other media for a period of time that would 44 permit absorption or other interactions causing harm to the test organism. These 45 surfactants may themselves affect the test organisms, and thereby the test outcome. 46 47

Control treatments used for comparisons should therefore take this into account.

#### **Toxicity of Particles** 28.2 48

Particle toxicity can be rather different from toxicity of soluble substances or ions. 49 This is due to the strong barriers against the uptake of particles that most organisms 50 possess. Ions enter organisms through channels (transporters) in the cell mem-51 branes, which can discriminate the uptake based on characteristics like charge 52 and size. Organic molecules may either pass through uptake channels for organic 53 nutrients or cross bilayer membranes due to their hydrophobicity. Nanoparticles fit 54 none of these routes of transport, and are therefore relegated to entering cells by 55 56 random "back doors", like compromised cell membranes or accidental passive uptake. The size of the particles in question is then of course of major importance 57 for such uptake. Most nanoparticles used for remediation are found among the 58 larger nanoparticles, typically >50–200 nm. To put this in perspective, a 20 nm 59 silver nanoparticle would contain 750,000 atoms (Oughton et al. 2008), each atom 60 61 approximately twice as big as, e.g., a zinc ion that enters cells through ion channels.



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Needless to say, most of the regular paths for entering into cells are not permitting 62 the entry of nanoparticles. One exception here may be endocytosis. 63

Yet, some nanoparticles find their way through cell walls and membranes, and 64 end up inside cells. Here they represent quite a different type of toxicity than 65 dissolved ions of the same elements, partly because they constitute discrete particles 66 rather than diffusive ions that may spread among cells. A particle is likely to stay 67 inside the cell it has entered and end up in lysosomes (in the case of eukaryotic cells), 68 but may, e.g., dissolve and represent a steady source of dissolved ions that may be 69 harmful. The concentration of such ions is likely to be substantially higher in a cell 70 containing a nanoparticle than in a cell being exposed (along with neighboring cells) 71 to a similar level of dissolved elements that may move more or less freely within the 72 exposed tissue.

Nanoparticle toxicity mechanisms include effects of dissolved ions from metallic 74 nanoparticles (common for Ag nanoparticles that release  $Ag^+$ ), induction of reactive 75 oxygen species (ROS and other types of oxidative stress; such effects have been 76 shown for iron-based nanoparticles (Lewinski et al. 2008)) and damages directly 77 related to the surface and shape of nanoparticles (e.g., nanotubes and their asbestos-78 like induction of damages on cells). For Fe-based nanoparticles, only mechanisms 79 related to ROS-formation and oxidative stress have been described as direct effects. 80 These have been reviewed recently (Lei et al. 2018), and will not be detailed further 81 here. Indirect effects on  $O_2$  availability is another mechanism, but as we argue 82 below, this is not a nanoparticle-specific toxic effect. 83

## 28.2.1 Ageing and Other Time-Dependent Modifications of Toxicity 85

A particular aspect of nanoparticle toxicity testing that is relevant for nano-sized 86 zero-valent iron (nZVI) and other nanoparticles to be used for remediation is 87 reduction in toxicity over time. While all nanoparticles are subject to changes in 88 surface properties as a result of interactions with environmental matrices, 89 nanoparticles for remediation are frequently designed to lose their reactivity by 90 interacting with environmental pollutants. Including temporal changes in toxicity 91 during testing is therefore a particularly relevant aspect that should be assessed for 92 such materials. Reduced toxicity as a result of ageing in soil has indeed been 93 demonstrated for nZVI aged for 30 days in soil, using growth (body weight) of 94 earthworms as an endpoint (El-Temsah and Joner 2012a) and partly for rice after 95 2-4 weeks ageing (Wang et al. 2016). In the natural environment, living organisms 96 will mostly be exposed to aged nZVI, and not to pristine particles. This is important 97 to keep in mind when designing toxicity studies. Not only may such tests show that 98 adverse effects of the nanoparticles are short-lived, but it may also be helpful in 99 designing nanoparticles for remediation as toxicity and reactivity against pollutants 100 are likely to be strongly linked (Hjorth et al. 2017). 101



Different types of nanomaterials may affect organisms differently. This is yet another aspect to consider when choosing how a given nanomaterial is tested with respect to ecotoxicity. For approval of new nanomaterials or for conducting risk assessments, a set of minimum three tests with contrasting organisms must be carried out (Baun et al. 2009).

#### 107 28.3 Choice of Test Organisms

The choice of test organisms is important for several reasons, and may ultimately 108 determine the outcome of a testing scheme. First, the choice of organisms must be 109 relevant for the matrix to be treated. If nanomaterials for treating polluted soil are to 110 be tested, soil organisms should be chosen. Similarly, freshwater and marine organ-111 isms are relevant to their native habitats. Within these three major organism habitats, 112 there may be some overlap, or it may be relevant to include organisms from two 113 groups as a remediation situation can affect more than one matrix: treated soil may 114 lead to nanomaterials ending up in nearby ponds and streams, or streams and rivers 115 may reach brackish or saltwater habitats. 116

117 When choosing test organisms within these major groups, there are at least three 118 key aspects to consider:

119 • How contact with the tested material may occur

120 • Which endpoints are available to assess effects

121 • Which trophic level the organism belongs to (and how this will affect exposure).

Ecotoxicity can be strongly affected by the mode of exposure. Dermal contact is 122 commonly affecting an organism less than ingestion or interference with respiratory 123 organs. This distinction is less relevant for, e.g., microorganisms and plants, but even 124 for microorganisms and plants that have no intestines, internalization may occur and 125 cause different toxicity than surface contact. In many cases, the nature of the 126 organism's natural habitat and the test design will determine the mode of exposure. 127 Plants may, e.g., be exposed in an aqueous suspension (seed germination tests and 128 hydroponic plant cultures), or in solid matrices with more or less resemblance to a 129 real soil at a site to be treated (El-Temsah and Joner 2012b). While exposure in 130 aqueous suspensions may say something about the inherent toxicity of the 131 nanoparticles tested, it will give a far higher exposure than equivalent tests using 132 soil, and should thus include appropriate exposure estimates when questions of risks 133 are addressed. When testing toxicity of nanoparticles to plants or soil organisms 134 using soil as an exposure medium, the choice of a test soil is also decisive, as the 135 136 relative amount of different soil constituents may vary considerably and affect both bioavailability of particles and whether plants or soil organisms thrive in them. 137 Using a soil with a minimum of soil organic matter will go a long way to ensure 138 that plants germinate and grow in them, or that earthworms are active in ingesting 139



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soil during a test. But organic matter in soil may also result in a different availability 140 of nanoparticles compared to a sub-soil void of humus, which is far more representative of soils being remediated using such particles. This is a trade-off situation 142 where test organisms and test media should be selected as to be appropriate for the 143 purpose of the test. 144

Another example of exposure control concerns earthworms. Dermal exposure of 145 earthworms is usually measured by dissolving or suspending the material to be tested 146 in water that imbibes a filter paper lining a glass vial where worms are placed (OECD 147 1984). Exposure through ingestion, on the other hand, uses a soil matrix where the 148 material to be tested is mixed in. As in the example of exposing plant roots to 149 nanoparticles in water or soil, exposure conditions for earthworms also differ greatly 150 between water and soil. However, for worms the exposure matrix also determines 151 mode of contact: nanoparticles suspended in water mainly result in dermal exposure, 152 while nanoparticles mixed into soil or feed result in intestinal exposure plus dermal 153 contact (Lapied et al. 2010). To relate data from the rapid and inexpensive dermal 154 tests to test made with soil where bioavailability of nanoparticles is reduced by 155 interactions with the soil components, one may perform dermal contact tests in soil 156 by preventing worms from ingesting soil by gluing shut their mouths using super 157 glue. The contribution from intestinal exposure may then be found by comparing 158 worms with and without glued mouths. 159

For many test organisms, exposure through ingested material may differ widely 160 according to how nanoparticles are introduced into the test system. Here, 161 nanoparticles mixed into feed may result in far higher exposure than if directly 162 mixed into soil. While certain earthworms (epigeic and anecic worms) seek out 163 organic debris when they forage, other worms (endogeic worms) ingest soil and feed 164 on the evenly distributed organic matter therein. Thus, exposing earthworms, e.g., to 165 nanoparticles contained in organic feed or mixed homogeneously into soil that 166 represents a volume that would frequently be at least 50 times higher may result in 167 very different rates of uptake. Similarly, nematodes may be exposed to nanoparticles 168 adsorbed onto bacteria upon which they feed, or through a suspension of 169 nanoparticles where no prior association between bacteria and nanoparticles has 170 occurred (Kleiven et al. 2018). To maximize ingestion by nematodes or other 171 particle feeders (e.g., Daphnia), the test may omit the feed (e.g., bacteria and 172 algae, respectively), but this may cause constipation and blockage of the digestive 173 tract of the test organisms, and adverse outcomes that are caused by excessively high 174 availability of nanoparticles (Roberts et al. 2007). In a more realistic exposure 175 situation, the organisms would ingest mainly digestible particles that would ensure 176 normal gut passage. 177

Aggregation (including agglomeration) of nanoparticles during exposure in aqueous media is a major determinant of exposure when particle uptake is sizedependent. Both medium constituents, particularly divalent ions like  $Ca^{2+}$ , excretions from test organisms and pH changes may cause this (Keller et al. 2012; Baker et al. 2014). Benthic organisms typically feeding on larger particles may potentially experience higher exposure due to aggregation, as discrete nanoparticles may be too small to be perceived as food. As mentioned above, surface-active compounds may 184 185 reduce aggregation and counteract aggregation effects, and even organisms may 186 cause dispersion by producing organics that stabilize nanoparticles in suspension 187 (Unrine et al. 2012).

#### 188 28.3.1 Endpoint Selection

An ecotoxicological endpoint is the parameter measured as a response to a poten-189 tially toxic substance. Numerous endpoints may be used when assessing the effects 190 on a test organism. For many organisms, mortality or growth rate are rather coarse 191 endpoints used for testing acute toxicity, while enzymatic activities (e.g., of anti-192 oxidative enzymes), genetic mutations, or expression of genes related to damage 193 repair or stress are gradually more sensitive test endpoints, permitting detection of 194 more subtle and chronic adverse effects at lower concentrations (Walker et al. 2001). 195 The interpretations of the responses to the endpoints with less obvious toxicity 196 functions are however a minefield. Are they, e.g., altered behavior, avoidance, 197 expression of stress-related genes, or enhanced frequencies of apoptosis indicators 198 of toxicity? For example, if nZVI is introduced into soil or a test system, the redox 199 conditions may change rapidly as to cause oxidative stress (or irreversible organis-200 mal damage) due to reactions of free  $O_2$  with nZVI. But frequently such conditions 201 are short-lived (El-Temsah et al. 2013), and oxygen will diffuse in from the border 202 zones and re-establish oxygenated conditions and alleviate the stress caused by the 203 reduced O<sub>2</sub> availability. For those organisms that have survived the period of 204 reduced O<sub>2</sub> availability, the effects may be fully reversible, with no negative impacts 205 on populations or communities (Nguyen et al. 2018a). Thus, if the exposure ceases 206 and the organism has avoided it or only passed through a period with sub-optimal 207 living conditions (stress) due to the nanoparticle exposure, it is not appropriate to 208 interpret this as toxicity. 209

A particular consideration to make when it comes to testing of nZVI and other 210 nanoparticles for remediation that may affect oxygen availability to organisms is the 211 fact that these nanoparticles may cause a lack of oxygen needed by aerobic organ-212 isms during respiration. nZVI may, e.g., react with the available O<sub>2</sub> in the test 213 medium as to render the conditions anoxic, thus asphyxiating the test organisms. 214 This is particularly relevant for exposure in water and wet soil where  $O_2$  diffusion 215 and replenishment is slow. Such induction of anaerobic conditions and its detrimen-216 tal effects on aerobic organisms is not a nano-specific effect (though the dynamics of 217 218  $O_2$  consumption may differ between nanoparticles and similarly reductive chemicals/bulk-size particles due to the specific surface area and chemical reactiv-219 ity). The effects nZVI may have on alternative electron acceptors (NO<sub>3</sub>, SO<sub>4</sub>, 220 oxidized forms of Mn, etc.) can similarly preclude the use of anaerobic test organ-221 isms to circumvent the need for O<sub>2</sub> during testing. 222

Testing for nanoparticle-specific effects when the nanoparticles to be tested cause changes in the availability of  $O_2$  or other electron acceptors require the use of control treatments that have comparable redox conditions, or the use of test systems or



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exposure matrices that buffer against such changes, coupled with appropriate monitoring of redox potentials during the tests. 227

Nanoparticles that form colored suspensions may lead to a particular set of 228 confounding effects related to shading of light. This is relevant for algae and other 229 photosynthetic organisms that may experience lower light availability when used in 230 tests where nanoparticle suspensions are dense enough to reduce transmittance 231 (Handy et al. 2012b; Hjorth et al. 2016; Nguyen et al. 2018b). Algal growth rates 232 or measurements of photosynthesis, chlorophyll content and related endpoints 233 should thus account for confounding effects of shading. 234

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#### 28.3.2 Trophic Interactions

The trophic level of an organism can determine the way it is exposed to 236 nanomaterials. This is partly due to the feeding habits of organisms at different 237 trophic levels. Free-living microalgae may absorb nanoparticles directly from the 238 water suspending the particles, but particles may also affect the algae by affecting the 239 amount of available nutrient ions, or by shading the algae from light as to reduce 240 photosynthesis. A filter feeder grazing on these algae may experience a similar 241 concentration of nanomaterials through contact with water, but will in addition 242 ingest, e.g., the aforementioned microalgae that may contain nanomaterials. 243 Depending on whether bioaccumulation (increased concentrations in organisms 244 with increasing lifetime) or biomagnification (predators accumulating higher con- 245 centrations than found in their prey) occurs, the next level predator may experience 246 different exposure through the ingested food. So far, bioaccumulation has been 247 observed for some nanomaterials (Petersen et al. 2008; Wang et al. 2013), and in 248 some cases even biomagnification (Judy et al. 2011; Majumdar et al. 2016; Gupta 249 et al. 2017). No such studies have been made with nanomaterials used for remedi- 250 ation. In some cases, the bioaccumulation may be due to the fact that an element 251 contained in the nanoparticles in question is a micronutrient that the organism needs 252 and scavenges for, as observed for cobalt nanoparticles (Coutris et al. 2012). Iron, 253 found in many nanoparticles used for remediation, is likely to behave similarly if test 254 organisms are experiencing sub-optimal iron supply. 255

#### **28.4** Standardized or Non-standardized Tests?

A major part of the research on nanoparticle toxicity has been made using 257 non-standardized tests, in the sense that they do not follow test protocols approved 258 by standardization organizations like the Organisation for Economic Co-operation 259 and Development (OECD) and International Organization for Standardization (ISO). 260 Non-standardized tests have the advantage of choosing freely among organisms, 261 endpoints, and exposure media. This allows for exposure optimization and 262



263 exploitation of the vast knowledge on biota, ranging from their behavior, physiology, metabolism, reproduction, and genetic peculiarities to community dynamics 264 and ecosystem functions when interacting with their habitat. Non-standardized tests 265 may thus be best suited to elucidate toxicity mechanisms or describe pertinent 266 environmental consequences of spreading potentially toxic nanoparticles. In com-267 parison, standardized tests are limited to easily culturable organisms exposed under 268 well-defined conditions, using a limited number of rather crude endpoints. The 269 advantages of using standardized tests are that the results can easily be compared 270 with those obtained for other chemicals, which in turn permits hazard classification, 271 and that standardized test results can easily be used for product documentation when 272 chemicals are used in commercial products requiring approval regarding possible 273 negative environmental effects. 274

### 275 28.4.1 Standardized Testing Methods

OECD and ISO have published a number of test guidelines (TGs) that describe in all details how chemicals testing for approval of new chemicals should be conducted. These tests have been developed for soluble chemicals and have not taken into account considerations that may be important for toxicity testing of nanoparticles. Yet, the OECD has concluded that the approaches for testing and assessment of traditional chemicals are in general appropriate for assessing nanomaterials, but that

the tests may have to be adapted to the specificities of nanomaterials (see ref. OECD). This concerns, e.g., methods of sample preparation, particularly regarding homogenization and distribution of the nanoparticles in the test media. Similarly, adaptations may be needed for certain test guidelines.

The first step is the preparation of a stable suspension of nanomaterials (see references in Hund-Rinke et al. 2016). This can be obtained through the use of surface-active agents reducing the attractive forces between particles and causing them to remain suspended in water or other media for a period of time that would permit absorption or other interactions causing harm to the test organism. These surfactants may themselves affect the test organisms, and thereby the test outcome, and should therefore be included in control treatments.

It is a common requirement of standardized test guidelines that exposure con-293 centrations should remain stable (often stated as no more than 20% deviation 294 between exposure concentrations and nominal concentrations) over the duration of 295 the test. This is a challenge with nanomaterials, which easily agglomerate and 296 sediment out of a water column (to mention the case of aquatic tests), exposing 297 pelagic organisms to lower concentrations and benthic organisms to higher concen-298 trations than originally intended. Several factors influence the agglomerating and 299 settling behavior of particles, such as agitation of the test system, ionic strength, pH, 300 presence of specific ligands/chelating agents, and organic matter content of the 301

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exposure medium. The following modifications have been proposed by Hund-Rinke 302 et al. (2016) for maintaining (more) constant exposure conditions in test systems: 303

- Conduct the OECD TG 202—acute immobilization of *Daphnia magna* test 304 (OECD 2004) at pH values enabling more stable dispersions of nanomaterials 305 and use a growth medium with very low ionic strength, e.g., very soft EPA 306 medium. This medium has been shown to allow normal growth and reproduction 307 of *D. magna*.
- In the OECD TG 210—fish, early-life stage toxicity test (OECD 2013) with 309 zebrafish, improve nanomaterial dispersion by using exposure chambers coupled 310 with water changes every 24 h.
- In tests using spiked soil or sediment, i.e., OECD TG 216—nitrogen transforma- 312 tion test (OECD 2000a); OECD TG 217—carbon transformation test (OECD 313 2000b); OECD TG 220—enchytraeid reproduction test (OECD 2016a); OECD 314 TG 222—earthworm reproduction test (Eisenia fetida/Eisenia andrei) (OECD 315 2016b); OECD TG 225—sediment-water lumbriculus toxicity test (OECD 316 2007), add nanomaterials to each replicate, to ensure homogeneity of spiking. 317 Exceptions can be made for low concentrations, where this modification can be 318 difficult to implement.

There has been a concern that some OECD test guidelines were not suited for the 320 detection of toxic effects, in the sense that they would underestimate the potential 321 toxicity of some nanomaterials. One way this underestimation could occur is by 322 reduction of the bioavailability of nanomaterials and their transformation products 323 due to sorption to organic matter or the elevated pH. Underestimation of the toxicity 324 of nanomaterials can also occur when the duration of the test is too short compared to 325 the slow transformation of nanomaterials in soil, which can be the source of toxic 326 chemical species. The following modifications proposed by (Hund-Rinke et al. 327 2016) may minimize the interference of the nanomaterials with the components of 328 the test media or the toxicity endpoints: 329

- OECD TG 201—freshwater alga and cyanobacteria, growth inhibition test 330 (OECD 2011): the chelating agent EDTA can interfere with metal nanomaterials 331 and a modified EDTA-free version of the OECD algal medium (OECD-M) for 332 *Raphidocelis subcapitata* is proposed. 333
- For the OECD TG 201 (OECD 2011), it is recommended to measure biomass by 334 determination of in vitro chlorophyll a, instead of optical density and in vivo 335 fluorescence measurements or cell counting by hemocytometry.
- OECD TG 216—nitrogen transformation test (OECD 2000a): for the testing of 337 ion-releasing metal nanomaterials, the pH of the soils should be at the lower end 338 of the range accepted according to the test guideline (pH 5.5). It is also proposed 339 to extend the duration of the test to 56 days, since some nanomaterials only show 340 effects after ageing, and to include multiple short-term measurements of the 341 potential ammonium activity, instead of single measurements at the start and 342 the end of the test. 343

• OECD TG 217—carbon transformation test (OECD 2000b): similar modifications are proposed for this test, except the multiple short-term measurements.

#### 346 28.4.2 Fe-Based Nanoparticles Exempt from Nano-Fear

nZVI and Fe-oxide-based nanoparticles have, to some extent, dodged the skepticism 347 that clings to other types of nanoparticles. This is partly because nanoparticles for 348 remediation are used to treat and remove the harmful effects of toxic pollutants, thus 349 reducing the exposure of humans and the environment to highly toxic and mobile 350 chemicals like TCE (trichloroethylene) for which there are no doubts of adverse 351 effects or environmental exposure. Further, Fe-based nanoparticles have repeatedly 352 been shown to have limited or even very limited mobility in the environment, 353 restricting movement away from the treated areas (Johnson et al. 2013), which are 354 often fenced in and unavailable to the public. Thus, risks appear confined to the 355 treated areas. A third point in favor of Fe-based nanoparticles comes from the fact 356 that most natural environments contain ample amounts of Fe, even in forms similar 357 to those coming out of nanoremediation treatments. The products of nZVI aged in 358 aerated water are mainly Fe<sub>3</sub>O<sub>4</sub> (magnetite) and  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (maghemite), accompanied 359 by  $\gamma$ -FeOOH (goethite). If corrosion continues, the products are predominantly 360  $\gamma$ -FeOOH, with small amounts of Fe<sub>3</sub>O<sub>4</sub> and  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (Liu et al. 2015). The final 361 aqueous corrosion product of nZVI is FeOOH (Pullin et al. 2017; Lei et al. 2018). 362 Finally, Fe-based nanoparticles have been scrutinized in several research projects in 363 parallel to their development, and the outcome of the ecotoxicity measurements as 364 well as practitioners feedbacks indicate that Fe-based nanoparticles are causing low 365 concern, if any (Bardos et al. 2011; Hjorth et al. 2017). 366

### 367 28.4.3 Ecotoxicity Does Not Equal Risk

Risk is the product of hazard (ecotoxicity) multiplied by the probability of encountering hazard (exposure). As mobility of nZVI and other Fe-based nanoparticles is limited, and as their prescribed use targets subsoils at several meters depth in industrial brownfields, the risk to humans (apart from workers exposed during production, transport, and deployment) and wildlife is extremely low or inexistent. The low hazard level due to the low inherent toxicity of the Fe-based nanomaterials of course also contributes to the low risks. Author's Proof

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