



Cite this: *RSC Adv.*, 2017, 7, 37065

Fate and transformation of nanoparticles (NPs) in municipal wastewater treatment systems and effects of NPs on the biological treatment of wastewater: a review

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Many types of nanoparticles (NPs) have been reported to be toxic to organisms. Wastewater treatment plants are among the important pathways for the migration and transformation of NPs in the environment. Based on a summary of the toxicity of NPs to microorganisms, this review discusses the advances in the fate and transformation of NPs in wastewater treatment plants (WWTPs) and the impacts of NPs on the biological treatment of wastewater. These NPs primarily include Ag NPs, Cu NPs, CuO NPs, ZnO NPs, TiO₂ NPs, Al₂O₃ NPs, SiO₂ NPs, carbon nanotubes (CNTs) and magnetic NPs. This review observes that the different properties of these NPs result in significant differences in the fate, transformation, antimicrobial activities and effects on biological wastewater treatment. Although many studies have been undertaken in this field, more studies are needed to address the adverse effects of NPs on organic and nutrient removal, as well as their toxic mechanisms.

Received 20th May 2017

Accepted 5th July 2017

DOI: 10.1039/c7ra05690g

rsc.li/rsc-advances

1. Introduction

Nanomaterials and nanotechnology have affected human life in many ways. Many types of nanoparticles (NPs), such as carbon nanotubes (CNTs), zinc oxide (ZnO NPs), titanium dioxide (TiO₂ NPs), manganese oxide (MnO₂ NPs), copper oxide (CuO NPs) and copper (Cu NPs), and silver (Ag NPs), have been widely used in such applications as medicines, pesticides, alternative energy, catalysts, and consumer products.¹ For example,² Ag

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NPs have been used as an antibacterial agent due to their antiseptic properties in children's toys, clothing and washing machines. Cu NPs have been applied in bioactive coatings, air and liquid filtration, skin care products, wood preservatives, textiles, and coatings on integrated circuits. During the production and applications of nanomaterials, a portion of them will likely enter the environment. Owing to their large surface area and small size (1 to 100 nm), nanoparticles (NPs) possess special physical and chemical properties. The ecotoxicological evaluation of NPs is an equally important problem.³ In recent years, a number of ecotoxicological studies have been carried out on NPs in aquatic organisms, such as bacteria, cells, algae, and zooplankton.⁴⁻⁷

It is known that wastewater treatment plants (WWTPs) are one of the most important pathways for NPs to enter the environment.^{8,9} Activated sludge in WWTPs, a suspended microbial aggregate playing a primary role in pollutants removal, contains many types of microorganisms and their extracellular products. Because NPs have been shown to be toxic, their impacts on the functional microorganisms of activated sludge are worth studying. Numerous published studies have investigated this subject.

The present review begins with a summary of the toxicity of various NPs to microorganisms. We summarize and review the fate and transformation of NPs in wastewater treatment systems, as well as the effects of NPs on the performance of biological nitrogen removal, phosphorus removal and microbial communities for the degradation of organic pollutants. We also advance our own views on the aspects that are incomplete and warrant further research.

2. Effects of NPs to microorganisms

The increasing use of metal and metal oxide nanomaterials leads to the discharge of nanomaterials into the environment. Many researchers have evaluated these materials' toxicity to microorganisms.¹⁰ The inhibitory effect of Ag NPs and Au NPs on microorganisms in a modern WWTP was reported using realistic activated sludge.¹¹ This research indicated that 100 mg L⁻¹ Ag NPs is intermediately toxic to ordinary heterotrophic organisms (OHO) (by 33% of inhibition), while Au NPs show no or very low toxicity to OHO, ammonia oxidizing bacteria (AOB), and anaerobic biomass at the concentration of 90 mg L⁻¹. At low concentrations (<5 mg L⁻¹), Ag NPs do not affect anaerobic degradation, resulting in functional redundancy built within the microbial community.¹² In addition, the toxicity of Ag NPs to different microorganisms greatly differs, being the lowest (0.1 mg L⁻¹) for crustaceans and algae, and the highest (26 mg L⁻¹) for mammalian cells.¹³ The toxicity of Ag NPs is also affected by coexisting substances, e.g., sulphide. The presence of sulphide reduces the toxicity of Ag NPs to *C. riparius*, because sulphide can decrease the release of Ag⁺ from Ag NPs.¹⁴

ZnO NPs toxicity is related to the species of bacteria. 10 mg L⁻¹ ZnO NPs can cause significant growth inhibition (up to 90%) in *Bacillus subtilis* but only cause 22% of the growth inhibition in *Escherichia coli* (*E. coli*).¹⁵ Moreover, ZnO NPs can induce the production of oxidative stress, causing cell

membrane damage and cytoplasmic leakage owing to lipid peroxidation (LPO) at concentrations higher than 1.3×10^{-3} M (105 mg L⁻¹) in a liquid culture medium.¹⁶ In addition, ZnO NPs can also cause microbial biomass reduction, bacterial community shift and diversity decline in bacterial communities at 60 mg L⁻¹ in activated sludge in an SBR.¹⁷

TiO₂ NPs have been detected in wastewater and sewage sludge. Even though the predicted concentration of TiO₂ NPs is at the µg L⁻¹ level¹⁸ in the aquatic environment, their release into the environment increases because of their increasing production and application. Most results indicate that TiO₂ NPs do not have a strong toxicity to microorganisms, even at the concentration of 500 mg L⁻¹ in short-term treatments.¹⁹ TiO₂ NPs also do not show significant toxicity²⁰⁻²² at concentrations of <50 mg L⁻¹ in long-term treatments.

The toxicity of CuO NPs to some bacteria has been reported in recent years. CuO NPs (2.5 mg L⁻¹) can decrease the total nitrogen (TN) removal rate by 22.4%, but the removal can be recovered under certain conditions.²³ CuO NPs exhibit high toxicity to *Escherichia coli* at the concentration of 10 mg L⁻¹, but fulvic acid (FA) can reduce the toxicity significantly.²⁴ Additionally, the exposure to 50 mg L⁻¹ CuO NPs results in a significant inhibition to the respiration activity of the bacteria of biofilms.²³

Only a limited number of reports are found examining the toxicity of Al₂O₃ NPs to certain organisms. Specifically, 20 mg L⁻¹ Al₂O₃ NPs can induce a mortality rate of 57% in *Bacillus subtilis*, 36% in *E. coli*, and 70% in *Pseudomonas fluorescens* using 1 g L⁻¹ NaCl as the experimental medium,²⁵ while in a nutrient enriched test medium, Al₂O₃ NPs can create an antimicrobial property towards *Escherichia coli* and exhibit a nominal growth inhibitory effect only at very high concentrations (>1 g L⁻¹).²⁶ For a dominant bacteria, *Bacillus licheniformis*, isolated from freshwater (lake water), exposure to 1 mg L⁻¹ Al₂O₃ NPs for 2 h can cause a decrease in cell viability by 17%.²⁷ Al₂O₃ NPs present mild-level toxic effects on the growth of bacteria only at very high concentrations, which results from surface charge interactions between the bacteria and Al₂O₃ NPs. The ROS scavenging properties of Al₂O₃ NPs prevent cell wall disruption and drastic antimicrobial action. In addition, Al₂O₃ NPs can significantly inhibit *Escherichia coli* growth and prevent the adsorption of Al₂O₃ NPs onto the cell surface.²⁸ The intracellular Al₂O₃ NPs can interact with cellular biomolecules and cause adverse effects, eventually triggering cell death.

SiO₂ nanomaterials are widely used in domestic products and the potential toxicity to model organisms has been reported in zebrafish embryos, lung epithelial cells and other cell lines.^{29,30} Some studies have explored the potential toxicity of SiO₂ NPs to bacteria. One study reports that SiO₂ NPs can kill 40% of *Bacillus subtilis*, 58% of *Escherichia coli*, and 70% of *Pseudomonas fluorescens* in experimental culture mediums when the exposure concentration reaches 20 mg L⁻¹.²⁵ At the concentrations of 0.05 to 1.0 g L⁻¹, SiO₂ NPs have different effects on the antiradical activity of the culture medium (CM) of *Bacillus subtilis* IMV B-7023, and low doses of SiO₂ NPs can stimulate the reducing power of the CM of bacteria and strongly inhibit it in consequence.³¹ By comparison, SiO₂ NPs have less



power than Ag NPs against the oral pathogenic species *Streptococcus mutans*.³²

Similar to magnetic nanoparticles (MNPs), nano zero-valent iron (nZVI) and magnetite (Fe₃O₄) are widely used in water treatment. These applications make MNPs likely to enter the environment, and then get into WWTPs.³³ nZVI can penetrate cellular membranes and get into living cells,³⁴ which results in the accumulation of NPs in living organisms. Two other toxicological studies^{35,36} showed that nZVI can rapidly inactivate *Escherichia coli*, *Pseudomonas fluorescens* and *Bacillus subtilis* var. *niger*, but *Aspergillus versicolor* is not affected by them. Some microorganisms can be positively affected by nZVI. For example, methanogens are significantly stimulated by nZVI, and methane production is increased while dechlorinating organisms are inhibited.³⁷

There are two types of carbon nanotubes (CNTs), namely, multi-walled carbon nanotubes (MWCNTs) and single-walled carbon nanotubes (SWCNTs). The microbial toxicity of CNTs has been demonstrated under different conditions. The toxicity of CNTs to microorganisms is believed to be related to physical properties (length and diameter), concentration, and types of functionalization.^{38–41} It was reported that *Staphylococcus warneri* and *Staphylococcus aureus* are not able to grow over MWCNTs films.⁴² Both SWCNTs and MWCNTs at concentrations of 5 mg L⁻¹ are demonstrated to have antimicrobial activity towards *Escherichia coli*.⁴³ Long SWCNTs (5–30 μm) can cause significantly greater reductions in the viability of entrapped cells of *Escherichia coli* than short SWCNTs at concentrations of 5–200 mg L⁻¹.⁴⁴ When a phenol wastewater treatment system is exposed to SWCNTs (2.5 g L⁻¹) over an extended period, the structure, richness and diversity of microbial communities are shifted.⁴⁵ The effect of MWCNTs on upflow anaerobic sludge blanket (UASB) microflora shows that the reduction in a colony-forming unit (CFU) reaches 29% and 58% at MWCNTs concentrations of 1 and 100 mg L⁻¹, respectively.⁴⁶

In summary, most of the NPs mentioned above are toxic to microorganisms at certain concentrations, which makes it necessary to identify the possible adverse effects of them on the microbial activities of activated sludge in wastewater treatment plants.

3. Fate and transformation of NPs during the biological treatment of municipal wastewater

NPs released from different nanomaterials are finding their way to wastewater treatment tanks and end up in wastewater sludge.^{47–49} It is essential to investigate the fate and transformation of NPs to evaluate the potential environmental impacts on wastewater treatment systems. Many research findings indicate that for the removal of NPs, such as Ag NPs, Cu NPs, ZnO NPs, TiO₂ NPs, CuO NPs and CeO₂ NPs, from wastewater, multiple mechanisms are involved, including the physical entrapment of NPs into the floc, active cellular uptake, and binding to extracellular polymers substances (EPS) or

bacteria surfaces.^{50,51} Regardless of the mechanisms of removal, most NPs can be removed from suspension with the primary and/or secondary solids that are settled, wasted, and dewatered to form biosolids. Namely, only a very small fraction of NPs are discharged with the treated effluent.⁵² For example, it is estimated that only 6% of CeO₂ NPs are transmitted through a WWTP,⁵³ and 95% of Cu NPs are removed from wastewater.⁵⁴ The predominant mechanisms of copper removal appear to be aggregation and settling (Cu NPs), rather than biosorption. In addition, ZnO NPs are also efficiently removed from wastewater, with an insignificant portion being released into the environment through the treated effluent.⁵⁵ A large percentage of ZnO NPs settle out into the sludge. Under wastewater conditions, most NPs aggregate and are attached to the biomass.

Silver can bind strongly to sulphur (both organic and inorganic) in WWTPs, and most Ag NPs are concentrated and then removed.⁵⁶ Additionally, most of the Ag NPs in both effluent and sludge are present in the form of Ag₂S.⁵⁷ The almost complete removal of citrate-coated Ag NPs is achieved from a simulated sequencing batch reactor (SBR).⁵⁸ Ag NPs are efficiently transported into sewer systems without substantial losses to the sewer biofilm.⁵⁹ Therefore, Ag NPs can be efficiently removed from wastewater treatment systems.

Raw sewage titanium concentrations range from 181 to 1233 μg L⁻¹ in 10 representative WWTPs in the United States. The WWTPs can remove more than 96% of the influent TiO₂ NPs, and the effluent titanium concentrations are less than 25 μg L⁻¹ in all WWTPs.⁶⁰

In many cases, >90% of NPs from sewage appear to be removed, but poor removal efficiencies of SiO₂ NPs have been reported. For example, the flocculation and sedimentation processes used in primary wastewater treatment are shown to be ineffective in removing 56 nm SiO₂ NPs at 2470 mg L⁻¹ due to their stability and slow settling rate.⁶¹ In another study,⁶² 96% of SiO₂ NPs was effectively removed from the wastewater during the first 6 d, while the concentration of SiO₂ NPs in the effluent gradually increased afterwards, and the NPs discharge was as high as 65% of the input after 30 d of SiO₂ NPs dosing. The poor removal of the SiO₂ NPs is related to the high colloidal stability of the SiO₂ NPs in the wastewater and their limited propensity to biosorption.

There are many environmental factors affecting the migration and transformation of NPs. Dissolution, aggregation and agglomeration are the main factors affecting the state of metal NPs in suspensions.⁶³ Additionally, the salinity also affects the activated sludge and its microorganisms.⁶⁴ These processes will consequently impact the bioavailability, uptake, and toxicity of NPs.^{65,66} Various properties of the exposure media can influence the dissolution and aggregation of metal NPs, e.g. ionic strength, pH and the presence of natural organic matters (NOM).⁶⁷

The release of Zn²⁺ from ZnO NPs in wastewater is more significant under acidic conditions and low ionic strength, but under alkaline conditions, most of the ZnO NPs show a strong affinity for the sewage sludge rather than dissolved or dispersed in the filtrate.^{68,69} It was found that the dissolution or ion release of Cu NPs occurs in wastewater treatment systems.⁷⁰ Ag NPs can



Table 1 Summary of fate and transformation of some NPs during the biological treatment of municipal wastewater

NPs	Transformation	Removal efficiency	References
Ag NPs	Release Ag ⁺ , Cu ²⁺ , Zn ²⁺	High removal efficiency for NPs	50, 51, 54–59, 63, 64, 68, 70–72
Cu NPs		Low removal efficiency for released ions	
ZnO NPs	No transformation found	High removal efficiency	50, 51–54, 60, 65, 67
TiO ₂ NPs			
CuO NPs			
CeO ₂ NPs			
SiO ₂ NPs	No transformation found	Low removal efficiency	61, 62

be transformed into Ag⁺ under certain conditions, which were found to be more toxic than Ag NPs to the microbial communities in activated sludge and the nitrification process. However, DOM can significantly stabilize Ag NPs.⁷¹ Sulphidation can lead to Ag NPs and ZnO NPs aggregation.⁷² ZnO NPs are slightly soluble and can dissolve to form Zn²⁺ under aqueous conditions. This finding suggests that the released Zn²⁺ is responsible for the toxicity of ZnO NPs to *Mycobacterium smegmatis*⁷³ and *Pseudokirchneriella subcapitata*.⁷⁴ However, *Shewanella oneidensis* MR-1 and *Escherichia coli* are insensitive to the released Zn²⁺ even though the concentrations of ZnO NPs are higher than 40 mg L⁻¹.⁷³

Overall, the data collected indicate that biological WWTPs operated using activated sludge can effectively remove most types of NPs from wastewaters. Such NPs as Ag NPs, ZnO NPs, and TiO₂ NPs can be efficiently removed through their interaction with biomass. Ag NPs, Cu NPs and ZnO NPs would be partially transformed into Ag⁺, Cu²⁺ and Zn²⁺, respectively, under certain conditions. The presence of both EPS and ionic compounds can effectively enhance the NPs removal. The instability of the NPs resulting from the ionic strength in sewage and the entrapment of NPs by EPS play important roles in NPs removal by activated sludge. Table 1 is a summary of the fate and transformation of NPs during the biological treatment of municipal wastewater.

4. Effects of NPs on performances of biological treatment of municipal wastewater

4.1. Effects of NPs on nitrification and denitrification

Many studies have been conducted on the effects of Ag NPs on functional microorganisms and microbial communities in biological wastewater treatment processes. For example, among nitrogen-cycling bacteria, the nitrifier *Nitrosomonas europaea* is more sensitive to Ag NPs when the concentration is higher than 1.44 mg L⁻¹ compared to the denitrifier *Pseudomonas stutzeri* and the nitrogen fixer *Azotobacter vinelandii*.⁷⁵ Ag NPs at concentrations >2 mg L⁻¹ can induce damage of the cell wall and membrane of *Nitrosomonas europaea*, and disorganize the nucleoids.⁷⁶ In addition, Ag NPs can inhibit important protein functions, including biosynthesis, energy production, gene expression and nitrification in the test media at 5 mg L⁻¹.⁷⁷ García *et al.*¹¹ reported the inhibitory effects of Ag NPs on the

activity of microbial communities in activated sludge systems through respiration tests and biogas-production analysis. Ag NPs at the concentration of 0.1 mg mL⁻¹ cause an intermediate inhibition in biogas production (within 33–50%). In biofilm systems, the original wastewater biofilms are highly tolerant to Ag NPs. With the application of 200 mg L⁻¹ Ag NPs, the reduction in biofilm bacteria measured by heterotrophic plate counts is insignificant after 24 h. However, after the removal of loosely bound EPS, the viability of wastewater biofilms is reduced when treated under the same conditions.⁷⁸

Most of the results indicate that low concentrations of Ag NPs (*e.g.*, <2 mg L⁻¹) do not have significant effects on nitrogen removal in activated sludge systems. However, Ag NPs only have a mild impact on NH₄⁺-N removal, but the denitrification-related species are inhibited by 0.5 mg L⁻¹ Ag NPs, including *Diaphorobacter* species, *Thauera* species and those in the *Sphaerotilus–Leptothrix* group.⁷⁹ Ag NPs at concentrations >2 mg L⁻¹ disrupt ammonia removal in activated sludge in an SBR, while Ag-ligand formation and microbial adaptation (chloride or sulphide groups) can reduce their overall impacts.⁸⁰ When the concentration reaches 2 mg L⁻¹ of total Ag, biological nutrient removal can be disrupted, whereas Ag NPs do not exhibit a significant effect on ammonia removal during a 2 mg L⁻¹ spike. However, nitrification recovers quickly, especially at lower concentrations of Ag NPs. In addition, Doolette *et al.*⁸¹ also suggested that the nitrification rate is not affected by Ag NPs at the concentration of 2.5 mg L⁻¹ in an SBR. Ag NPs are found to be transformed into Ag₂S in activated sludge, and Ag₂S has no adverse effects on nitrification. However, a change in niche populations occur in aerobic sludge, indicating Ag NPs can negatively affect sub-dominant wastewater microbial communities.

Additionally,⁸² the continuous loading of Ag NPs (0.1 mg L⁻¹) for more than 60 d does not show adverse impacts on bacterial activity in a membrane bioreactor (MBR) activated sludge system. The nitrifying bacterial community structure is stable after long-term exposure to Ag NPs, while an abundance of the silver resistance gene *silE* in the MBR is increased after 41 d exposure, and then decreased with continuous exposure. These two results above indicate that activated sludge can effectively reduce the toxicity of Ag NPs by microbial adaptation and adsorbing or precipitating Ag NPs.

In summary, low concentrations of Ag NPs (<2 mg L⁻¹) mostly do not have significant effects on nitrification and denitrification in conventional activated sludge systems, while



Ag NPs at high concentrations ($>2 \text{ mg L}^{-1}$) can mostly inhibit nitrogen removal with long-term exposure.

Compared with Ag NPs, only a small number of studies have been conducted on the impact of Cu NPs on activated sludge. At concentrations from 0.1 to 10 mg L^{-1} , TN removal is promoted (improved from 60.6% to $>70\%$) and N_2O generation is reduced, but ammonia removal is not affected.⁸³ It was found that most Cu NPs are absorbed onto activated sludge, but the activated sludge surface is not damaged. The released Cu^{2+} is the main cause of TN removal improvement and N_2O reduction.

It was also found that the activities of ammonia mono-oxygenase and nitrite oxidoreductase are not affected by Cu NPs, whereas the activities of denitrification enzymes are increased, and Cu NPs increase the number of denitrifiers but decrease nitrite accumulation. All these results are in correspondence with the enhancement of TN removal and the decrease in N_2O production. Overall, Cu NPs do not show much of a negative impact on biological nitrogen removal. Short-term exposure to Cu NPs causes slight toxicity to ammonia oxidizing bacteria, but TN removal is promoted and N_2O generation is reduced by long-term exposure at certain concentrations of Cu NPs.

Several studies reveal the impacts of ZnO NPs on nitrification and denitrification. For example, short-term exposure to ZnO NPs can decrease TN removal from 81.5% to 75.6% and 70.8%, respectively, at the concentrations of 10 and 50 mg L^{-1} .⁸⁴ It was suggested that the inhibition of nitrogen removal mainly results from the release of Zn^{2+} from ZnO NPs dissolution and the increase of ROS production, which decreases nitrate reductase (NAR) activity. Similarly, the denitrification-related species were inhibited by 5 mg L^{-1} ZnO NPs, including *Diaphorobacter* species, *Thauera* species and those in the *Sphaerotilus-Leptothrix* group, although the $\text{NH}_4^+\text{-N}$ removal efficiency was either slightly reduced or not reduced at all.⁸⁵ In addition, ZnO NPs slightly decrease TN removal efficiency and reduce $\text{NH}_4^+\text{-N}$ removal by inhibiting the respiration of nitrifying microorganisms.⁸⁶ Overall, ZnO NPs affect TN removal efficiency by inhibiting the activity of NAR and the denitrification related to bacteria.

Several reports have described the impact of TiO_2 NPs on nitrification and denitrification. The concentrations of 1 and 50 mg L^{-1} TiO_2 NPs do not have acute effects on wastewater nitrogen removal after short-term exposure (1 d) in an anaerobic-low dissolved oxygen SBR, whereas 50 mg L^{-1} TiO_2 NPs significantly decreases TN removal efficiency by 56% after 70 d exposure. Mechanism analysis reveals that TiO_2 NPs at 50 mg L^{-1} obviously reduces the abundance of nitrifying bacteria and the diversity of microbial communities in activated sludge. In particular, ammonia-oxidizing bacteria are greatly decreased after long-term exposure to 50 mg L^{-1} TiO_2 NPs, resulting from the serious deterioration of ammonia oxidation. In addition, TiO_2 NPs at 50 mg L^{-1} can inhibit the activities of ammonia mono-oxygenase (AMO) and nitrite oxidoreductase (NOR) after long-term exposure.⁸⁷ TiO_2 NPs reduce the microbial diversity in activated sludge at concentrations from 100 mg L^{-1} to 200 mg L^{-1} , and the abundance of denitrifying bacteria is obviously decreased, which explains the inhibition.

Summarily, exposure time is one of the important factors of the adverse effects of TiO_2 NPs. Low concentrations (e.g. $<50 \text{ mg L}^{-1}$) of TiO_2 NPs do not have acute effects on wastewater nitrogen removal after short-term exposure, but high concentrations (e.g. $100\text{--}200 \text{ mg L}^{-1}$) of TiO_2 NPs can markedly decrease TN removal efficiency. Different from short-term exposure, 50 mg L^{-1} TiO_2 NPs can significantly decrease TN removal efficiency after long-term exposure. TiO_2 NPs inhibit both nitrifying bacteria and denitrifying bacteria, as well as AMO and NOR.

Only a small number of reports are found on the impact of Al_2O_3 NPs on wastewater nitrogen removal. Most of the Al_2O_3 NPs can adsorb onto activated sludge, but Al_2O_3 NPs have no adverse effects on the surface integrity and viability of activated sludge.⁸⁸ Additionally, short-term exposure to Al_2O_3 NPs does not present adverse impacts on wastewater nitrification and denitrification. However, the long-term exposure to 50 mg L^{-1} Al_2O_3 NPs can decrease the TN removal efficiency by 17.9%, owing to a suppressed denitrification process. The abundance of denitrifying bacteria is also decreased, and the activities of key denitrifying enzymes, such as nitrite reductase (NIR), are inhibited. Therefore, only long-term exposure to Al_2O_3 NPs has adverse impacts on wastewater nitrification and denitrification.

The acute and chronic effect of SiO_2 NPs on the nutrient removal performance of activated sludge was reported. Zheng *et al.*⁸⁹ noted that the presence of SiO_2 NPs at an environmentally relevant concentration (e.g. 1 mg L^{-1}) caused no adverse acute and chronic effects on sludge viability and wastewater nitrogen removal. Nevertheless, long-term exposure to 50 mg L^{-1} SiO_2 NPs can depress the TN removal efficiency by 18% after 70 d exposure, resulting from the declined activities of denitrifying enzymes and NIR. Further research revealed that the microbial community structure is changed, and the abundance of denitrifying bacteria is reduced after chronic exposure. In summary, SiO_2 NPs at low concentrations cause no significant adverse effects on nitrogen removal, but high concentrations of SiO_2 NPs affect TN removal efficiency.

We identified two reports regarding the most widely used magnetic NPs (nZVI, magnetite (Fe_3O_4) and maghemite ($\gamma\text{-Fe}_2\text{O}_3$) NPs). Wu *et al.*⁹⁰ examined the impacts of nZVI on nitrogen removal under continuous aerobic/anaerobic conditions in an activated sludge system and found no measurable effect of nZVI on nitrogen removal at the concentrations of 50 mg L^{-1} and below. However, 200 mg L^{-1} of nZVI inhibited $\text{NH}_4^+\text{-N}$ removal. The microbial activities were inhibited on exposure to nZVI; the adenosine triphosphate (ATP) content was decreased by approximately 13%, 31% and 43% at the nZVI concentrations of 20, 50, and 200 mg L^{-1} , respectively; and ROS production was increased. Lower concentrations of nZVI (20 and 50 mg L^{-1}) boosted the microbial activity, whereas 200 mg L^{-1} nZVI caused an approximate 19% depression in dehydrogenase activity. In addition, the acute exposure to $50\text{--}200 \text{ mg L}^{-1}$ Fe_3O_4 NPs can decrease TN removal efficiencies, resulting from the inhibition of nitrifying bacteria and the secretion of EPS.⁹¹ In contrast, chronic exposure to 50 mg L^{-1} magnetic NPs can clearly improve TN removal, resulting from the increased abundance of nitrifying bacteria and the improved activities of



Table 2 Summary of effects of NPs on biological treatment of municipal wastewater

Types of effects	NPs	Comments	Concentrations	References
Effects of NPs on nitrification and denitrification	Ag NPs	<i>Nitrosomonas europaea</i> is more sensitive than denitrifier <i>Pseudomonas stutzeri</i> and the nitrogen fixer <i>Azotobacter vinelandii</i>	>1.44 mg L ⁻¹	73
		Cause damage of cell wall and the membrane of <i>Nitrosomonas europaea</i> and make the nucleoids disorganized; disrupt ammonia removal	>2 mg L ⁻¹	74, 78
		Inhibit important protein functions, including biosynthesis, energy production, gene expression and nitrification	5 mg L ⁻¹	75
		Cause an intermediate inhibition in biogas production (within 33–50%)	0.1 mg mL ⁻¹	10
		Activated sludge can reduce Ag NPs toxicity	>2 mg L ⁻¹	78, 79
	Cu NPs	TN removal is promoted and N ₂ O generation is reduced; ammonia removal is not affected	0.1–10 mg L ⁻¹	81
	ZnO NPs	Decreases TN removal Denitrification species are inhibited; slightly decrease TN removal	10 and 50 mg L ⁻¹ 5 mg L ⁻¹	82 83, 84
	TiO ₂ NPs	Reduce the diversity of microbial community in activated sludge the abundance of nitrifying bacteria	>50 mg L ⁻¹ (long-term); 100–200 mg L ⁻¹ (short-term)	21, 85
	Al ₂ O ₃ NPs	No adverse effects on TN removal efficiency on (short-term exposure); decrease TN removal efficiency (long-term exposure)	50 mg L ⁻¹ ; 50 mg L ⁻¹	86
	SiO ₂ NPs	No adverse acute and chronic effects on nitrogen removal; depresses TN removal (long-term exposure)	1 mg L ⁻¹ ; 50 mg L ⁻¹	87
nZVI	No measurable effect on nitrogen removal; inhibit NH ₄ ⁺ -N removal	≤50 mg L ⁻¹ ; 200 mg L ⁻¹	88	
Fe ₃ O ₄ and Fe ₂ O ₃ NPs	Decreases TN removal (short-term exposure)	50–200 mg L ⁻¹	89	
Effects of NPs on biological phosphorus removal	Ag NPs	No significant effect on biological phosphorus removal; Ag ⁺ shows stronger toxicity (>1 mg L ⁻¹)	<5 mg L ⁻¹	85, 90, 91
	ZnO NPs	Inhibition of phosphorus removal is due to the release of zinc ions	<50 mg L ⁻¹	82
	TiO ₂ NPs	No acute effects on phosphorus removal	<50 mg L ⁻¹	22
	Al ₂ O ₃ NPs	Induce marginal influences on wastewater phosphorus removal	<50 mg L ⁻¹	86
	SiO ₂ NPs	Phosphorus removal is insensitive	<50 mg L ⁻¹	87
	Cu NPs	Phosphorus removal is unaffected	0.1–10 mg L ⁻¹	81
	nZVI	Improves phosphorous removal by boosting the microbial activity	20 mg L ⁻¹	88
Effects of NPs on organic-degradation microbial community	Ag NPs	Impact the activated sludge microbial community and cell culturability	1 mg L ⁻¹	92
		Methane production are not affected	2.9 g per (kg TS)	79
		COD removal is moderately affected and can be recovered significantly	0.2 and 2 mg L ⁻¹	77
		COD removal will diminish as the increased addition of Ag NPs	50 mg L ⁻¹	93
	Negligible impact on anaerobic digestion and methanogenic assemblages	≤40 mg L ⁻¹	49	
	ZnO NPs	Highly toxic and rapidly disrupt the methanogenic activity	34.5 mg Zn per L	94
	Inhibit the microbial activities in the outer layer of biofilms	50 mg L ⁻¹	95	
EPS of AGS and the methane production are not significantly affected; EPS of AGS and the methane production are decreased	<50 mg g ⁻¹ -TSS; 200 mg g ⁻¹ -TSS	96		



Table 2 (Contd.)

Types of effects	NPs	Comments	Concentrations	References
	CuO NPs	Marked negative effect on the performance of methanogenesis in UASB reactor	1.4 mg Cu per L	99
	Cu NPs	Cause inhibition of acetoclastic methanogens but not H ₂ -utilizing methanogens	62 mg L ⁻¹	97
		No significant effect on sludge fermentative volatile fatty acids (VFA)	50 mg L ⁻¹	100
	SiO ₂ NPs	Affect the biological activity in activated sludge through the inhibition of total oxygen uptake	>50 mg L ⁻¹	101

NAR and NIR. The release of iron ions from the dissolution of magnetic NPs may also be responsible for the increase in bacteria activities. In summary, magnetic NPs have little adverse effects on nitrification and denitrification at ≤ 50 mg L⁻¹ of magnetic NPs, whereas high concentrations of NPs can cause a depression in dehydrogenase activity.

Overall, exposure time (short-term or long-term) is one of the important factors affecting the adverse effects of many types of NPs. The adverse effects of many types of NPs are of concern for the nitrification and denitrification in activated sludge. Most NPs at certain concentrations exhibit some adverse effects on nitrogen removal by inhibiting the key enzymes of nitrifying and denitrifying bacteria. Some adverse effects result from the release of ions from dissolved NPs.

4.2. Effects of NPs on biological phosphorus removal

Some studies have been reported on the effects of NPs on biological phosphorus removal. It was reported⁹² that Ag NPs (< 5 mg L⁻¹) do not present obvious effects on biological phosphorus removal in both acute and chronic exposure. However, the removal rate of phosphorus was decreased to 48.8% at 1 mg L⁻¹ of Ag⁺ at short-term exposure, and almost zero phosphorus removal was achieved when the concentration of Ag⁺ was greater than 2 mg L⁻¹. Ag⁺ shows stronger toxicity to polyphosphate accumulating organisms (PAOs) than glycogen accumulating organisms (GAOs). In addition, the decrease in anaerobic phosphorus release by Ag⁺ is caused by the inhibition of the activities of adenylate kinase and exopolyphosphatase (PPX), whereas the decrease in aerobic phosphorus uptake is due to the suppression of energy generation for phosphorus uptake. A comparison was conducted on the effects of Ag NPs and Ag⁺ at concentrations from 1 to 5 mg L⁻¹ on the performance of enhanced biological phosphorus removal (EBPR). Ag NPs showed no obvious effect on EBPR at both acute and chronic exposure,⁹³ whereas phosphorus removal was decreased by a sudden increase in Ag⁺ concentration and was gradually recovered after long-term culture. The mechanism study showed that the transformation of polyhydroxyalkanoates (PHA) and glycogen, and the activity of PPX decline when Ag⁺

was suddenly increased, but they were recovered after long-term culture.⁸²

The effects of NPs such as ZnO, TiO₂, Al₂O₃, and SiO₂ NPs on biological phosphorus removal are of concern. ZnO NPs at concentrations of 10 and 50 mg L⁻¹ can inhibit phosphorus removal and the inhibition result from the release of zinc ions from ZnO NPs dissolution, as well as ROS production, causing an inhibitory effect on PAOs and decreased activities of PPX and polyphosphate kinase (PPK).⁸⁴ It was reported that 1 and 50 mg L⁻¹ TiO₂ NPs do not exhibit significant impacts on phosphorus removal at both acute exposures (1 d) and chronic exposure (70 d).²² Short-term exposure to 1 and 50 mg L⁻¹ Al₂O₃ NPs can induce marginal influence on wastewater phosphorus removal.⁸⁸ In addition, transformations of intracellular PHA and glycogen were not significantly changed in the presence of 1 and 50 mg L⁻¹ Al₂O₃ NPs.

Zheng *et al.* reported that wastewater phosphorus removal is insensitive to 1 and 50 mg L⁻¹ SiO₂ NPs at both acute and chronic exposures because the important factors related to phosphorus removal, such as the activities of PPX and PPK and the intracellular transformations of PHA and glycogen, were not obviously changed.⁸⁹ As illustrated above,⁸³ the presence of Cu NPs in activated sludge affects the nitrogen removal by increasing the activity of NAR and NIR while the effects of Cu NPs are insignificant on the activities of AMO, NOR, PPX, and PPK. This means that phosphorus removal is unaffected by Cu NPs.⁸³ As illustrated above,⁹⁰ nZVI does not exhibit any measurable influence on nitrogen removal at 50 mg L⁻¹ and below, whereas the addition of nZVI at 10–20 mg L⁻¹ significantly improves phosphorus removal by boosting microbial activity.⁹⁰

In summary, the inhibition of phosphorus removal induced by ZnO NPs is due to the release of zinc ions from ZnO NPs dissolution, as well as ROS production. Most other NPs, such as Cu NPs, TiO₂ NPs, Al₂O₃ NPs and SiO₂ NPs, do not present significantly adverse effects on biological phosphorus removal at the concentrations of 50 mg L⁻¹ and below, because they have no significant impacts on the activities of PPX and PPK, and the transformations of intracellular PHA and glycogen. Low



concentrations of nZVI (10–20 mg L⁻¹) can improve phosphorous removal.

4.3. Effects of NPs on organic-degradation microbial communities

Many studies have commenced investigations into the effects of NPs on activated sludge microbial communities. Ag NPs (1 mg L⁻¹) can affect activated sludge microbial communities and cell culturability depending on the physical structure of the activated sludge flocs, the spatial distribution of microorganisms in activated sludge flocs, and the community structures in the activated sludge.⁹⁴ Doolette *et al.*⁸¹ found methane production was not affected by Ag NPs (~3 g kg⁻¹-TS) in an anaerobic batch test. Ag NPs were found to be transformed into Ag₂S in activated sludge, and transformed Ag NPs have no adverse effects on methanogenesis. However, a change in niche populations was found, suggesting that Ag NPs may adversely impact sub-dominant wastewater microbial communities. In addition, Chen *et al.*⁷⁹ suggested that COD removal is moderately affected by low concentrations of Ag NPs. When Ag NPs were added at 0.2 and 2 mg L⁻¹, the removal percentages were decreased from 99% to approximately 80%, but COD removal was remarkably recovered 3 d later, and a COD removal efficiency of approximately 93% was achieved. Additionally, a remarkable change occurred in the microbial community structure, which can be interpreted as an improvement in recovery. However, it was reported that the COD removal rate will decrease with the increased addition of Ag NPs and that only 20% of COD is removed at an Ag NPs concentration of 50 mg L⁻¹.⁹⁵ Yang *et al.* suggested that Ag NPs at moderate concentrations (≤40 mg L⁻¹) have negligible effects on anaerobic digestion and methanogenic assemblages,⁴⁹ whereas 50 mg L⁻¹ Ag NPs significantly decreased the COD removal efficiency.

ZnO NPs at 34.5 mg L⁻¹ are highly toxic and rapidly disrupt the methanogenic activity during UASB operation, and the toxicity was more severe for acetoclastic methanogens compared with hydrogenotrophic methanogens.⁹⁶ In addition, sub-ppm levels of ZnO NPs present a remarkable effect over the long-term, as evidenced by the gradual decrease in the methanogenic activity of the biomass. This finding may be attributable to two factors: the enhanced toxicity with exposure time, and the accumulation of ZnO NPs during wastewater treatment that may increase the effective concentration of ZnO NPs in direct contact with the microorganisms in sludge. Also, 50 mg L⁻¹ ZnO NPs can inhibit the microbial activities only in the outer layer of biofilms, and bacteria present in the deeper parts of biofilms become even more active.⁹⁷ Methane production was not obviously affected by ZnO NPs at the concentrations of 10 and 50 mg g⁻¹-TSS, but it was decreased when ZnO NPs concentration was higher than 100 mg g⁻¹-TSS, because ZnO NPs caused a decrease of proteins in EPS by 69.6%.⁹⁸ Additionally, the C–O–C group of polysaccharides and carboxyl group of proteins in EPS were also changed. The average methane production was decreased by 43.5% at a ZnO NPs concentration of 200 mg g⁻¹-TSS. ZnO NPs can also cause severe inhibition of methanization.

Cu NPs can cause inhibition of acetoclastic methanogens, but not of H₂-utilizing methanogens at a Cu NPs concentration of 62 mg L⁻¹.⁹⁹ Therefore, ZnO NPs, CuO NPs and Cu NPs can induce inhibition of methane production during anaerobic biological wastewater processes. The production of ROS and the release of LDH showed that exposing the activated sludge to CuO NPs caused oxidative stress and damaged cell membranes in the sludge flocs and the production of EPS; in particular, polysaccharides were enhanced by 89.7% with exposure to 50 mg L⁻¹ CuO NPs.¹⁰⁰ Long-term exposure of the microorganisms retained in the sludge bed to CuO NPs (1.4 mg Cu per L) can cause significant inhibition of *Acetoclastic methanogenesis*, which eventually leads to the complete disruption of the bioreactor performance.¹⁰¹ The inhibition is likely induced by both CuO NPs and released copper ions.

The addition of Cu NPs to biological wastewater treatment systems presented no significant effects on sludge fermentative volatile fatty acids (VFA), the preferred carbon source for wastewater biological nutrient removal production, as the sludge solubilization was increased, and the acidification was decreased.¹⁰² When Cu NPs are directly added to the anaerobic fermentation reactor, both the hydrolysis and acidification are inhibited at a Cu NPs concentration of 50 mg L⁻¹, resulting in an ultimate VFA production much less than that of the control by approximately 45% after 10 d exposure.

The adverse impacts of SiO₂ NPs on activated sludge was studied using the activated sludge respiration inhibition test and it was suggested that SiO₂ NPs affect the biological activity in activated sludge through the inhibition of total oxygen uptake at concentrations higher than 50 mg L⁻¹.¹⁰³ Transmission Electron Microscopy (TEM) image analysis suggests some of the SiO₂ NPs are adsorbed on and/or apparently embedded in the microbial cell membrane.

Summarily, most NPs at certain concentrations have adverse effects on microorganisms for organic degradation in activated sludge (Table 2). After exposure, a degree of inhibition might occur to microbial communities and to the activities of certain key enzymes in activated sludge, subsequently resulting in the abnormal performance of activated sludge systems.

5. Conclusions and future directions

Many studies have been conducted on the fate of NPs, as well as their effects on biological wastewater treatment, and many achievements have been reported. It can be concluded from this review as follows:

(1) Most NPs, such as Ag, ZnO, CuO, Al₂O₃, SiO₂, CNTs and magnetic NPs, can cause varying degrees of damage to microorganisms at certain concentrations. Ag, Cu and ZnO NPs present relatively significant toxic effects compared with other NPs at similar exposure concentrations. In particular, TiO₂ NPs do not show strong toxicity to microorganisms at both short-term (even at 500 mg L⁻¹) and long-term exposures (<50 mg L⁻¹).

(2) WWTPs operated using activated sludge have the potential to remove most types of NPs, such as Ag, Cu, ZnO, CuO, and TiO₂, but not SiO₂, by interaction with biomass. Ag NPs, Cu NPs



and ZnO NPs can be partially transformed into Ag^+ , Cu^{2+} and Zn^{2+} , respectively, under certain conditions. The poor removal of SiO_2 NPs is due to the high colloidal stability of SiO_2 NPs in the wastewater and their limited propensity to biosorption.

(3) Most NPs have varying degrees of impacts on the performance of the biological treatment of wastewater, including nitrogen, phosphorus and organic pollutants removal. The impacts are dose-dependent under most conditions, and the exposure time (short-term or long-term) is also an important factor of adverse effects. (a) For nitrogen removal, Ag, CuO and ZnO NPs have slight or moderate effects on TN removal at relatively low concentrations; TiO_2 NPs show some inhibitory effects on nitrifying and denitrifying bacteria, and AMO and NOR; and Al_2O_3 and SiO_2 NPs have adverse impacts on nitrification and denitrification. (b) For phosphorus removal, Al_2O_3 , TiO_2 and SiO_2 NPs do not have significant adverse effects, but ZnO NPs can result in net phosphorus removal failure at certain concentrations. (c) For organics removal, Ag NPs can decrease COD removal. ZnO NPs and CuO NPs can inhibit methane production during anaerobic biological wastewater processes. On the whole, the impacts are dose-dependent under most conditions. Some of the impacts are due to the NPs themselves, and some of impacts result from released ions such as Ag^+ , Cu^{2+} , and Zn^{2+} .

Although many studies have been conducted in this field, the achievements are far from complete. For instance, as to Au NPs and CNTs, no reports are found on their effects on biological nitrogen and phosphorus removal. There are only a few studies found that report the effects of NPs on organics removal. NPs removed from sewage will accumulate in biosolids, and more research is required to understand the fate of NPs upon the ultimate disposal of biosolids. The effects of NPs on biosolids disposal process also need more research. In addition, long-term effects of NPs at low concentrations on the functional microorganisms need more concern.

Acknowledgements

The Project is sponsored by the Scientific Research Foundation for the Returned Overseas Chinese Scholars, State Education Ministry, China. This work is also supported by the Key Laboratory of Urban Water Resource and Environment of Harbin institute of technology, China (ES201608).

References

- 1 Y. Ju-Nam and J. R. Lead, Manufactured nanoparticles: An overview of their chemistry, interactions and potential environmental implications, *Sci. Total Environ.*, 2008, **400**, 396–414.
- 2 J. Wackerlig and R. Schirhagl, Applications of Molecularly Imprinted Polymer Nanoparticles and their Advances toward Industrial Use: A Review, *Anal. Chem.*, 2016, **88**(1), 250–261.
- 3 S. J. Klaine, P. J. J. Alvarez, G. E. Batley, T. F. Fernandes, R. D. Handy, D. Y. Lyon, S. Mahendra, M. J. McLaughlin and J. R. Lead, Nanomaterials in the environment: Behavior, fate, bioavailability, and effects, *Environ. Toxicol. Chem.*, 2008, **27**(9), 1825–1851.
- 4 S. K. Brar, M. Verma, R. D. Tyagi and R. Y. Surampalli, Engineered nanoparticles in wastewater and wastewater sludge – Evidence and impacts, *Waste Manag.*, 2010, **30**, 504–520.
- 5 V. K. Sharma, K. M. Siskova, R. Zboril and J. L. Gardea-Torresdey, Organic-coated silver nanoparticles in biological and environmental conditions: Fate, stability and toxicity, *Adv. Colloid Interface Sci.*, 2014, **204**, 15–34.
- 6 L. A. S. Bahari and H. Hamishehkar, The impact of variables on particle size of solid lipid nanoparticles and nanostructured lipid carriers: A comparative literature review, *Adv. Pharm. Bull.*, 2016, **6**(2), 143–151.
- 7 B. Song, J. Liu, X. L. Feng, *et al.*, A review on potential neurotoxicity of titanium dioxide nanoparticles, *Nanoscale Res. Lett.*, 2015, **10**, 342–358.
- 8 A. Lazareva and A. A. Keller, Estimating Potential Life Cycle Releases of Engineered Nanomaterials from Wastewater Treatment Plants, *ACS Sustainable Chem. Eng.*, 2014, **2**, 1656–1665.
- 9 A. A. Keller and A. Lazareva, Predicted Releases of Engineered Nanomaterials: From Global to Regional to Local, *Environ. Sci. Technol. Lett.*, 2014, **1**, 65–70.
- 10 L. S. Dorobantu, C. Fallone, A. J. Noble, J. Veinot, G. Ma, G. G. Goss and R. E. Burrell, Toxicity of silver nanoparticles against bacteria, yeast, and algae, *J. Nanopart. Res.*, 2015, **17**, 172–184.
- 11 A. García, L. Delgado, J. A. Torà, E. Casals, E. González, V. Puentes, X. Font, J. Carrera and A. Sánchez, Effect of cerium dioxide, titanium dioxide, silver, and gold nanoparticles on the activity of microbial communities intended in wastewater treatment, *J. Hazard. Mater.*, 2012, **199**–**200**, 64–72.
- 12 A. Gitipour, S. W. Thiel, K. G. Scheckel and T. Tolaymat, Anaerobic toxicity of cationic silver nanoparticles, *Sci. Total Environ.*, 2016, **557**–**558**, 363–368.
- 13 A. Ivask, I. Kurvet, K. Kasemets, *et al.*, Size-dependent toxicity of silver nanoparticles to bacteria, yeast, algae, crustaceans and mammalian cells *in vitro*, *PLoS One*, 2014, **9**(7), e102108.
- 14 S.-W. Lee, S.-Y. Park, Y. H. Kim, H. Im and J. Choi, *Sci. Total Environ.*, 2016, **553**, 565–573.
- 15 L. K. Adams, D. Y. Lyon and P. J. J. Alvarez, *Water Res.*, 2006, **40**, 3527–3532.
- 16 R. Brayner, R. Ferrari-Iliou, N. Brivois, S. Djediat, M. F. Benedetti and F. Fiévet, *Nano Lett.*, 2006, **6**, 866–870.
- 17 S. Wang, M. C. Gao, Z. L. She, D. Zheng, C. J. Jin, L. Guo, Y. G. Zhao, Z. W. Li and X. J. Wang, *Bioresour. Technol.*, 2016, **216**, 428–436.
- 18 F. Gottschalk, T. Sonderer, R. W. Scholz and B. Nowack, *Environ. Sci. Technol.*, 2009, **43**, 9216–9222.
- 19 L. K. Adams, D. Y. Lyon and P. J. J. Alvarez, *Water Res.*, 2006, **40**, 3527–3532.
- 20 T. J. Battin, F. V. D. Kammer, A. Weilhartner, S. Ottofuelling and T. Hofmann, *Environ. Sci. Technol.*, 2009, **43**, 8098–8104.



- 21 C. Supha, Y. Boonto, M. Jindakaraked, J. Ananpattarachai and P. Kajitvichyanukul, *Sci. Technol. Adv. Mater.*, 2015, **16**, 1–12.
- 22 X. Zheng, Y. Chen and R. Wu, *Environ. Sci. Technol.*, 2011, **45**, 7284–7290.
- 23 Y. L. Su, Y. G. Chen and X. Zheng, *Water Res.*, 2016, **99**, 216–224.
- 24 J. Zhao, Z. Wang, Y. Dai and B. Xing, *Water Res.*, 2013, **47**(12), 4169–4178.
- 25 W. Jiang, H. Mashayekhi and B. Xing, *Environ. Pollut.*, 2009, **157**, 1619–1625.
- 26 I. M. Sadiq, B. Chowdhury, N. Chandrasekaran and A. Mukherjee, *Biol. Med.*, 2009, **5**, 282–286.
- 27 S. Pakrashi, S. Dalai, D. Sabat, S. Singh, N. Chandrasekaran and A. Mukherjee, *Chem. Res. Toxicol.*, 2011, **24**, 1899–1904.
- 28 M. A. Ansari, H. M. Khan and A. A. Khan, *J. Appl. Microbiol.*, 2014, **116**(4), 772–783.
- 29 T. J. Brunner, P. Wick, P. Manser, P. Spohn, R. N. Grass, L. K. Limbach, A. Bruinink and W. J. Stark, *Environ. Sci. Technol.*, 2006, **40**, 4374–4381.
- 30 J. Chang, K. L. B. Chang, D. Hwang and Z. Kong, *Environ. Sci. Technol.*, 2007, **41**, 2064–2068.
- 31 I. O. Skorochod, A. O. Roy and I. K. Kurdish, *Nanoscale Res. Lett.*, 2016, **11**, 139–144.
- 32 A. Besinis, T. D. Peralta and R. D. Handy, *Nanotoxicology*, 2014, **8**(1), 1–16.
- 33 M. Stefaniuk, P. Oleszczuk and Y. Sik Ok, *Chem. Eng. J.*, 2016, **287**, 618–632.
- 34 M. P. Herrling, S. Lackner, O. Tatti, G. Guthausen, M. Delay, M. Franzreb and H. Horn, *Sci. Total Environ.*, 2016, **544**, 722–729.
- 35 C. Lee, J. Y. Kim, W. I. Lee, K. L. Nelson, J. Yoon and D. L. Sedlak, *Environ. Sci. Technol.*, 2008, **42**, 4927–4933.
- 36 M. Diao and M. Yao, *Water Res.*, 2009, **43**, 5243–5251.
- 37 Z. Xiu, Z. Jin, T. Li, S. Mahendra, G. V. Lowry and P. J. J. Alvarez, Iron particles on a mixed culture, *Bioresour. Technol.*, 2010, **101**, 1141–1146.
- 38 D. Kerfahi, B. M. Tripathi, D. Singh, H. Kim, S. Lee, J. Lee and J. M. Adams, *PLoS One*, 2015, **10**, e0123042.
- 39 P. Oleszczuk, I. Joško and B. Xing, *J. Hazard. Mater.*, 2011, **186**, 436–442.
- 40 D. Goyal, X. J. Zhang and J. N. Rooney-Varga, *Lett. Appl. Microbiol.*, 2010, **51**, 428–435.
- 41 L. A. Luongo and X. Zhang, *J. Hazard. Mater.*, 2010, **178**, 356–362.
- 42 R. J. Narayan, C. J. Berry and R. L. Brigmon, *Mater. Sci. Eng., B*, 2005, **123**, 123–129.
- 43 S. Kang, M. Herzberg, D. F. Rodrigues and M. Elimelech, *Langmuir*, 2008, **24**, 6409–6413.
- 44 T. T. A. Le, J. McEvoy and E. Khan, *Sci. Total Environ.*, 2016, **565**, 787–794.
- 45 Y. Y. Qu, X. W. Zhang, W. L. Shen, *et al.*, *Bioresour. Technol.*, 2016, **211**, 209–215.
- 46 T. Yadav, A. A. Mungray and A. K. Mungray, *Environ. Sci. Pollut. Res.*, 2016, **23**, 4063–4072.
- 47 H. Mu, Y. Chen and N. Xiao, *Bioresour. Technol.*, 2011, **102**, 10305–10311.
- 48 H. Mu and Y. Chen, *Water Res.*, 2011, **45**, 5612–5620.
- 49 Y. Yang, Q. Chen, J. D. Wall and Z. Hu, *Water Res.*, 2012, **46**, 1176–1184.
- 50 H. Park, H. Y. Kim, S. Cha, C. H. Ahn, J. Roh, S. Park, S. Kim, K. Choi, J. Yi, Y. Kim and J. Yoon, *Chemosphere*, 2013, **92**, 524–528.
- 51 E. F. C. Chauque, J. N. Zvimba, J. C. Ngila and N. Musee, *Phys. Chem. Earth*, 2014, **67–69**, 140–144.
- 52 P. K. Westerhoff, M. A. Kiser and K. Hristovski, *Environ. Eng. Sci.*, 2013, **30**, 109–117.
- 53 L. K. Limbach, R. Bereiter, E. Müller, R. Krebs, R. Gälli and W. J. Stark, *Environ. Sci. Technol.*, 2008, **42**, 5828–5833.
- 54 R. Ganesh, J. Smeraldi, T. Hosseini, L. Khatib, B. H. Olson and D. Rosso, *Environ. Sci. Technol.*, 2010, **44**, 7808–7813.
- 55 E. F. C. Chauque, J. N. Zvimba, J. C. Ngila and N. Musee, *Water SA*, 2016, **42**(1), 72–80.
- 56 C. Levard, E. M. Hotze, G. V. Lowry and G. E. Brown, *Environ. Sci. Technol.*, 2012, **46**, 6900–6914.
- 57 R. Kaegi, A. Voegelin, B. Sinnet, S. Zuleeg, H. Hagedorfer, M. Burkhardt and H. Siegrist, *Environ. Sci. Technol.*, 2011, **45**, 3902–3908.
- 58 L. Hou, K. Li, Y. Ding, Y. Li, J. Chen, X. Wu and X. Li, *Chemosphere*, 2012, **87**, 248–252.
- 59 R. Kaegi, A. Voegelin, C. Ort, B. Sinnet, B. Thalmann, J. Krismer, H. Hagedorfer, M. Elumelu and E. Mueller, *Water Res.*, 2013, **47**, 3866–3877.
- 60 P. Westerhoff, G. X. Song, K. Hristovski and M. A. Kiser, *J. Environ. Monit.*, 2011, **13**(5), 1195–1203.
- 61 H. P. Jarvie, H. Al-Obaidi, S. M. King, M. J. Bowes, M. J. Lawrence, A. F. Drake, M. A. Green and P. J. Dobson, *Environ. Sci. Technol.*, 2009, **43**, 8622–8628.
- 62 L. Otero-González, J. A. Field, I. A. C. Calderon, C. A. Aspinwall, F. Shadman, C. Zeng and R. Sierra-Alvarez, *Water Res.*, 2015, **77**, 170–178.
- 63 H. He, Y. Cheng, C. Yang, *et al.*, *J. Environ. Sci.*, 2017, **54**, 135–141.
- 64 H. He, Y. Chen, X. Li, *et al.*, *Int. Biodeterior. Biodegrad.*, 2017, **119**, 520–527.
- 65 S. K. Misra, A. Dybowska, D. Berhanu, S. N. Luoma and E. Valsami-Jones, *Sci. Total Environ.*, 2012, **438**, 225–232.
- 66 L. Otero-González, I. Barbero, J. A. Field, F. H. Shadman and R. Sierra-Alvarez, *Water Sci. Technol.*, 2014, **70**(9), 1533–1539.
- 67 N. M. Franklin, N. J. Rogers, S. C. Apte, G. E. Batley, G. E. Gadd and P. S. Casey, *Environ. Sci. Technol.*, 2007, **41**(24), 8484–8490.
- 68 E. Lombi, E. Donner, E. Tavakkoli, T. W. Turney, R. Naidu, B. W. Miller and K. G. Scheckel, *Environ. Sci. Technol.*, 2012, **46**, 9089–9096.
- 69 N. Musee, J. N. Zvimba, L. M. Schaefer, N. Nota, L. M. Sikhwihilu and M. Thwala, *J. Environ. Sci. Health, Part A: Toxic/Hazard. Subst. Environ. Eng.*, 2014, **49**, 59–66.
- 70 Y. Liu, J. Baas, W. J. G. M. Peijnenburg and M. G. Vijver, *Environ. Sci. Technol.*, 2016, **50**, 5328–5337.
- 71 S. Yu, Y. Yin, J. Chao, M. Shen and J. Liu, *Environ. Sci. Technol.*, 2014, **48**, 403–411.



- 72 R. Ma, C. Levard, F. M. Michel, G. E. Brown and G. V. Lowry, *Environ. Sci. Technol.*, 2013, **47**, 2527–2534.
- 73 B. Wu, Y. Wang, Y. Lee, A. Horst, Z. Wang, D. Chen, R. Sureshkumar and Y. J. Tang, *Environ. Sci. Technol.*, 2010, **44**, 1484–1489.
- 74 N. M. Franklin, N. J. Rogers, S. C. Apte, G. E. Batley, G. E. Gadd and P. S. Casey, *Environ. Sci. Technol.*, 2007, **41**, 8484–8490.
- 75 Y. Yang, J. Wang, Z. Xiu and P. J. J. Alvarez, *Environ. Toxicol. Chem.*, 2013, **32**(7), 1488–1494.
- 76 Z. Yuan, J. Li, L. Cui, B. Xu, H. Zhang and C. Yu, *Chemosphere*, 2013, **90**, 1404–1411.
- 77 T. S. Radniecki, D. P. Stankus, A. Neigh, J. A. Nason and L. Semprini, *Chemosphere*, 2011, **85**, 43–49.
- 78 Z. Sheng and Y. Liu, *Water Res.*, 2011, **45**, 6039–6050.
- 79 J. Chen, Y. Tang, Y. Li, Y. Nie, L. Hou, X. Li and X. Wu, *Chemosphere*, 2014, **104**, 141–148.
- 80 C. L. Alito and C. K. Gunsch, *Environ. Sci. Technol.*, 2014, **48**, 970–976.
- 81 C. L. Doolette, M. J. McLaughlin, J. K. Kirby, D. J. Batstone, H. H. Harris, H. Q. Ge and G. Comelis, *Chem. Cent. J.*, 2013, **7**, 1–18.
- 82 C. Zhang, Z. Liang and Z. Hu, *Water Res.*, 2014, **50**, 350–358.
- 83 Y. Chen, D. Wang, X. Zhu, X. Zheng and L. Feng, *Environ. Sci. Technol.*, 2012, **46**, 12452–12458.
- 84 X. Zheng, R. Wu and Y. Chen, *Environ. Sci. Technol.*, 2011, **45**, 2826–2832.
- 85 L. Hou, J. Xia, K. Li, J. Chen, X. Wu and X. Li, *Water Sci. Technol.*, 2013, **67**(2), 254–260.
- 86 S. T. Wang, S. P. Li, W. Q. Wang and H. You, *RSC Adv.*, 2015, **5**, 67335–67342.
- 87 D. Li, F. Cui, Z. Zhao, D. Liu, Y. Xu, H. Li and X. Yang, *Biodegradation*, 2014, **25**, 167–177.
- 88 Y. Chen, Y. Su, X. Zheng, H. Chen and H. Yang, *Water Res.*, 2012, **46**, 4379–4386.
- 89 X. Zheng, Y. Su and Y. Chen, *Environ. Sci. Technol.*, 2012, **46**, 7182–7188.
- 90 D. Wu, Y. Shen, A. Ding, Q. Mahmood, S. Liu and Q. Tu, *J. Hazard. Mater.*, 2013, **262**, 649–655.
- 91 S. Ni, J. Ni, N. Yang and J. Wang, *Bioresour. Technol.*, 2013, **143**, 555–561.
- 92 H. Chen, X. Zheng, Y. Chen and H. Mu, *RSC Adv.*, 2013, **3**, 9835–9842.
- 93 Y. Chen, H. Chen, X. Zheng and H. Mu, *J. Hazard. Mater.*, 2012, **239–240**, 88–94.
- 94 X. Sun, Z. Sheng and Y. Liu, *Sci. Total Environ.*, 2013, **443**, 828–835.
- 95 E. Jeong, W. Im, D. Kim, M. Kim, S. Kang, H. Shin and S. Chae, *J. Environ. Sci. Health, Part A: Toxic/Hazard. Subst. Environ. Eng.*, 2014, **49**, 685–693.
- 96 L. Otero-González, J. A. Field and R. Sierra-Alvarez, *J. Environ. Manage.*, 2014, **135**, 110–117.
- 97 J. Hou, L. Z. Miao, C. Wang and P. F. Wang, *J. Hazard. Mater.*, 2014, **276**, 164–170.
- 98 H. Mu, X. Zheng, Y. Chen, H. Chen and K. Liu, *Environ. Sci. Technol.*, 2012, **46**, 5997–6003.
- 99 J. Gonzalez-Estrella, R. Sierra-Alvarez and J. A. Field, *J. Hazard. Mater.*, 2013, **260**, 278–285.
- 100 J. Hou, L. Z. Miao, C. Wang, P. F. Wang, Y. H. Ao and B. W. Lv, *Bioresour. Technol.*, 2015, **176**, 65–70.
- 101 L. Otero-González, J. A. Field and R. Sierra-Alvarez, *Water Res.*, 2014, **58**, 160–168.
- 102 H. Chen, Y. Chen, X. Zheng, X. Li and J. Y. Luo, *Water Res.*, 2014, **63**, 125–134.
- 103 M. Sibag, B.-G. Choi, C. Suh, K. H. Lee, J. W. Lee, S. K. Maeng and J. Cho, *J. Hazard. Mater.*, 2015, **283**, 841–846.

