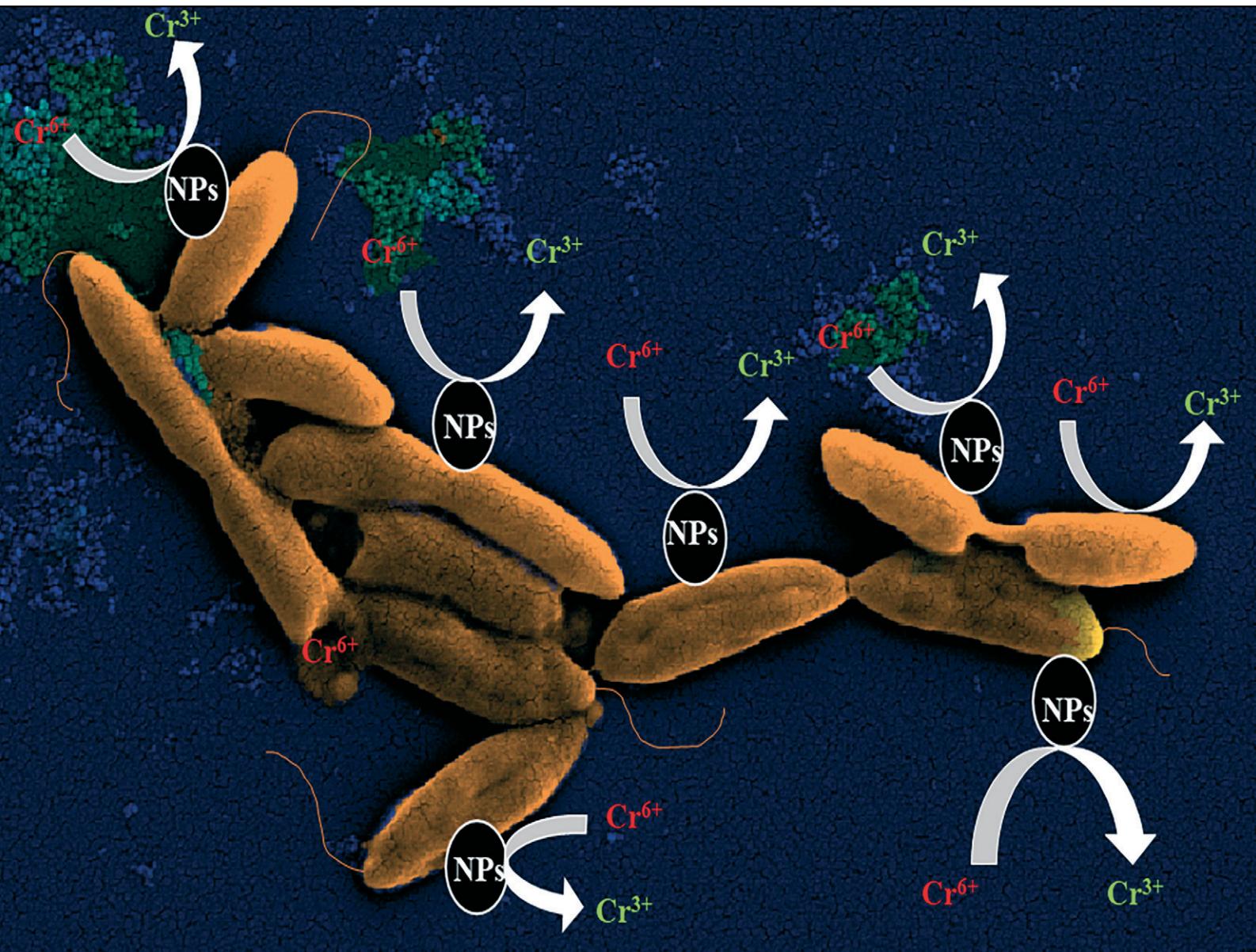


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## Protection of *Shewanella oneidensis* MR-1 by manganese ferrite nanoparticles during chromate bio-reduction†

Diana S. Raie, <sup>ab</sup> Ioannis Tsonas, <sup>c</sup> Stefanos Moudrikoudis, <sup>ab</sup> Evangelia Delli, <sup>d</sup> Antonios Makridis, <sup>d</sup> Lena Ceric<sup>e</sup> and Nguyen Thi Kim Thanh <sup>\*</sup><sup>ab</sup>

*Shewanella oneidensis* (*S. oneidensis*) MR-1 is a metal-reducing bacterium that can bio-reduce the carcinogenic hexavalent chromium ( $\text{Cr}^{6+}$ ) to a less toxic trivalent form ( $\text{Cr}^{3+}$ ). The bacteriocidal effect of  $\text{Cr}^{6+}$  challenges the above bio-reduction process. This work aims to illustrate the protective role of manganese ferrite nanoparticles ( $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs) to *S. oneidensis* MR-1 bacteria during the bio-reduction of  $\text{Cr}^{6+}$ . Nanostructures were characterised by transmission electron microscopy (TEM) and X-ray diffraction (XRD). The interaction between *S. oneidensis* MR-1,  $\text{Cr}^{6+}$  and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs was monitored by X-ray photoelectron spectroscopy (XPS), which helped to unravel the oxidation states of Cr. The XPS analysis provided key insights into the oxidation states of Mn and Fe, confirming the redox interactions facilitating  $\text{Cr}^{6+}$  reduction.  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs boosted the detoxification of the removed  $\text{Cr}^{6+}$  by 2.1 and 1.4 times compared to using *S. oneidensis* MR-1 alone and NPs alone, respectively. Scanning electron microscopy (SEM) imaging evaluated the changes in the morphology of bacterial cells. After exposure to  $\text{Cr}^{6+}$ , *S. oneidensis* MR-1 cells revealed their inability to produce nanofibers, which are electrically conductive bacterial appendages. Yet,  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs provoked the formation of bacterial nanofibers. These findings highlight the potential of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs for enhancing the bioremediation of  $\text{Cr}^{6+}$  contaminated environments.

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### Environmental significance

Carcinogenic hexavalent chromium leaks from industrial sites due to improper wastewater treatment into surface and groundwater, exposing flora and fauna to danger. The metal-reducing bacterium, *Shewanella oneidensis* MR-1, can reduce  $\text{Cr}^{6+}$  into less toxic  $\text{Cr}^{3+}$ ; bacteria lose their viability during treatment due to the toxicity of  $\text{Cr}^{6+}$ . The novelty of this work is the discovery of a protective role of Mn-ferrite nanoparticles to *S. oneidensis* MR-1 bacteria during  $\text{Cr}^{6+}$  bio-reduction. We show that  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs induced bacterial cell elongation and promoted nanofiber formation. Such morphological changes improve bacterial cell viability in response to the sub-lethal dose of  $\text{Cr}^{6+}$  and enhance their detoxification capability. Our findings provide a promising application of using nano- $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  in the bioremediation of  $\text{Cr}^{6+}$ -contaminated environments.

## 1. Introduction

Contamination of air, soil and water with heavy metals is hazardous to human health and the environment due to their

toxicity, even at low concentrations<sup>1</sup> as they are non-biodegradable materials.<sup>2</sup> The Agency for Toxic Substances and Disease Registry (ATSDR) ranked chromium (Cr) the 17th on the substance priority list among many heavy metals.<sup>3</sup>

Cr mainly occurs in two valence states: hexavalent ( $\text{Cr}^{6+}$ ) and trivalent ( $\text{Cr}^{3+}$ ). Human exposure to  $\text{Cr}^{6+}$  can cause liver damage, pulmonary congestion, oedema, skin irritation, ulcer formation,<sup>4</sup> neurotoxicity,<sup>5</sup> and carcinogenesis.<sup>6</sup> U.S. Environmental Protection Agency (EPA) and WHO guidelines reported a permissible limit of  $\text{Cr}^{6+}$  in drinking water of 50 ppb.<sup>7</sup> According to the EU drinking water directive, the regulation limit for the total Cr will be 25  $\mu\text{g L}^{-1}$  by 12 January 2036.<sup>8</sup> Since  $\text{Cr}^{3+}$  has low mobility, limited bio-absorptivity, and lower toxicity than  $\text{Cr}^{6+}$ ,<sup>9</sup>  $\text{Cr}^{6+}$  should be reduced to  $\text{Cr}^{3+}$  for its safe removal.<sup>10</sup>

<sup>a</sup> Biophysics Group, Department of Physics and Astronomy, University College London (UCL), London, WC1E 6BT, UK

E-mail: nt.k.thanh@ucl.ac.uk; Web: <https://www.ntk-thanh.co.uk>

<sup>b</sup> UCL Healthcare Biomagnetics and Nanomaterials Laboratories, 21 Albemarle Street, London, W1S 4BS, UK

<sup>c</sup> UCL Electronic and Electrical Engineering, UCL, Gower Street, London, WC1E 7JE, UK

<sup>d</sup> Laboratory of Advanced Materials and Devices, Department of Physics, Aristotle University of Thessaloniki, 54124 Thessaloniki, Greece

<sup>e</sup> Healthy Infrastructure Research Group, Department of Civil, Environmental & Geomatic Engineering, UCL, Gower Street, WC1E 6BT London, UK

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Bio-reduction of  $\text{Cr}^{6+}$  is a cost-effective and environmentally friendly method, attracting widespread interest.<sup>11</sup> Some bacteria can reduce metals, acting as terminal electron acceptors under anaerobic conditions.<sup>12</sup> So, metal-reducing bacteria can be used for the biotic reduction of heavy metals for detoxification purposes. Such a natural process is applicable for the biological reduction of the carcinogenic  $\text{Cr}^{6+}$  into less toxic  $\text{Cr}^{3+}$  form.<sup>13</sup>

*Shewanella oneidensis* MR-1 is a model metal-reducing bacteria for detoxifying  $\text{Cr}^{6+}$ .<sup>14–18</sup> *S. oneidensis* MR-1 can employ  $\text{Cr}^{6+}$  as a terminal electron acceptor under anaerobic conditions.<sup>14,15,19</sup> The biosafety of *S. oneidensis* MR-1 is an essential criterion for selecting bioremediation biological agents. In contrast, *Pseudomonas aeruginosa* bacteria can be used for  $\text{Cr}^{6+}$  removal but are not preferred for bioremediation because they cause diseases in humans and animals.<sup>20,21</sup> Yet, the lethal effect of  $\text{Cr}^{6+}$  on the microbes during their respiration limited the bioremediation of  $\text{Cr}^{6+}$ .<sup>22</sup>

$\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs showed a higher adsorption capacity for  $\text{Cr}^{6+}$  than  $\text{Fe}_2\text{O}_3$  NPs and other tested  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs.<sup>23</sup> This chemical structure improved the bacterial viability and microbial detoxification of  $\text{Cr}^{6+}$ .<sup>23</sup> The adsorption of  $\text{Cr}^{6+}$  can limit the availability of the toxic cations to cells, which could improve their viability and bio-reduction efficiency.

Herein, to the best of our knowledge, we showed for the first time the protective role of the  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs to *S. oneidensis* MR-1 during the bio-reduction of  $\text{Cr}^{6+}$ . Raie *et al.*<sup>23</sup> primarily investigated the adsorption and bio-removal of  $\text{Cr}^{6+}$  using  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs and *S. oneidensis* MR-1, respectively.

This article builds upon findings by Raie *et al.*,<sup>23</sup> and elucidates the reduction process of  $\text{Cr}^{6+}$  using XPS. In addition, this work presents bacterial imaging to visualise morphological changes in response to  $\text{Cr}^{6+}$  and NPs, providing deeper insights into the mechanism of  $\text{Cr}^{6+}$  reduction.

XPS revealed the possible reduction of  $\text{Cr}^{6+}$  to  $\text{Cr}^{3+}$  due to its interaction with  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs. This allowed us to confirm the redox-based interaction among  $\text{Cr}^{6+}$  and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs. In addition, SEM showed the morphological change response of *S. oneidensis* MR-1 as a coping strategy in response to the toxic  $\text{Cr}^{6+}$  in the presence of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs. This article will advance the treatment of  $\text{Cr}^{6+}$  by demonstrating its removal, unravelling its reduction mechanism and the biological implications, thereby contributing novel insights and practical advancements to nanobiotechnology and environmental applications.

## 2. Materials and methods

### 2.1 NPs preparation and characterisation

$\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs were prepared by an adapted polyol solvothermal synthetic process<sup>24,25</sup> at 250 °C as described in our recent work.<sup>23</sup> In 20 mL of tetraethylene glycol (TEG), 0.3 M of iron(III) acetylacetone (2.1 g) and 0.1 M manganese(II) acetylacetone (0.5 g) were added. The mixture was added into a 45 mL Teflon-lined stainless-steel autoclave after being homogenised by vortex and sonication to be placed in an oven (Memmert, model UFP400) and heated within 30 min up to 250

°C for a 6 h hold at that temperature. In polyol synthesis, metal precursors are reduced by TEG, which acts as a high-temperature capping agent, solvent, and reductant. The formed metal nuclei grow and controllably coalesce together to produce the desired particles.<sup>26,27</sup> The produced black dispersion underwent characterisation and functionalisation by tri-sodium citrate *via* ligand exchange.<sup>23</sup> A JEOL JEM 1200-EX microscope operating at an acceleration voltage of 120 kV was employed to investigate the shape and size of the produced particles. The polydispersity index (PDI) is the ratio between the standard deviation and the mean nanoparticle diameter. To determine the crystal phase and the average crystallite size, we used XRD (PANalytical XPERT PRO MPD) coupled with Co K $\alpha$  radiation source ( $\lambda = 1.789$  Å) and an X'Celetor detector operated at 40 kV and 40 mA. An Optima 3100 XL Perkin Elmer Inductively Coupled Plasma Atomic Emission (ICP-AES) spectrometer was employed to determine the chemical composition of  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  particles. To quantify the iron content of the functionalised NPs dispersed in water, a colorimetric phenanthroline method was applied for the acid-digested NPs using a spectrophotometer (SpectraMax M2e, Molecular Devices, UK).

### 2.2 Sources for bacteria of interest

A freeze-dried culture of *S. oneidensis* MR-1 (LMG 19005) was purchased from BCCM/LMG bacteria collection.

### 2.3 Viability of *S. oneidensis* MR-1 to $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$ NPs

The impact of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs on the viability of the *S. oneidensis* MR-1 was assessed using Guava easyCyte® flow cytometer (Merck, UK) following a protocol previously utilised by Raie *et al.*,<sup>23</sup> under anaerobic conditions overnight. A homogeneous bacterial cell suspension (10  $\mu\text{L}$  with OD measured at  $\lambda = 600$  nm equal to 0.1) was added to 80  $\mu\text{L}$  of M9 minimal salts ( $\times 2$ ) medium, containing 20 mM sodium lactate as a sole electron source, 5 mL  $\text{L}^{-1}$  each of vitamins and minerals and pH was adjusted to 7.2 by 10 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES) buffer.<sup>23,28</sup> Sodium fumarate (20 mM) was used as a terminal electron acceptor.<sup>23,28</sup>  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs (10  $\mu\text{L}$ ) were added to the mixture. The tested concentrations of NPs ranged from 1–60 mg  $\text{mL}^{-1}$  with an approximate total Fe content from 0.7 mg  $\text{mL}^{-1}$  to 40.6 mg  $\text{mL}^{-1}$ .

### 2.4 The exposure of *S. oneidensis* MR-1 to $\text{Cr}^{6+}$ and $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$ NPs

*S. oneidensis* MR-1 was exposed to  $\text{Cr}^{6+}$  and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs individually and also in a combined way overnight, in conditions similar to that mentioned in Section 2.3.  $\text{Cr}^{6+}$  (as a terminal electron acceptor) and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs were added to this medium with concentrations of 50 mg  $\text{L}^{-1}$  (sub-lethal dose, as reported by Raie *et al.*)<sup>23</sup> and 1 mg  $\text{mL}^{-1}$ , respectively.<sup>23</sup>



## 2.5 Analysis of oxidation state of $\text{Cr}^{6+}$ , $\text{Mn}^{x+}$ , and $\text{Fe}^{y+}$

The oxidation states of Mn and Fe in  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs and Cr were investigated after being incubated together or separately with *S. oneidensis* MR-1 by XPS; a Kratos Analytical AXIS Ultra<sup>DLD</sup> system with aluminium X-ray source ( $\lambda_{\text{Ka}} = 1486.6$  eV) was used, operated under ultra-high vacuum conditions ( $10^{-9}$  torr). The experimental curves were best fitted by combining Gaussian (70%) and Lorentzian (30%) distributions, while background subtraction was performed using the Shirley equation. A normalised peak area of each element is calculated by dividing its area by the sensitivity factor.<sup>29</sup> To determine the redox interaction between  $\text{Cr}^{6+}$  and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs, we compared the normalised peak areas of  $\text{Mn}^{2+}$  to  $\text{Mn}^{3+}$ ,  $\text{Fe}^{2+}$  to  $\text{Fe}^{3+}$  and  $\text{Cr}^{3+}$  to  $\text{Cr}^{6+}$  in the high-resolution Mn 2p, Fe 2p and Cr 2p spectra, respectively, while only the ratios between that peak areas of  $\text{Cr}^{3+}$  to  $\text{Cr}^{6+}$  were analysed in the case of applying bacterial cells. The relative fold increase in  $\text{Cr}^{6+}$  bio-reduction was calculated by its equivalent atomic fraction to the reference values.

## 2.6 Imaging bacteria by SEM

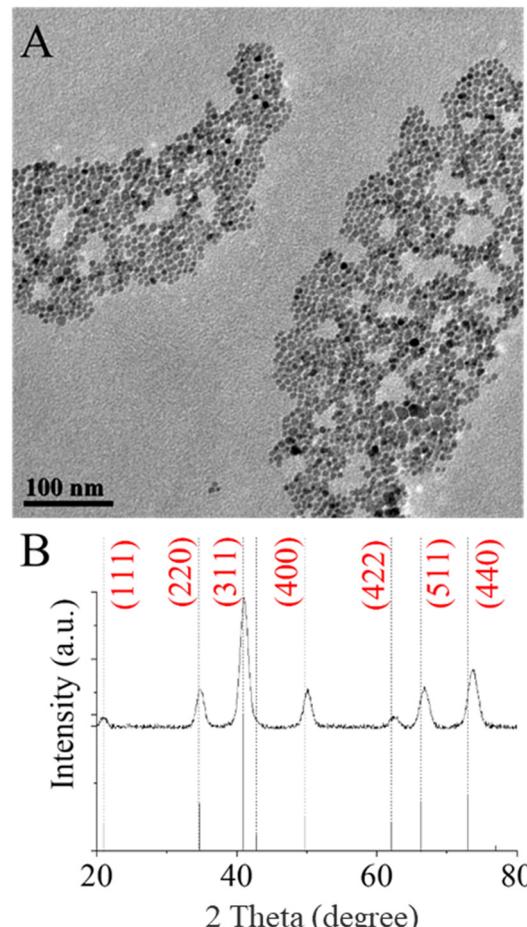
To acquire SEM images, 50  $\mu\text{L}$  from the untreated or  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs treated *S. oneidensis* MR-1 bacteria cell suspension were deposited on a microscope cover glass (Fisher, UK). The samples were imaged using Philips XL30 FEG SEM (FEI, Eindhoven, Netherlands), which operates at an accelerating voltage of 5 keV. Cell fixation was performed using glutaraldehyde (2.5% v/v in 0.01 M PBS) for 30 min at room temperature. Samples were washed three times in phosphate-buffered saline (PBS, 0.01 M) and dehydrated for 5 min in ethanol aqueous solutions. The concentrations of ethanol aqueous solutions were 10% v/v, 30% v/v, 50% v/v, 70% v/v, 90% v/v, 100% v/v, sequentially. A double-sided carbon tape (Agar Scientific, UK) was used to attach the glass slide with the SEM specimens onto aluminium stubs. Samples were then sputter-coated with gold-palladium at 20 mA and 1.25 kV for 90 s (Palaron E5000 sputter coater).

## 3. Results and discussion

### 3.1 Characterisation of NPs

**3.1.1 Morphology of NPs.** Regarding the obtained spherical  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs (Fig. 1A), our results agree with Raie *et al.*,<sup>23</sup> Vamvakidis *et al.*,<sup>25</sup> and García-Soriano *et al.*,<sup>30</sup> who used the polyol solvothermal technique for producing spherical  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs.<sup>23,25,30</sup> The mean size of  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs is  $7.4 \pm 1.3$  nm. The PDI is 0.18, which indicates a relatively narrow size distribution.<sup>31</sup> Similarities in spherical shape and small size range (approximately 7–9 nm) are attributed to the specific procedure where sole polyols were used to prepare the NPs.<sup>23,25,30</sup>

**3.1.2 Crystal structure of NPs.** Powder XRD patterns for the prepared  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs recorded at room temperature are illustrated in Fig. 1B. All the diffraction peaks show the presence of the face-centred cubic (FCC) crystal structure, while no impurity phase was observed. So, the formation of



**Fig. 1**  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs prepared at 250 °C for 6 h: (A) TEM images and (B) XRD patterns, and XRD reference for  $\text{MnFe}_2\text{O}_4$  (PDF card no 00-010-0319).

$\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs was obtained through a facile polyol solvothermal process with reaction times of 6 h.

**3.1.3 Elemental analysis of  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs.** The formed  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs have a low Mn content ( $x = 0.2$ ), based on ICP-AES results. Etemadi & Plieger,<sup>32</sup> Oberdick *et al.*,<sup>33</sup> and Raie *et al.*<sup>23</sup> reported similar results of low Mn doping levels because  $\text{Mn}(\text{acac})_2$  is more thermally stable than  $\text{Fe}(\text{acac})_3$ .<sup>34</sup>

### 3.2 Interaction of $\text{Cr}^{+6}$ with $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$ NPs

$\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs adsorbed  $16.8 \pm 1.6 \text{ mg g}^{-1}$  (around 61%) of  $\text{Cr}^{6+}$ .<sup>23</sup> The possible reduction of the adsorbed  $\text{Cr}^{6+}$  by  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs was explored here by studying the oxidation state of Mn, Fe, and Cr of the adsorbent and adsorbate by XPS, as shown in Fig. 2A.

**3.2.1 Oxidation state of Mn in  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs after  $\text{Cr}^{6+}$  adsorption.** In Fig. 2B, the position of binding energy (BE) for Mn 2p was slightly shifted from 640.45 eV<sup>23</sup> to higher BE (641.80 eV), which could be attributed to the possible oxidation of  $\text{Mn}^{2+}$  into  $\text{Mn}^{3+}$  upon interacting with  $\text{Cr}^{6+}$ . The dissolved  $\text{Mn}^{3+}$  could generate manganese oxide ( $\text{MnO}_x$ ), which provides more adsorption sites for  $\text{Cr}^{6+}$  removal.<sup>35</sup>



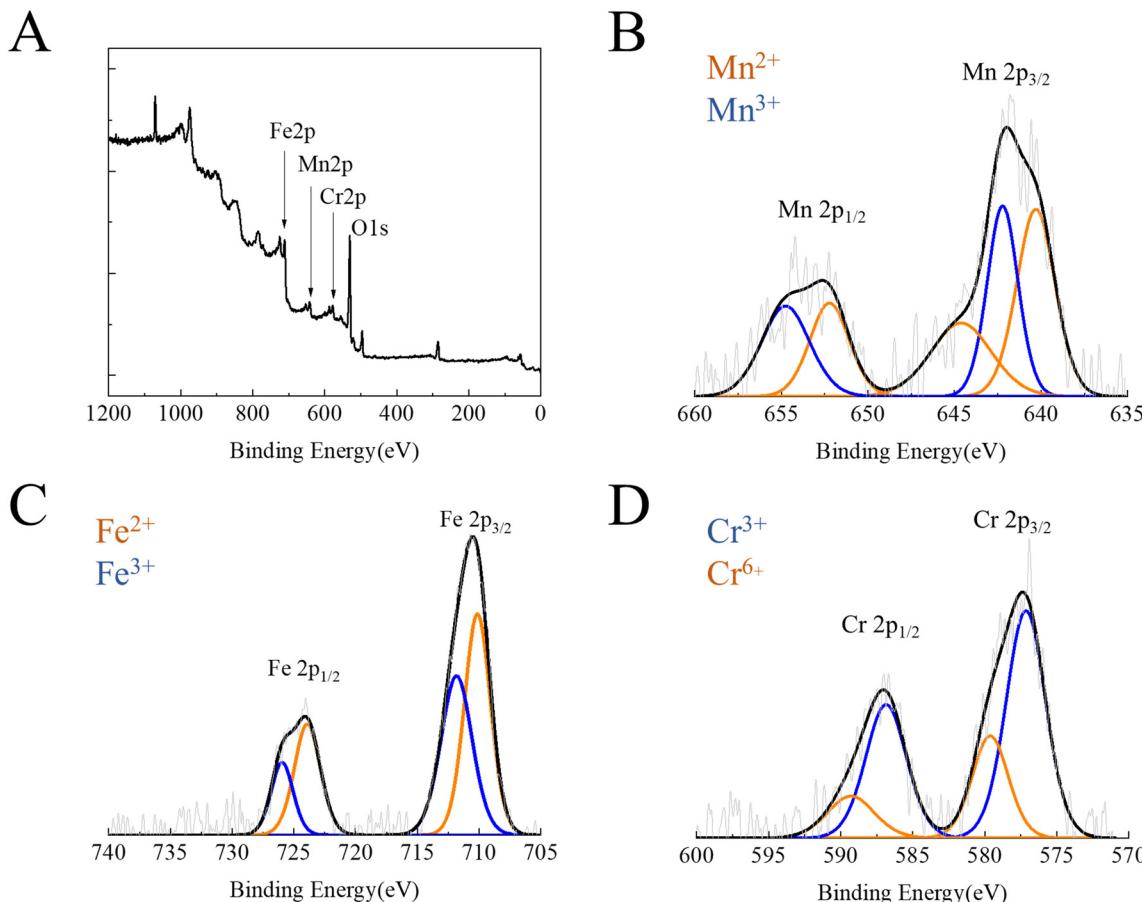


Fig. 2  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs treated by  $\text{Cr}^{6+}$ : (A) wide scan XPS spectrum, and high-resolution XPS spectra of (B) Mn 2p, (C) Fe 2p, and (D) Cr 2p.

In  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs,<sup>23</sup> Mn 2p peak, in Fig. 2B, was fitted by 5 contributions at 640.3 eV, 642.2 eV, 644.61 eV, 652.2 eV and 654.8 eV. Mn 2p<sub>3/2</sub> was deconvoluted into 640.35 eV and 642.25 eV peaks, representing  $\text{Mn}^{2+}$  and  $\text{Mn}^{3+}$ , respectively, as shown in Fig. 2B. The peak of Mn 2p<sub>1/2</sub> was fitted into two contributions of  $\text{Mn}^{2+}$  and  $\text{Mn}^{3+}$  at 652.15 eV and 654.6 eV, respectively.<sup>36–39</sup> The fifth small satellite peak at 645.2 eV was assigned to  $\text{Mn}^{2+}$  of  $\text{MnO}$ .<sup>38</sup> Since stoichiometric  $\text{MnFe}_2\text{O}_4$  can be expressed as  $\text{MnO}\text{-Fe}_2\text{O}_3$ , this pointed to the formation of  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs.

**3.2.2 Oxidation state of Fe after  $\text{Cr}^{6+}$  interaction with  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs.** After  $\text{Cr}^{6+}$  adsorption on  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs, XPS (in Fig. 2C) showed the position of BE for Fe 2p at 711 eV. A peak of Fe 2p<sub>3/2</sub> was spotted at 710.75 eV, and the asymmetric peaks are situated at 723.9 eV, attributed to 2p<sub>1/2</sub>.<sup>39,40</sup> The observed signals at these BE positions probably correspond to the formation of iron oxide phase, *i.e.*, hematite or maghemite phase.<sup>41</sup> Unlike untreated  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs,<sup>23</sup> Fe 2p missed the satellite peak at 718 eV as shown in Fig. 2C, which was due to the presence of  $\text{Fe}_3\text{O}_4$ .<sup>40</sup> The ratio between Mn and Fe was doubled from 0.24 to 0.44 (as was reported by XPS and range based on elemental analysis by ICP in our recent work) compared to untreated  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs,<sup>23</sup> which can be ascribed to the release of iron in the medium.

**3.2.3 Reduction of  $\text{Cr}^{6+}$  by  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$ .** In Fig. 2D, XPS spectra of Cr 2p showed two different peaks, corresponding to the Cr 2p<sub>3/2</sub> (576.0 eV–578.0 eV) and Cr 2p<sub>1/2</sub> (585.0 eV–587.0 eV) orbits. After fitting peaks with the use of the Gauss–Lorentz algorithm, two peaks arised with the BE of 577 eV relating to  $\text{Cr}^{3+}$  2p<sub>3/2</sub> and 586 eV belonging to  $\text{Cr}^{3+}$  2p<sub>1/2</sub>,<sup>42,43</sup> which mainly corresponds to the precipitation of insoluble  $\text{Cr}^{3+}$  species,  $\text{Cr}(\text{OH})_3$  and  $\text{Cr}_2\text{O}_3$ . The adsorbed  $[\text{CrO}_4]^{2-}$  on NPs<sup>37</sup> explained the presence of peaks at BE of 579.6 eV and 589 eV, representing  $\text{Cr}^{6+}$  2p<sub>3/2</sub> and 2p<sub>1/2</sub>, respectively.<sup>43</sup> The ratio of  $[\text{Cr}^{3+}]/[\text{Cr}^{6+}]$  was estimated to be equal to 2.56. Our results point out a significant finding: the interaction between  $\text{Cr}^{6+}$  and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs involved a redox reaction in addition to what was stated in our recent work regarding adsorption.<sup>23</sup> Raie *et al.* reported that the oxidation state of Mn in  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  was mainly  $\text{Mn}^{2+}$  with a minor fraction of  $\text{Mn}^{4+}$ , and that of Fe was a mixture of  $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$ .<sup>23</sup> In the present study, the possible oxidation of  $\text{Mn}^{2+}$  to  $\text{Mn}^{3+}$  and  $\text{Fe}^{2+}$  to  $\text{Fe}^{3+}$ , besides the iron release, is due to the redox reaction between  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs and  $\text{Cr}^{6+}$ . The absence of  $\text{Mn}^{4+}$  XPS related peak after interaction with  $\text{Cr}^{6+}$  was attributed to the ability of  $\text{Fe}^{2+}$  to reduce  $\text{Mn}^{4+}$ , yielding  $\text{Fe}^{3+}$  and  $\text{Mn}^{2+}$ .<sup>44</sup> In addition to being a stabilising agent, citrate can act as a chelating agent<sup>45</sup> and as a reductant for  $\text{Cr}^{6+}$ ,<sup>46</sup> due to its ability to donate electrons through ligand–metal electron transfer.<sup>46</sup>  $\text{Mn}^{2+}$  catalyses the reduction reaction.<sup>47</sup>



### 3.3 $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$ NP-assisted bacterial respiration of $\text{Cr}^{6+}$

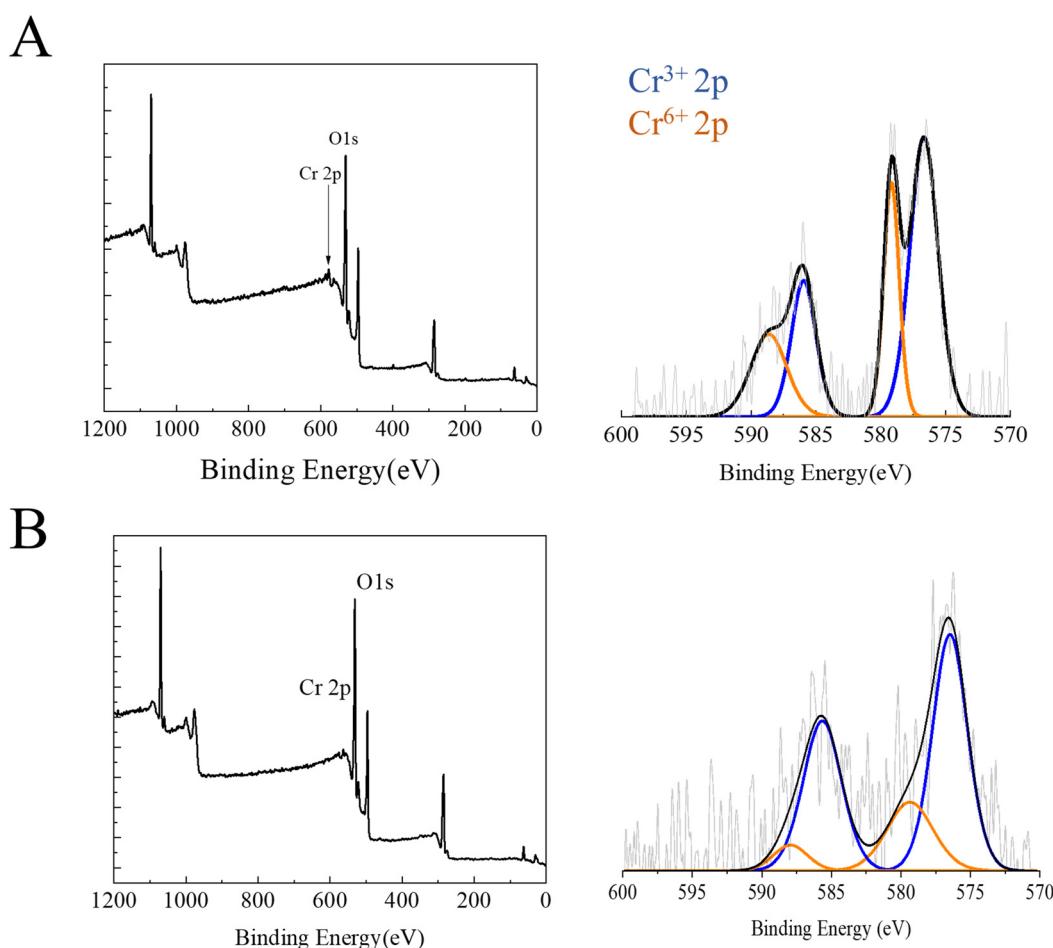
*S. oneidensis* MR-1 can respire  $\text{Cr}^{6+}$  under anaerobic conditions.<sup>48–50</sup> The adsorption of  $\text{Cr}^{6+}$  ( $9 \pm 1.5 \text{ mg g}^{-1}$ , *i.e.*  $30 \pm 0.5\%$  of removal) by  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs supported microbial survival in media supplemented with the tested *S. oneidensis* MR-1.<sup>23</sup> The mechanism of bio-removal of  $\text{Cr}^{6+}$  can be attributed to the respiration of  $\text{Cr}^{6+}$  into  $\text{Cr}^{3+}$  (ref. 48–50) or bio-sorption<sup>51,52</sup> by bacterial cells. Examining the oxidation state of Cr element *via* XPS analysis determines the interaction between  $\text{Cr}^{6+}$ ,  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs and *S. oneidensis* MR-1, as illustrated in Fig. 3, which positively related to the enhanced  $\text{Cr}^{6+}$  bio-reduction by 2.7–3.6 fold.<sup>23</sup> The reported significant drop in the XPS revealed the presence of peaks related to both  $\text{Cr}^{6+}$  and  $\text{Cr}^{3+}$  after exposing *S. oneidensis* MR-1 to  $\text{Cr}^{6+}$ .

Peaks of Cr 2p XPS were observed at BE 576.7 eV and 585.9 eV, which were related to  $\text{Cr}^{3+}$ , while peaks at 579.2 eV and 588.6 eV were assigned to  $\text{Cr}^{6+}$ , as presented in Fig. 3A. *S. oneidensis* MR-1 can reduce  $\text{Cr}^{6+}$  into  $\text{Cr}^{3+}$ , as confirmed by our XPS results in Fig. 3A and supported by the literature.<sup>48,53</sup>

Our findings reveal an extracellular interaction between  $\text{Cr}^{6+}$  and *S. oneidensis* MR-1 bacteria. A portion of  $\text{Cr}^{6+}$  was reduced

to  $\text{Cr}^{3+}$ , resulting in a  $[\text{Cr}^{3+}]/[\text{Cr}^{6+}]$  ratio of 1.7, while the remaining 41% of  $\text{Cr}^{6+}$  is adsorbed on the bacterial cell surface. The extracellular reduction of  $\text{Cr}^{6+}$  can occur *via* direct contact of  $\text{Cr}^{6+}$  with the metal-reducing protein complex on the cell surface and nanofiber. Also, *S. oneidensis* MR-1 can produce electron shuttles to promote mediated electron transfer between the cell and  $\text{Cr}^{6+}$ . *S. oneidensis* MR-1 can uptake  $\text{Cr}^{6+}$  to be reduced inside the cell to  $\text{Cr}^{3+}$ , but our results could not confirm the intracellular reduction of  $\text{Cr}^{6+}$  due to the depth limitation of XPS (7–10 nm).

Our XPS results revealed peaks related to both  $\text{Cr}^{6+}$  and  $\text{Cr}^{3+}$  after being incubated with *S. oneidensis* MR-1 in the presence of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs. Peaks of Cr 2p XPS observed at BE 576.7 eV and 585.9 eV denote the presence of  $\text{Cr}^{3+}$ .  $\text{Cr}^{6+}$  is represented by one peak at 579.2 eV,<sup>38</sup> as illustrated in Fig. 3B. Similar results were reported due to using  $\text{Cr}^{6+}$  as a terminal electron acceptor during the respiration process of *S. oneidensis* MR-1.<sup>48,53</sup> The ratio between extracellular  $\text{Cr}^{3+}$  and  $\text{Cr}^{6+}$  was equal to 3.5. Bacteria can reduce  $\text{Fe}^{3+}$  to  $\text{Fe}^{2+}$ , and biogenic  $\text{Fe}^{2+}$  can detoxify  $\text{Cr}^{6+}$  to  $\text{Cr}^{3+}$ .<sup>54,55</sup> The affinity of  $\text{MnFe}_2\text{O}_4$  NPs to proteins on the bacterial outer membrane can improve the contact area between a single bacterium and  $\text{Cr}^{6+}$  as an external electron acceptor.<sup>56–59</sup>



**Fig. 3** Wide scan and high-resolution XPS spectra of (A) Cr 2p treated by *S. oneidensis* MR-1 alone, (B) Cr 2p after incubation of *S. oneidensis* MR-1 and  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs.



In this work, the presence of both *S. oneidensis* MR-1 and  $Mn_{0.2}Fe_{2.8}O_4$  NPs removed  $Cr^{6+}$  1.37 times more than using the NPs alone. Some possible scenarios could explain how NPs enhanced the bio-reduction of *S. oneidensis* MR-1 from  $Cr^{6+}$  to  $Cr^{3+}$ . By adsorption, NPs can bridge the bacterial cell and  $Cr^{6+}$  to promote electron transfer.  $Cr^{6+}$  is adsorbed onto the  $MnFe_2O_4$  NPs *via* partial chemisorption<sup>60,61</sup> and partial physisorption.<sup>37</sup> The Mn in  $MnFe_2O_4$  can interact *via* ionic bonding with the O atoms of  $HCrO_4^-/CrO_4^{2-}$ , facilitating  $Cr^{6+}$  adsorption.<sup>60,61</sup>  $Mn^{2+}$  can reduce  $Cr^{6+}$  to  $Cr^{3+}$  and be oxidised to  $Mn^{3+}$ . The disproportionation of oxidised  $Mn^{3+}$  produced  $Mn^{2+}$ , causing  $Mn^{2+}$  to continue participating in the  $Cr^{6+}$  reduction.  $Cr^{3+}$  is deposited on the  $MnFe_2O_4$  surface as  $Cr(OH)_3$  colloids.<sup>60,61</sup>

The limited availability of adsorbed  $Cr^{6+}$  improved the efficiency of microbial respiration,<sup>48,54</sup> as was indicated by our results. Since  $MnFe_2O_4$  NPs have electrochemical properties,<sup>59,62</sup> metal oxides can link *S. oneidensis* MR-1 with  $Cr^{6+}$  to promote direct electron transfer and act as an electron mediator from the cell to  $Cr^{6+}$ , a terminal electron acceptor.<sup>63</sup>

In addition, NPs can act as physical shields for bacterial surfaces from  $Cr^{6+}$ , which could reduce the direct damage to bacteria caused by heavy metals. Encapsulating *S. loihica* by biochar reported that it could avoid the lethal effect of  $Cr^{6+}$ .<sup>63</sup> In addition,  $Mn_{0.2}Fe_{2.8}O_4$  NPs can sustain bacterial viability, as shown in Fig. S1† and supported by the literature.<sup>64</sup> The Mn content in the chemical structure of  $Mn_{0.2}Fe_{2.8}O_4$  NPs improved the anti-oxidant activity of NPs and, in turn, cell viability.<sup>65</sup> Substituting  $Fe^{2+}$  by  $Mn^{2+}$  in  $Mn_{0.2}Fe_{2.8}O_4$  NPs decreased the lethal effect of  $Fe^{2+}$  on bacterial viability. This explains how  $Mn_{0.2}Fe_{2.8}O_4$  NPs improved the viability of *S. oneidensis* MR-1 under the sub-lethal concentration of  $Cr^{6+}$  by 3.3 times.<sup>23</sup>

### 3.4 Boosting the bacterial tolerance to $Cr^{6+}$ by $Mn_{0.2}Fe_{2.8}O_4$ NPs

SEM imaging monitored the alterations in the morphology of bacterial cells following the bio-reduction.

**3.4.1 Morphology of untreated bacterial cells.** The untreated tested *S. oneidensis* MR-1 demonstrated their viability under anaerobic redox conditions, as shown in Fig. 4A. In the absence of both  $Cr^{6+}$  and  $Mn_{0.2}Fe_{2.8}O_4$  NPs, bacterial cells of *S. oneidensis* MR-1 were observed as rod-shaped with smooth surfaces as commonly described.<sup>22,49,63,66–68</sup> The formation of the division ring (Z-ring) at the division site at the mid-cell on the bacteria was an indicator of cell division, as depicted in Fig. 4A. Parker *et al.*<sup>22</sup> reported that the delay in separating daughter cells could be ascribed to the minimum availability of nutrients in the media.<sup>22</sup> The presence of bacterial nanofibers as extensions of the outer membrane and periplasm (Fig. 4A) was demonstrated to be increased under oxygen-limited conditions.<sup>66</sup> Nanofibers were reported to have the multiheme cytochromes responsible for the extracellular electron transport pathway for linking the respiratory chain of bacteria to an external electron acceptor.<sup>66</sup> Electrons are transferred along nanofibers of *S. oneidensis* MR-1 between the close cytochromes *via* an electron-hopping mechanism.<sup>67,68</sup>

**3.4.2 Rupture of *S. oneidensis* MR-1 cells in response to a sub-lethal dose of  $Cr^{6+}$ .** The impact of exposure of *S. oneidensis* MR-1 to  $Cr^{6+}$  was observed on the rupture on one pole of a cell, as shown in Fig. 4B. A shrunken surface and crack formation in bacteria cells were also observed after the reaction with  $Cr^{6+}$ .<sup>22,49,69</sup> As in the case of untreated cells, attempts of cell division were still observed for cells exposed to  $Cr^{6+}$ , as demonstrated in Fig. 4B. The presence of cell division septa was an indicator for the initial phase of cell division of *S. oneidensis* cells.<sup>22</sup> SEM images of *S. oneidensis* MR-1 revealed the inability to produce nanofibers after exposure to  $Cr^{6+}$ . The variation in the length of cells exposed to  $Cr^{3+}$  is presented in Fig. 4B. Bacterial cells modified their shape as a coping strategy for tolerating the stress induced by  $Cr^{6+}$ .<sup>55,63</sup>

**3.4.3 Cellular compatibility of  $Mn_{0.2}Fe_{2.8}O_4$  NPs.** Fig. S1† shows the biocompatibility of different concentrations of  $Mn_{0.2}Fe_{2.8}O_4$  NPs towards *S. oneidensis* MR-1. Our findings were supported by Desai *et al.*, who reported that  $MnFe_2O_4$  NPs showed no antimicrobial activity against some pathogenic bacteria.<sup>70</sup> *Shewanella* can survive upon exposure to 50 mg mL<sup>-1</sup> of magnetite ( $Fe_3O_4$ ) with approximately 36.2 mg mL<sup>-1</sup> of total iron content under anaerobic conditions. Such tolerance to high iron concentrations was due to the cellular attachment to magnetite for  $Fe^{3+}$  acquisition.<sup>71</sup> The tolerance of *S. oneidensis* MR-1 to such concentrations of  $Mn_{0.2}Fe_{2.8}O_4$  NPs could be attributed to the presence of  $Mn^{2+}$  in the chemical structure of NPs, which improved the antioxidant activity, cell viability, and ability to respire metal.<sup>65</sup> In addition, the  $Mn^{2+}$  content in  $Mn_{0.2}Fe_{2.8}O_4$  NPs lowered  $Fe^{2+}$  concentration, which could decrease the lethal effect of  $Fe^{2+}$  on the viability of the tested bacterial strain. The presence of  $Fe^{3+}$  in  $Mn_{0.2}Fe_{2.8}O_4$  NPs<sup>23</sup> has less toxicity than  $Fe^{2+}$  under physiological conditions.<sup>72</sup> The resistance of *S. oneidensis* MR-1 to  $Fe^{2+}$  depends on the ClpXP protease complex, which removes the mis-metallated protein. ClpX is an unfoldase, and ClpP is a peptidase that degrades damaged or misfolded proteins.<sup>73</sup>

The capability of *Shewanella* to produce nanofibers in the presence of  $Mn_{0.2}Fe_{2.8}O_4$  NPs is shown in Fig. 4C. The poles of *Shewanella* cells were reported to be attractive to the metal oxide/hydroxides under both aerobic and anaerobic conditions,<sup>74,75</sup> which explains the polar rupture of some cells in Fig. 4C.

**3.4.4 Enhanced tolerance of *Shewanella* to  $Cr^{6+}$  by  $Mn_{0.2}Fe_{2.8}O_4$  NPs.** In the presence of  $Mn_{0.2}Fe_{2.8}O_4$  NPs, the surface of the treated *S. oneidensis* MR-1 cells by  $Cr^{6+}$  retained a smooth surface but with an elongated morphology (see Fig. 4D). Such stretching in the shapes of cells was observed by *S. loihica* PV-4 in response to  $Cr^{6+}$  in a mixture containing biochar and  $\alpha$ - $Fe_2O_3$  together.<sup>63</sup> The morphological changes observed in the bacteria are adaptive strategies for coping with environmental stresses like the presence of toxic  $Cr^{6+}$ . Inhibiting cell division while maintaining cell growth leads to increased cell length<sup>76</sup> and boosts the extracellular electron transfer by *S. oneidensis* MR-1.<sup>77</sup>



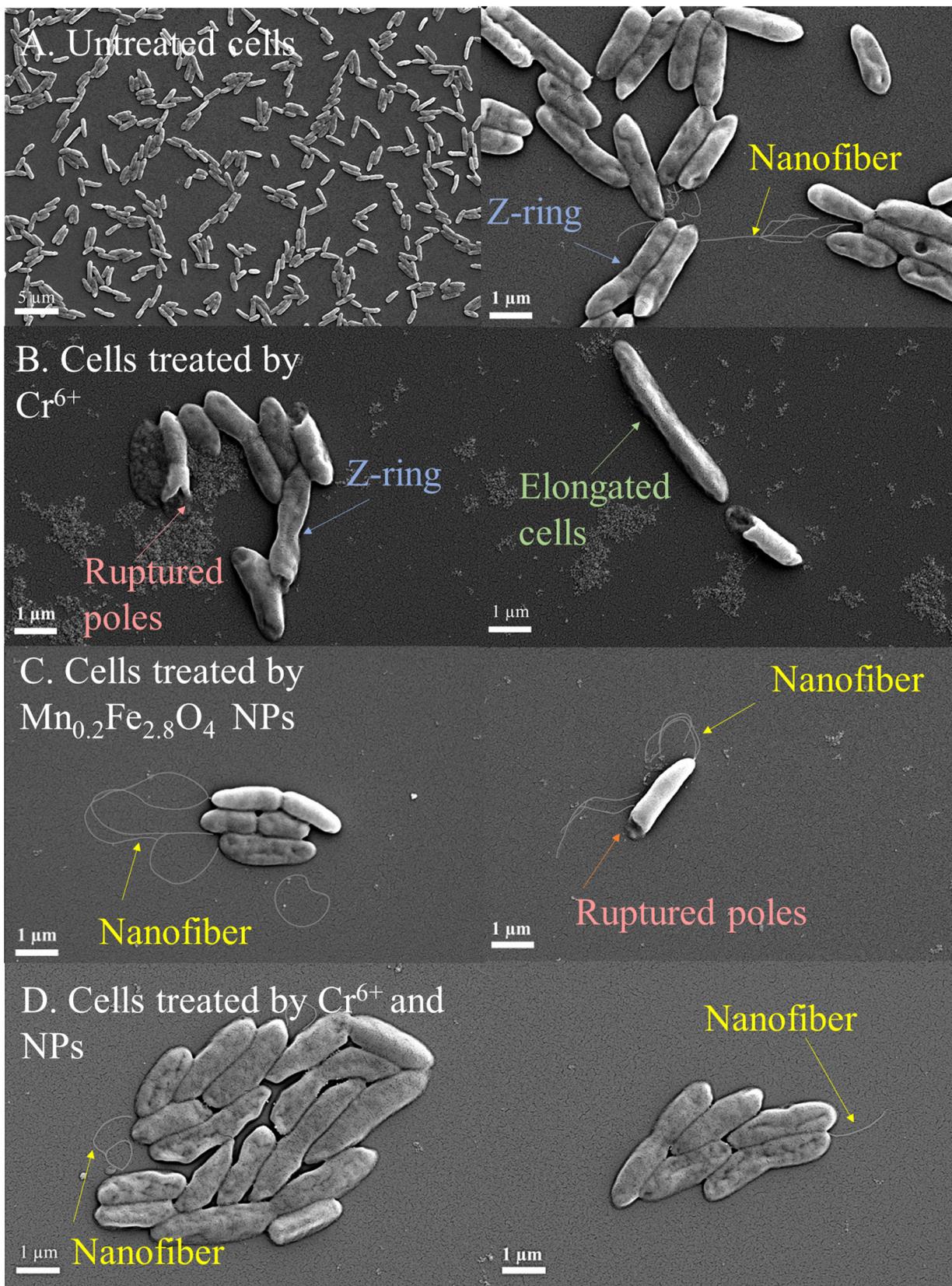


Fig. 4 SEM micrographs of (A) untreated *S. oneidensis* MR-1 cells, (B & C) treated cells by Cr<sup>6+</sup> alone and NPs alone, respectively, and (D) treated cells by both Cr<sup>6+</sup> and NPs.

$\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs provoked the formation of bacterial nanofiber in the presence of  $\text{Cr}^{6+}$ , as depicted in Fig. 4D. Nanofibers are extensions of the outer membrane and periplasm, which are the extracellular electron transport components.<sup>66</sup> Nanofibers are important for long-range extracellular electron transfer.<sup>53,66</sup> The ability of NPs to regenerate bacterial nanofiber production agrees with the findings reported by Yu *et al.*<sup>53</sup> Such observation in response to the interaction between  $\text{Mn}_x\text{Fe}_{3-x}\text{O}_4$  NPs and cells was confirmed in the present work by electron microscopy.

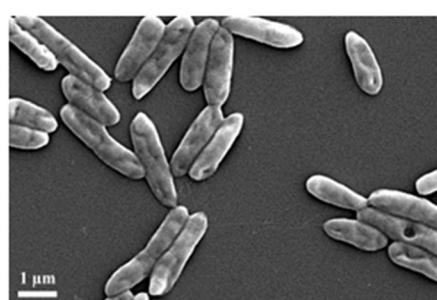
So, Fig. 5 summarises the protective role of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs to *S. oneidensis* MR-1 bacterial cells during  $\text{Cr}^{6+}$ . The use of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs improved the viability of *S. oneidensis* MR-1 under a sub-lethal concentration of  $\text{Cr}^{6+}$  by 3.3 times, as shown in our previous report.<sup>23</sup> Employing  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs as the adsorbent can limit the availability of  $\text{Cr}^{6+}$  to *S. oneidensis* MR-1, boosting the tolerance to  $\text{Cr}^{6+}$ .<sup>18,69</sup> The positive adsorptive effect of NPs on  $\text{Cr}^{6+}$  concerning the viability of bacteria has been reported in the presence of goethite, humic acid,<sup>34</sup> and ferric oxyhydroxide mediators.<sup>18,69</sup> As reported in our recent investigation,<sup>23</sup>  $\text{Cr}^{6+}$  was adsorbed on  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs following the Langmuir adsorption isotherm model. Based on this model, the adsorption and desorption rates should be equal. Adsorption is the separation of molecules from the aqueous solution by being attached to the surface

of the adsorbent. The desorption is inversely related to adsorption processes, where adsorbates are transferred from the adsorbed state to bulk solution.<sup>78</sup> This possible continuous adsorption–desorption rate of  $\text{Cr}^{6+}$  can sustain a release of  $\text{Cr}^{6+}$  from the surface of NPs, which makes the exposure of cells to  $\text{Cr}^{6+}$  occur at a gradual rate.

Furthermore,  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs reduce  $\text{Cr}^{6+}$  into  $\text{Cr}^{3+}$ , as shown in Fig. 3B.  $\text{Cr}^{3+}$  is less toxic than  $\text{Cr}^{6+}$  towards *S. oneidensis* MR-1.<sup>22</sup> Bacterial cells exposed to  $\text{Cr}^{3+}$  experienced viability loss but maintained some enzymatic activity and cellular integrity,<sup>22</sup> which explains the morphological response of *S. oneidensis* MR-1 to  $\text{Cr}^{6+}$  in the presence of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs, as shown in Fig. 4B.

## 4. Conclusion

This study describes a possible protecting role of manganese ferrite nanoparticles ( $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs) to *Shewanella oneidensis* (*S. oneidensis*) MR-1 during hexavalent chromium ( $\text{Cr}^{6+}$ ) bio-reduction.  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs can reduce the highly toxic  $\text{Cr}^{6+}$  to less toxic  $\text{Cr}^{3+}$ . Under anaerobic conditions, we found that  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs induced the elongation of the bacterial cells and promoted the formation of nanofibers. Such morphological change could improve the viability of *S. oneidensis* MR-1 cells in response to the sub-lethal dose of  $\text{Cr}^{6+}$  and, in turn, enhance



*Shewanella oneidensis* MR-1

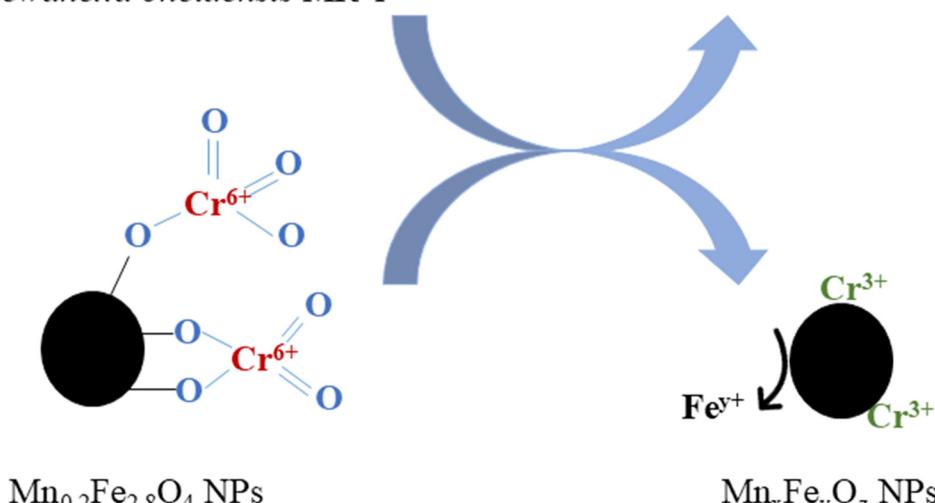


Fig. 5 Illustration of the protective role of  $\text{Mn}_{0.2}\text{Fe}_{2.8}\text{O}_4$  NPs to *S. oneidensis* MR-1 during  $[\text{CrO}_4]^{2-}$  bio-reduction.



their detoxification ability. Integrating both *S. oneidensis* MR-1 and  $Mn_{0.2}Fe_{2.8}O_4$  NPs enhanced  $Cr^{6+}$  detoxification by 2.1-fold compared to *S. oneidensis* MR-1 alone and 1.4-fold compared to NPs alone. Therefore, the present article provides evidence of  $Cr^{6+}$  bio-reduction and the bacterial response to  $Cr^{6+}$  and  $Mn_{0.2}Fe_{2.8}O_4$  NPs. This study will open a venue for applying nanotechnology in the bio-remediation of highly contaminated sites by heavy metals.

## Data availability

The data within this study is included in either the main article or ESI† figures.

## Author contributions

N. T. K. T. and L. C. devised and coordinated the project and provided resources. D. S. R. designed and did most of the experiments and wrote the manuscript. I. T. assisted in particle synthesis and data analysis. N. T. K. T. and S. M. provided expertise, revised the manuscript and helped to acquire funding. E. D. carried out XPS characterisation, processed data and corrected the manuscript. A. M. did a part of the characterisation and edited the manuscript.

## Conflicts of interest

The authors declare no competing financial interest.

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