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# Ti<sub>3</sub>C<sub>2</sub> nanosheets with broad-spectrum antioxidant activity for cytoprotection against oxidative stress†

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Redox regulation in biological systems represents a fascinating method for treatment and prevention of oxidative stress induced diseases. The key and difficult point is to find ideal materials with excellent antioxidant capability and good biocompatibility. To this end, ultra-thin two-dimensional MXene (Ti<sub>3</sub>C<sub>2</sub>) nanosheets (NSs) were investigated for their antioxidant capability. It is found that Ti<sub>3</sub>C<sub>2</sub> NSs can scavenge efficiently reactive oxygen and nitrogen species (\*OH, H<sub>2</sub>O<sub>2</sub>, O<sub>2</sub><sup>-</sup> and \*NO), ABTS<sup>•+</sup> and DPPH<sup>•</sup> free radicals in a concentration dependent manner, showing broad-spectrum antioxidant activities. Ti<sub>3</sub>C<sub>2</sub> NSs exhibit higher antioxidant activity and broader antioxidant capability than natural antioxidant molecules. The significant role of PEG modified Ti<sub>3</sub>C<sub>2</sub> with good stability in preventing cell damage against oxidative stress was demonstrated. Upon treatment of H<sub>2</sub>O<sub>2</sub> induced oxidative stress with Ti<sub>3</sub>C<sub>2</sub>, the intracellular ROS level decreases and the cell survival rate increases significantly. An antioxidant mechanism based on gradient oxidation was proposed to account for the superior antioxidant activity of Ti<sub>3</sub>C<sub>2</sub>. Our result proves that ultra-thin MXenes as antioxidants have great potential in preventing oxidative stress caused biological damage.

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## 1. Introduction

Oxidative stress is a state of imbalance between oxidation and antioxidation in biological systems, and is considered to be one of the critical factors leading to aging and many intractable diseases.<sup>1–3</sup> Reactive oxygen species (ROS) and reactive nitrogen species (RNS), including superoxide (O<sub>2</sub><sup>-</sup>), hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), hydroxyl radicals (\*OH), \*NO and ONOO<sup>-</sup>, play important roles in cellular life cycles.<sup>4</sup> Unfortunately, the over-production of ROS and RNS has been recognized as the direct cause of damage to cellular components (cell membranes, protein structures, and DNA) and their functions.<sup>5</sup> Redox regulation represents a fascinating way to relieve oxidative stress. The key point is to find suitable materials with excellent antioxidant capability and low toxicity.

There are effective ways to alleviate oxidative stress through antioxidant small molecule, enzymes and enzyme-like nanomaterials. A number of literatures based on *in vitro* and *in vivo* experiments provide evidences that antioxidant molecules

represent a rational curative strategy to reduce ROS/RNS as well as prevent and cure oxidative stress induced diseases.<sup>6,7</sup> Natural antioxidants suffer from their low antioxidant efficiency, and sometimes their oxidation can yield unfriendly products. Natural enzymes signify an important role in defense oxidative stress, but the “paradoxical” roles of antioxidant enzymes triggering metabolic disorders also was revealed.<sup>8</sup> In addition, the enzymes work only to catalyze the reduction of specific oxidative species, such as superoxide dismutase (SOD) and catalase. Enzymelike nanomaterials have been named as nanozymes because of their intrinsic activity to mimic natural enzymes.<sup>9–12</sup> Most reported nanozymes belong to the oxidoreductases family and show great potential in regulating ROS level.<sup>13–20</sup> In addition, nanozymes (*e.g.* Pt based and metal oxide) often function to mimic multiple redox enzymes (oxidase, SOD, peroxidase and catalase).<sup>21–25</sup> Such multiple enzymelike activity enables nanozymes with antioxidant activity while also having pro-oxidative functions. Therefore, it is highly required to develop ideal antioxidant nanomaterial with high efficiency and broad spectrum for biomedical study and clinical applications.

MXenes have been found as a family of two-dimensional (2D) metal carbides and nitrides, which show tunable electronic, optical and catalytic properties.<sup>26–30</sup> Because of good biocompatibility and low cytotoxicity, MXenes have been explored for biomedical applications, *e.g.* antibacterial, cancer therapeutics and diagnosis.<sup>31–33</sup> Especially, few or monolayer MXenes are very sensitive to be oxidized because of abundant low-valence metal species, and can be used as antioxidant to defense the oxidative damage.<sup>34</sup> Recently, V<sub>2</sub>C, Ti<sub>3</sub>C<sub>2</sub> and Nb<sub>2</sub>C MXenes have been

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reported to exhibit ROS scavenging activity and show potential in alleviation of ROS induced neurons and acute kidney injury.<sup>35–40</sup> But we still know very little about their antioxidant behavior, for example, the antioxidant mechanism and the antioxidant capability toward various active species and free radicals. To fully understand the antioxidant capability and mechanism of Ti<sub>3</sub>C<sub>2</sub> for better applications, we systematically evaluate their antioxidant capability toward ROS, RNS and free radicals as well as demonstrate the potential in preventing oxidative stress induced cell damage. The primary objective of this study is to investigate the high-efficiency and broad-spectrum antioxidant activity of Ti<sub>3</sub>C<sub>2</sub> nanosheets and understand their antioxidant mechanism.

## 2. Experimental

### 2.1 Chemical and materials

Ti<sub>3</sub>AlC<sub>2</sub> MAX powder (400 mesh) was purchased from 11 Technology, China. Hydrofluoric acid (HF), hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), horseradish peroxidase (HRP) and xanthine oxidase (XOD) were purchased from Sinopharm Chemical Reagent Co., Ltd (Beijing, China) and 2,2'-azinobis-(3-ethylbenzthiazoline-6-sulphonate) (ABTS), (S)-nitroso-N-acetylpenicillamine (SNAP), and ferrous sulfate (FeSO<sub>4</sub>) were commercially available from Aladdin Industrial Co. (CA, USA). 1,1-Diphenyl-2-picrylhydrazyl radical 2,2-diphenyl-1-(2,4,6-trinitrophenyl)hydrazyl (DPPH) was purified from Shanghai Macklin Biochemical Co., Ltd. Tetramethyl ammonium hydroxide (TMAOH), diethylenetriaminepentaacetic acid (DTPA) and xanthine were purchased from Sigma-Aldrich (Shanghai, China). 5,5-Dimethyl-1-pyrroline-N-oxide (DMPO) and 5-tert-butoxycarbonyl-5-methyl-1-pyrroline-N-oxide (BMPO) were purchased from Tongren Institute of Chemistry. 2-Phenyl-4,4,5,5-tetramethylimidazole-*q*-oxyl 3-oxide (PTIO) was obtained from Shanghai Maokang Biotechnology Co., Ltd. Milli-Q water (18 MΩ cm) was used in the preparation of all solutions.

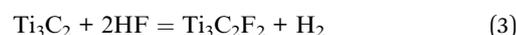
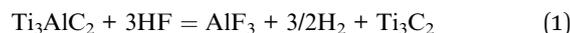
### 2.2 Characterization

UV-Vis absorption spectra were obtained using a Cary 5000 UV-Vis spectrometer (Varian, USA) and a matched quartz cuvette with a path length of 1 cm. The crystal structures of the Ti<sub>3</sub>C<sub>2</sub> NSs were characterized by X-ray diffraction (XRD, D8 Advance diffractometer, Bruker, Germany) using monochromatized Cu Kα radiation (λ = 1.5418 Å). Transmission electron microscope (TEM) and high resolution TEM (HRTEM) images were captured on a Tecnai G2 F20 U-Twin electron microscope with an accelerating voltage of 200 kV. X-ray photoelectron spectroscopy (XPS) was performed with a Thermo ESCALAB 250XI multi-functional imaging electron spectrometer (Thermo Fisher Scientific, USA) using 150 W Al K<sub>α</sub> radiation and a base pressure of approximately 3 × 10<sup>-9</sup> mbar. The binding energies were calibrated to the C 1s line at 284.8 eV. Raman spectra was performed with a Renishaw Raman microscope (inVia, Renishaw, UK) using an excitation laser with 532 nm wavelength. AFM images were tested by Dimension Icon produced by Bruker Germany and zeta potential was tested using Malvern Zetasizer

Nano ZS90. Electron spin-resonance (ESR) spectra were obtained using an EMX micro electron paramagnetic resonance spectrometer (Bruker, Germany). The measurements were carried out at room temperature under the following conditions: microwave power 2 mW, modulation amplitude 1.0 G, attenuation 30 dB, and scan range of 100 G.

### 2.3 Synthesis of multilayer Ti<sub>3</sub>C<sub>2</sub>

10 mL 50 wt% HF was added to a 25 mL Teflon tank, and 1 g Ti<sub>3</sub>AlC<sub>2</sub> was added to the solution gradually and carefully due to the exothermic reaction between Ti<sub>3</sub>AlC<sub>2</sub> and HF solution. The reaction between Ti<sub>3</sub>AlC<sub>2</sub> and HF follows the eqn (1)–(3).<sup>26</sup>



After stirring at room temperature for 24 h, the solution was centrifuged at 3500 rpm for 10 min until the pH > 6. The products were dried under vacuum at 80 °C for 24 h. HF is very dangerous, it should be extremely careful during operation.

### 2.4 Synthesis of 2D Ti<sub>3</sub>C<sub>2</sub> nanosheets

0.2 g multilayer Ti<sub>3</sub>C<sub>2</sub> powder was dispersed in 60 mL aqueous solution of 2 wt% TMAOH for 30 min, and 0.352 g AA was added. After mixing, the resulting mixture was transferred to a 100 mL Teflon stainless steel autoclave, sealed and placed in oven at 140 °C for 24 h.<sup>41</sup> Then, the TMAOH solution and AA were removed by centrifugation at 10 000 rpm for three times. The black precipitate was resuspended into 60 mL deionized water and transferred to a conical flask, and ultrasonic was performed in argon atmosphere for 2 h. Then, a colloidal solution of 2D Ti<sub>3</sub>C<sub>2</sub> nanosheets was collected by centrifugation for 20 min at 8000 rpm. The 2D Ti<sub>3</sub>C<sub>2</sub> nanosheets were collected *via* drying in a vacuum freeze dryer for at least 24 h. The resulting powder was dispersed to 1 mg mL<sup>-1</sup> in deionized water.

### 2.5 Synthesis of Ti<sub>3</sub>C<sub>2</sub>-PEG

In order to completely coat the Ti<sub>3</sub>C<sub>2</sub> surface with PEG molecules, an excessive amount of PEG was used with a PEG/Ti<sub>3</sub>C<sub>2</sub> mass ratio of 20 : 1. 0.1 g of PEG-6000 were dispersed into 5 mL water and mixed with 5 mL Ti<sub>3</sub>C<sub>2</sub> nanosheets solution (1 mg mL<sup>-1</sup>). The aqueous solution was kept stirring overnight. The product was centrifuged three times at 13 000 rpm for 30 min to remove excess PEG molecules. The collected black precipitate was freeze-dried, dispersed to 0.5 mg mL<sup>-1</sup> in deionized water.

### 2.6 Synthesis of Ti<sub>3</sub>C<sub>2</sub>-A and Ti<sub>3</sub>C<sub>2</sub>-B (the Ti<sub>3</sub>C<sub>2</sub> nanosheets after oxidation by ABTS<sup>•+</sup> and H<sub>2</sub>O<sub>2</sub>)

2 mL freshly made Ti<sub>3</sub>C<sub>2</sub> (1 mg mL<sup>-1</sup>) was stirred with excess 2,2'-azinobis-(3-ethylbenzthiazoline-6-sulphonate) free radicals (ABTS<sup>•+</sup>) overnight. Then, the precipitation was collected by centrifugation three times at 8000 rpm for 10 min. 4 mL deionized water was added into the precipitation to prepare



Ti<sub>3</sub>C<sub>2</sub>-A with a final concentration of 0.5 mg mL<sup>-1</sup>. Following the similar procedure, Ti<sub>3</sub>C<sub>2</sub>-B (Ti<sub>3</sub>C<sub>2</sub> reacting with H<sub>2</sub>O<sub>2</sub>) with concentration of 0.5 mg mL<sup>-1</sup> was prepared by stirring, centrifugation and constant volume.

## 2.7 Measurement of radical scavenging activity of Ti<sub>3</sub>C<sub>2</sub> NSs

**2.7.1 ABTS<sup>•+</sup> and DPPH<sup>•</sup> scavenging activity.** ABTS and DPPH assays are used to measure the total antioxidant capacity of Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG NSs. 0.0374 g ABTS powders was dissolved into 30 mL deionized water, followed by addition of 0.0066 g K<sub>2</sub>S<sub>2</sub>O<sub>8</sub> under stirring and dark reaction for 12 h. To test the ABTS<sup>•+</sup> reduction, 0.5 mg mL<sup>-1</sup> Ti<sub>3</sub>C<sub>2</sub> or Ti<sub>3</sub>C<sub>2</sub>-PEG with varied volume (3, 5, 7, 10, 15 μL) is added into 3 mL ABTS<sup>•+</sup> solution (O.D. = ~1.2). To test the DPPH<sup>•</sup> reduction, 2.5 mL of 1 mg mL<sup>-1</sup> DPPH<sup>•</sup> ethanol solution was added with 0.5 mL of deionized water. Then, 0.5 mg mL<sup>-1</sup> Ti<sub>3</sub>C<sub>2</sub> or Ti<sub>3</sub>C<sub>2</sub>-PEG with different volumes of (3, 5, 7, 10, 15 μL) were added. Two minutes later, the UV-Vis spectra were recorded to measure the reduction of ABTS<sup>•+</sup> and DPPH<sup>•</sup>.

**2.7.2 H<sub>2</sub>O<sub>2</sub> scavenging activity.** The H<sub>2</sub>O<sub>2</sub> scavenging ability of Ti<sub>3</sub>C<sub>2</sub> was analyzed by HRP-H<sub>2</sub>O<sub>2</sub>-TMB system. 20 μL H<sub>2</sub>O<sub>2</sub> (0.1 M) is firstly mixed with Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG with desirable concentrations in 3 mL H<sub>2</sub>O, 2 min later, 20 μL TMB (20 mM) and 10 μL HRP (1 μg mL<sup>-1</sup>) were added into the above mixture to initiate the TMB oxidation. After 3 min of reaction, the absorption spectra showing characteristic absorption at 652 nm was recorded by UV-Vis spectrometer.

**2.7.3 <sup>•</sup>OH scavenging activity.** The <sup>•</sup>OH was generated by classic Fenton reaction between Fe<sup>2+</sup> and H<sub>2</sub>O<sub>2</sub>, and the production of <sup>•</sup>OH is monitored by both UV-Vis spectra and ESR. For UV-Vis spectra test, 20 μL TMB (20 mM), 20 μL H<sub>2</sub>O<sub>2</sub> (0.1 M) and 20 μL FeSO<sub>4</sub> (2 mM) were mixed in 3 mL H<sub>2</sub>O, then (3, 5, 7, 10, 15 μL) 0.5 mg mL<sup>-1</sup> Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG solution was added. For ESR test, 10 mM H<sub>2</sub>O<sub>2</sub> was mixed with 50 mM DMPO and 10 mM Fe<sup>2+</sup> in the absence and presence of Ti<sub>3</sub>C<sub>2</sub> with different concentrations or antioxidant agents. At selected time intervals, the UV-Vis spectra and ESR spectra was recorded.

**2.7.4 SOD-like activity.** ESR was used to test SOD-like activity to scavenge superoxide of Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG. Typically, 10 μL xanthine (5 mM) is mixed with 10 μL DTPA (0.25 mM), 5 μL BMPO (250 mM) and Ti<sub>3</sub>C<sub>2</sub> or Ti<sub>3</sub>C<sub>2</sub>-PEG with different concentrations, then 5 μL XOD (0.4 U mL<sup>-1</sup>) was added to initiate the production of superoxide anion. The SOD-like enzyme activity of antioxidant was analyzed by BMPO/<sup>•</sup>OOH signal changes.

**2.7.5 <sup>•</sup>NO scavenging activity.** The <sup>•</sup>NO scavenging activity of Ti<sub>3</sub>C<sub>2</sub> was tested by ESR. <sup>•</sup>NO was produced SNAP and was trapped by PTIO<sup>•</sup> to form typical ESR signals. For the ESR measurement, 10 μL SNAP (2 mM) was mixed with 10 μL PTIO<sup>•</sup> (2 μM) in PBS solution in the absence and presence of Ti<sub>3</sub>C<sub>2</sub> or Ti<sub>3</sub>C<sub>2</sub>-PEG or antioxidant molecule.

## 2.8 Cell culture

Mouse fibroblast cell line NIH3T3 was purchased from the Cell Center of Institute of Basic Medical Sciences, Chinese Academy of Medical Sciences & Peking Union Medical College (Beijing,

China). NIH3T3 was cultured in high-glucose basal Dulbecco's modified Eagle's medium (DMEM) (Hyclone) containing 10% fetal bovine serum (FBS) (Invitrogen), 4500 mg L<sup>-1</sup> glucose, 4 mM L-glutamine, and 0.1% streptomycin and penicillin G at 37 °C with 5% CO<sub>2</sub>.

## 2.9 Cell viability assay

NIH3T3 were seeded into 96-well cell culture plates at a density of 1 × 10<sup>4</sup> cells per well and cultured 24 h for adherence. The cells were incubated with 100 μL DMEM containing different concentration of Ti<sub>3</sub>C<sub>2</sub>-PEG or/and 25 μM H<sub>2</sub>O<sub>2</sub> at 37 °C for 24 h, followed by washing with PBS. Photograph were acquired using an EVOS M7000 imaging system (Thermo Scientific). After the cells incubated with 110 μL fresh medium containing 10 μL Cell Counting Kit-8 (CCK-8, Dojindo Molecular Technologies, Inc.) for about 2 h, the absorbance of the medium was measured at 450 nm using the Varioskan LUX multimode microplate reader (Thermo Scientific). The measurements were carried out in three parallel lines, and the relative cell viability was expressed as a percentage of the control.

## 2.10 Intracellular ROS measurement

NIH3T3 were seeded at a density of 1 × 10<sup>5</sup> cells per well in the 24-well cell culture plate and kept overnight to adhere. The cells were treated with 25 μM H<sub>2</sub>O<sub>2</sub> in the absent and present of 5 μg mL<sup>-1</sup> Ti<sub>3</sub>C<sub>2</sub>-PEG for 24 h at 37 °C and then washed with PBS. After that, PBS containing 10 μM DCFH-DA (Sigma-Aldrich, USA) was added at 37 °C for 30 min and then washed with PBS twice. Photograph was acquired using an EVOS M7000 imaging system. Finally, the cells were detached from the plates with trypsin, and collected by centrifugation. Flow cytometric analysis was conducted on all the groups, using an excitation wavelength of 488 nm and emission wavelength of 525 nm (AccuriTM C6 flow cytometer, BD Biosciences, San Jose, CA). All groups had three replicate wells.

## 2.11 Statistical analysis

The data are shown as mean ± standard deviation (SD) for all treatment groups. Statistical significance was ascertained through one way ANOVA with SPSS software (SPSS17.0).

# 3. Results and discussion

## 3.1 Fabrication and characterization of Ti<sub>3</sub>C<sub>2</sub> NSs

Few-layered Ti<sub>3</sub>C<sub>2</sub> MXene nanosheets (NSs) were synthesized by a typical exfoliation and intercalation process (Fig. 1a).<sup>41,42</sup> The tight "sandwich" structure of the Ti<sub>3</sub>AlC<sub>2</sub> MAX-phase was confirmed by SEM image (Fig. 1b). Multiple-layered Ti<sub>3</sub>C<sub>2</sub> with accordion structure were obtained *via* selective etching of Al layer by using HF solution (Fig. 1c). Then, few-layered Ti<sub>3</sub>C<sub>2</sub> nanosheets were formed by further TMAOH intercalation, hydrothermal and ultrasonic treatment under anti-oxidation protection. The as-prepared Ti<sub>3</sub>C<sub>2</sub> nanosheets are well dispersed in aqueous solution due to abundant hydrophilic groups on surface. TEM image shows the few-layer structure of Ti<sub>3</sub>C<sub>2</sub> NSs with a lateral dimension of about 200 nm (Fig. 1d).



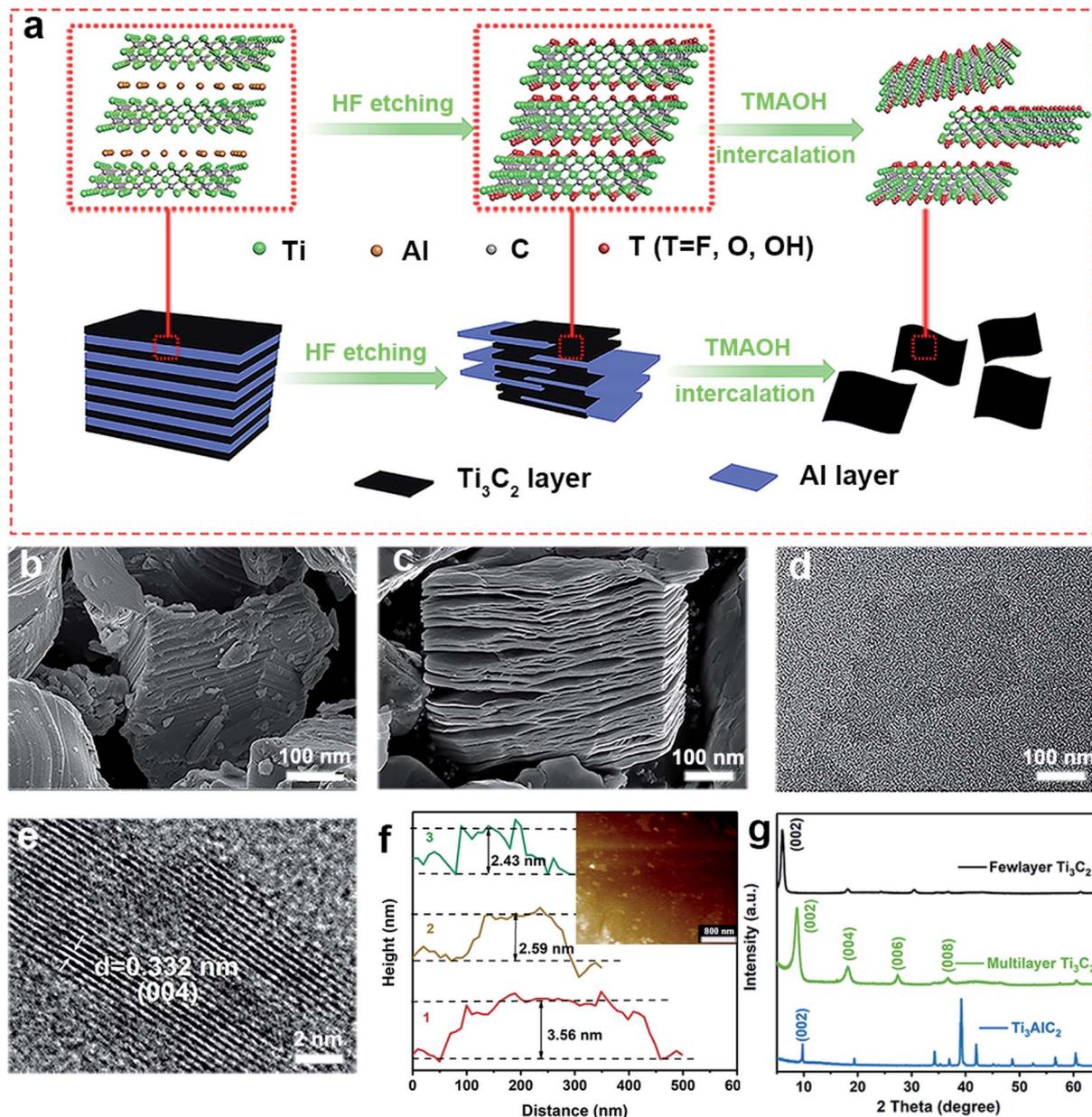


Fig. 1 Fabrication and characterization of  $\text{Ti}_3\text{C}_2$  MXene NSs. (a) Schematic diagram for the procedure to prepare few layered  $\text{Ti}_3\text{C}_2$  NSs, insets illustrate the corresponding chemical structures. SEM image of (b)  $\text{Ti}_3\text{AlC}_2$  and (c) multilayer  $\text{Ti}_3\text{C}_2$ , TEM (d), HRTEM (e) and AFM (f) image of  $\text{Ti}_3\text{C}_2$  NSs, (g) XRD patterns of  $\text{Ti}_3\text{AlC}_2$ , multilayer  $\text{Ti}_3\text{C}_2$  and few layered  $\text{Ti}_3\text{C}_2$  nanosheets.

HRTEM image shows the well-defined lattice planes of selected region (Fig. 1e), and the lattice spacing is calculated to be 0.332 nm, corresponding to the interplanar distance of (004) plane. The averaged thickness of  $\text{Ti}_3\text{C}_2$  NSs is  $\sim 2.59$  nm, denoting that the synthesized  $\text{Ti}_3\text{C}_2$  is composed of two or three layers (Fig. 1f). XRD pattern shows that after HF etching, the diffraction peak of  $\text{Ti}_3\text{AlC}_2$  at  $39^\circ$  disappeared, while the characteristic peak of  $\text{Ti}_3\text{C}_2$  at  $8.8^\circ$  appeared (Fig. 1g). This indicates that Al intercalation is etched away and  $\text{Ti}_3\text{AlC}_2$  is completely converted to multilayer  $\text{Ti}_3\text{C}_2$ . With further stripping, the characteristic peak (002) of multilayer  $\text{Ti}_3\text{C}_2$  shifts to  $6.4^\circ$ , finally forming the  $\text{Ti}_3\text{C}_2$  NSs.<sup>26,41,43</sup>

### 3.2 Antioxidant activity of $\text{Ti}_3\text{C}_2$ NSs by scavenging free radicals, ROS and RNS

The antioxidant efficiency is determined by the category and degree of removing oxidative free radicals.  $\text{ABTS}^{+\cdot}$  radicals is an oxidative product of ABTS.<sup>44</sup>  $\text{DPPH}^\cdot$  is a well-known stable free radical containing unpaired electrons, it has a strong absorption band in the visible region at 517 nm owing to its odd electron, which disappears when this electron is paired up or forms a H-bond in the presence of antioxidants.<sup>45</sup>  $\text{ABTS}^{+\cdot}$  and  $\text{DPPH}^\cdot$  are two standard assays commonly used to evaluate the total antioxidant capabilities of natural antioxidants. The reduction of  $\text{ABTS}^{+\cdot}$  is dominated by the electron transfer



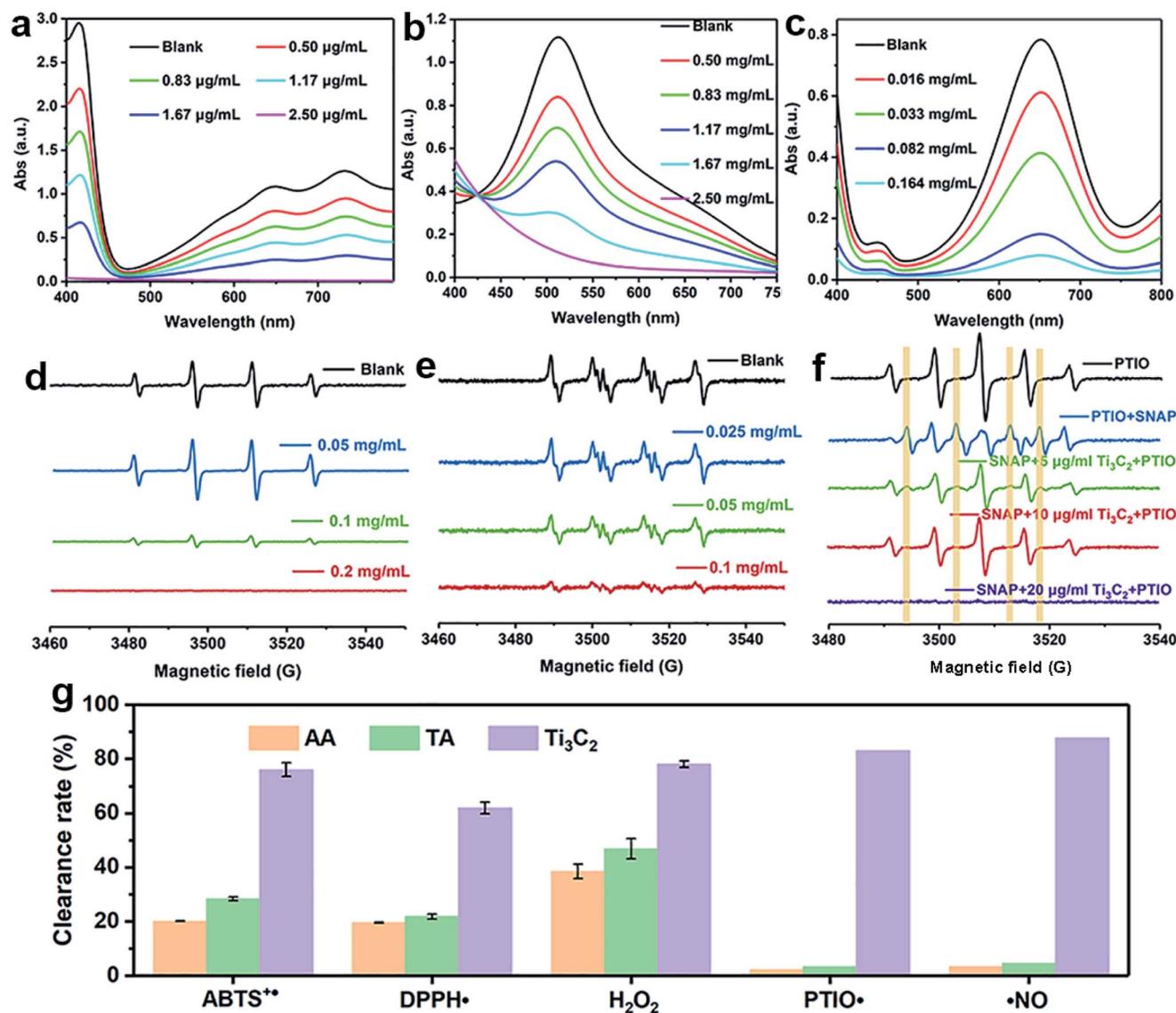


Fig. 2 Antioxidant activity of  $\text{Ti}_3\text{C}_2$ . ABTS<sup>•+</sup> and DPPH<sup>•</sup> scavenging activity: UV-Vis spectra of ABTS<sup>•+</sup> (a) and DPPH<sup>•</sup> solutions (b) in different concentration of  $\text{Ti}_3\text{C}_2$  NSs,  $\text{H}_2\text{O}_2$  scavenging activity (c) UV-Vis spectra of samples containing TMB, HRP and  $\text{H}_2\text{O}_2$  in different concentration of  $\text{Ti}_3\text{C}_2$  NSs, (d) ESR spectra of BMPO/<sup>•</sup>OH generated from the samples containing  $\text{H}_2\text{O}_2$ , BMPO,  $\text{Fe}^{2+}$  and with different concentration of  $\text{Ti}_3\text{C}_2$ , (e) ESR spectra of BMPO/<sup>•</sup>OOH generated from samples containing xanthine, XOD, BMPO and different concentration of  $\text{Ti}_3\text{C}_2$ , (f) ESR spectra of the PTIO<sup>•</sup> solution alone and the PTIO<sup>•</sup> solutions with 'NO-generator' SNAP in the absence or presence of different concentration of  $\text{Ti}_3\text{C}_2$ , (g) comparison of  $\text{Ti}_3\text{C}_2$  with TA and AA in scavenging free radicals, ROS and RNS.

process, while the reduction of DPPH<sup>•</sup> mainly involves the proton transfer process.<sup>46,47</sup> Both ABTS<sup>•+</sup> and DPPH<sup>•</sup> solution show typical color and characteristic absorption band at visible region, this provides a convenient way to compare the antioxidant activity of  $\text{Ti}_3\text{C}_2$  nanosheets by monitoring the change in absorbance. It shows clearly that the addition of  $\text{Ti}_3\text{C}_2$  can efficiently reduce ABTS<sup>•+</sup> and DPPH<sup>•</sup> free radicals (Fig. S1† and b). In contrast, pure ABTS<sup>•+</sup> or DPPH<sup>•</sup> are stable during testing period and do not degrade. The scavenging degree is found to be closely dependent on the concentration of  $\text{Ti}_3\text{C}_2$ . A higher concentration of  $\text{Ti}_3\text{C}_2$  results in more reduction of absorbance, indicating higher removal degree of ABTS<sup>•+</sup> (Fig. S1†) and DPPH<sup>•</sup> (Fig. S2†). This proves that  $\text{Ti}_3\text{C}_2$  has excellent

antioxidant capacity, and its antioxidant mechanism may be related to both electron and proton transfer pathways. Compared the DPPH<sup>•</sup> scavenging ability with other antioxidants or nanoparticles (Table S2†),  $\text{Ti}_3\text{C}_2$  can remove DPPH<sup>•</sup> in a smaller dose and shorter time, which indicates that  $\text{Ti}_3\text{C}_2$  has strong free radical scavenging ability and fast response.<sup>48–51</sup>

Considering ROS play the critical role in regulation of oxidative stress balance, three representative ROS in biological conditions,  $\text{H}_2\text{O}_2$ , <sup>•</sup>OH and  $\text{O}_2^{\cdot-}$ , were selected to investigate systematically the powerful ROS scavenging capabilities of  $\text{Ti}_3\text{C}_2$ . The ability of  $\text{Ti}_3\text{C}_2$  to reduce  $\text{H}_2\text{O}_2$  is evaluated by using HRP- $\text{H}_2\text{O}_2$ -TMB system, which is a classic enzymatic reaction. Under catalysis of HRP, TMB can be quickly oxidized by  $\text{H}_2\text{O}_2$  to



produce typical blue color and characteristic absorption. However, when  $\text{Ti}_3\text{C}_2$  was interacted with  $\text{H}_2\text{O}_2$ , the TMB oxidation degree decreased evidently as reflected by the reduction of characteristic absorption at 652 nm. This indicates that  $\text{Ti}_3\text{C}_2$  can reduce  $\text{H}_2\text{O}_2$  thus inhibit the oxidation of TMB. The  $\text{H}_2\text{O}_2$  reduction activity also shows concentration dependence on the concentration of  $\text{Ti}_3\text{C}_2$ . Higher amount of  $\text{Ti}_3\text{C}_2$  is used to react with  $\text{H}_2\text{O}_2$ , the oxidation of TMB is inhibited to a higher degree (Fig. 2c). In the interaction with nanomaterials, it is well recognized that the reduction of  $\text{H}_2\text{O}_2$  may go through 2 pathways, (1) disproportionation reaction produces  $\text{O}_2$  and  $\text{H}_2\text{O}$ , and (2) reduction reaction produces water. We used dissolved oxygen meter and ESR oximetry to detect whether the oxygen is produced from the mixture of  $\text{H}_2\text{O}_2$  and  $\text{Ti}_3\text{C}_2$ , but no signal from oxygen production was detected. This indicates  $\text{Ti}_3\text{C}_2$  NSs have high chemically reducing ability to  $\text{H}_2\text{O}_2$ .

The antioxidant potential of  $\text{Ti}_3\text{C}_2$  to clear  $\cdot\text{OH}$  and  $\text{O}_2^-$  was evaluated by ESR. Typically, the  $\cdot\text{OH}$  was generated by the Fenton reaction in  $\text{Fe}^{2+}$ - $\text{H}_2\text{O}_2$  system. The DMPO, was used to capture  $\cdot\text{OH}$  and form DMPO/ $\cdot\text{OH}$  spin adduct which shows characteristic ESR peaks with intensity of 1 : 2 : 2 : 1. Compared with control without  $\text{Ti}_3\text{C}_2$ , the addition of  $0.1 \text{ mg mL}^{-1}$   $\text{Ti}_3\text{C}_2$  results in more than 90% reduction of signal intensity,  $0.2 \text{ mg mL}^{-1}$   $\text{Ti}_3\text{C}_2$  can completely scavenge the  $\cdot\text{OH}$  signal (Fig. 2d). This result verifies that  $\text{Ti}_3\text{C}_2$  NSs have strong  $\cdot\text{OH}$  scavenging ability. It is noteworthy that the signal intensity of DMPO/ $\cdot\text{OH}$  increased considerably at the dosage of  $0.05 \text{ mg mL}^{-1}$ , which seems like contrary to the antioxidant capability of  $\text{Ti}_3\text{C}_2$ . This is due to that  $\text{Ti}_3\text{C}_2$  can reduce  $\text{Fe}^{3+}$  to  $\text{Fe}^{2+}$  thus accelerate Fenton

reaction to produce more hydroxyl radicals. The hydroxyl radical generating system by light irradiating  $\text{TiO}_2$  NPs was further employed to confirm the hydroxyl radical scavenging ability of  $\text{Ti}_3\text{C}_2$  in a concentration dependent manner (Fig. S3†).

To evaluate the  $\text{O}_2^-$  scavenging activity of  $\text{Ti}_3\text{C}_2$ , the  $\text{O}_2^-$  was generated by an enzymatic xanthine/XOD system. BMPO is used as a typical spin trap for superoxide. Addition of XOD to the solution containing xanthine, DTPA, and BMPO in  $\text{pH} = 7.4$  PBS buffer produces a strong ESR signal with peak intensity of 1 : 1 : 1 : 1 attributable to BMPO/ $\cdot\text{OOH}$  (Fig. 2e). The ESR signal intensity decreases greatly when  $\text{Ti}_3\text{C}_2$  was added, suggestive of its antioxidant ability to scavenge superoxide. The scavenging activity is highly dependent on the concentration of  $\text{Ti}_3\text{C}_2$ . In comparison,  $0.1 \text{ mg mL}^{-1}$   $\text{Ti}_3\text{C}_2$  NSs can scavenge almost all the superoxide produced, indicating their excellent superoxide scavenging activity.

In order to prove the broad-spectrum antioxidant ability to scavenge free radicals,  $\cdot\text{NO}$  as a representative RNS was used to investigate RNS elimination capabilities of  $\text{Ti}_3\text{C}_2$ . SNAP and PTIO were chosen as the  $\cdot\text{NO}$  donor and label molecule, respectively. PTIO is stable oxygen center free radicals, which has typical five-line ESR spectrum with relative intensity of 1 : 2 : 3 : 2 : 1. PTIO is also a common  $\cdot\text{NO}$  scavenger, which can react with  $\cdot\text{NO}$  to yield PTI, resulting in the reduction of the five-line spectrum and generation of seven-line spectrum. The addition of  $\text{Ti}_3\text{C}_2$  NSs into the mixture of PTIO and SNAP significantly inhibits the formation of PTI under  $\text{pH} 7.4$  (Fig. 2f), and a dosage of  $10 \mu\text{g mL}^{-1}$  can completely remove the  $\cdot\text{NO}$  signal, indicating the superior  $\cdot\text{NO}$  radical scavenging ability of

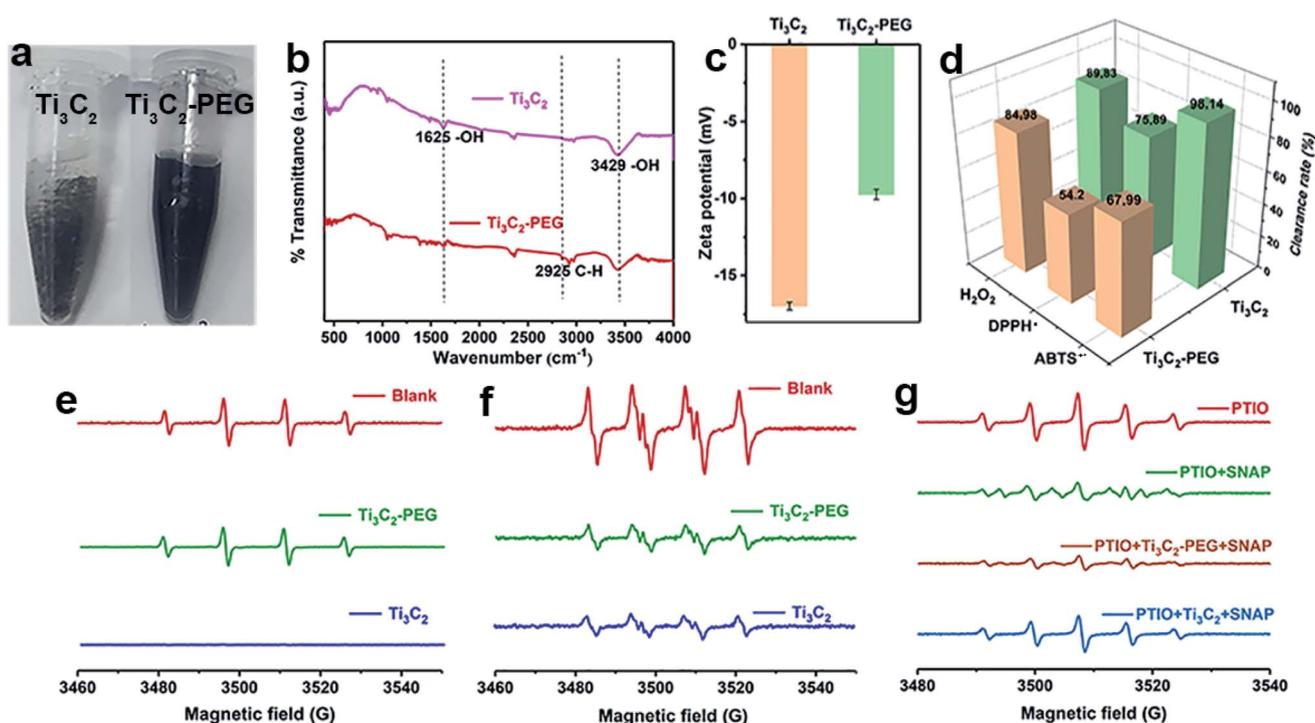


Fig. 3 Stability and antioxidant activity of PEG modified  $\text{Ti}_3\text{C}_2$  ( $\text{Ti}_3\text{C}_2\text{-PEG}$ ). (a) Photographs of pristine  $\text{Ti}_3\text{C}_2$  and  $\text{Ti}_3\text{C}_2\text{-PEG}$  suspended in 100 mM pH 7.4 PBS solution, (b) FTIR spectra and (c) zeta potential of  $\text{Ti}_3\text{C}_2$  and  $\text{Ti}_3\text{C}_2\text{-PEG}$ , (d) comparison of  $\text{Ti}_3\text{C}_2$  and  $\text{Ti}_3\text{C}_2\text{-PEG}$  for their scavenging activity toward  $\text{ABTS}^{\cdot+}$ ,  $\text{DPPH}^{\cdot}$  and  $\text{H}_2\text{O}_2$ , the scavenging activity toward (e)  $\cdot\text{OH}$ , (f)  $\text{O}_2^-$  and (g)  $\cdot\text{NO}$  of  $\text{Ti}_3\text{C}_2$  and  $\text{Ti}_3\text{C}_2\text{-PEG}$ .



Ti<sub>3</sub>C<sub>2</sub> NSs. Interestingly, it is found that the ESR signal of PTIO<sup>•</sup> also disappeared when more Ti<sub>3</sub>C<sub>2</sub> (20 μg mL<sup>-1</sup>) was added, this indicates that Ti<sub>3</sub>C<sub>2</sub> can also scavenge oxygen-centered free radical PTIO<sup>•</sup> (Fig. S4†).

Compared with natural antioxidant molecules at same dosage, we found that Ti<sub>3</sub>C<sub>2</sub> was 2.7 times more potent than TA and 3.3 times more potent than AA in scavenging ABTS<sup>•+</sup> (Fig. 2g). In addition, Ti<sub>3</sub>C<sub>2</sub> is more active than AA and TA in scavenging DPPH<sup>•</sup> and H<sub>2</sub>O<sub>2</sub>. Interestingly, AA and TA show weak ability to reduce PTIO<sup>•</sup> and <sup>•</sup>NO but Ti<sub>3</sub>C<sub>2</sub> can scavenge them efficiently, indicating that Ti<sub>3</sub>C<sub>2</sub> has a broader range scavenging ability than that of conventional antioxidants. Taken into together, Ti<sub>3</sub>C<sub>2</sub> NSs can not only reduce various ROS and RNS (e.g. H<sub>2</sub>O<sub>2</sub>, <sup>•</sup>OH, O<sub>2</sub><sup>-•</sup>, and <sup>•</sup>NO), but also scavenge nitrogen and oxygen-center free radicals, showing more efficient and broad antioxidant capability. The antioxidant capacity is higher than that of natural antioxidants, which is essential for reducing oxidative stress levels to prevent oxidative damage to biological structures.

### 3.3 Stability and antioxidant activity of Ti<sub>3</sub>C<sub>2</sub>-PEG NSs

As discussed above, Ti<sub>3</sub>C<sub>2</sub> with excellent antioxidant capabilities can be an ideal alternative to natural antioxidants. Unfortunately, the as-prepared Ti<sub>3</sub>C<sub>2</sub> NSs are easy to be aggregated in physiochemical environment (e.g. PBS buffer), which limit largely the biological application of Ti<sub>3</sub>C<sub>2</sub>. To overcome this problem, we modify Ti<sub>3</sub>C<sub>2</sub> surface with a biocompatible

molecule (PEG). Fig. 3a shows clearly that PEG modified Ti<sub>3</sub>C<sub>2</sub> (Ti<sub>3</sub>C<sub>2</sub>-PEG) are well dispersed in 100 mM PBS buffer (pH = 7.4) while Ti<sub>3</sub>C<sub>2</sub> NSs are aggregated obviously, demonstrating the successful modification of PEG on Ti<sub>3</sub>C<sub>2</sub> and increase of the stability. The Fourier transform infrared spectra of Ti<sub>3</sub>C<sub>2</sub>-PEG shows a new peak appeared at 2925 cm<sup>-1</sup>, which corresponds to C-H stretching vibration of PEG molecules (Fig. 3b). The surface potential of Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG was measured to be ~-17 and -9.7 mV, respectively (Fig. 3c). The loading of electrically neutral PEG molecules reduces the negative charge on Ti<sub>3</sub>C<sub>2</sub> surface, thus the zeta potential of Ti<sub>3</sub>C<sub>2</sub>-PEG is decreased. These data together give the direct evidences for the successful modification on PEG. To understand how the modification of PEG affect the antioxidant performance, we have further comparatively studied the bare Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG for their ability to scavenge the various free radicals. Fig. 3d–g display the scavenging effect of Ti<sub>3</sub>C<sub>2</sub> and Ti<sub>3</sub>C<sub>2</sub>-PEG at fixed concentration on ABTS<sup>•+</sup>, DPPH<sup>•</sup>, H<sub>2</sub>O<sub>2</sub>, <sup>•</sup>OH, O<sub>2</sub><sup>-•</sup> and <sup>•</sup>NO. Behave like Ti<sub>3</sub>C<sub>2</sub>, Ti<sub>3</sub>C<sub>2</sub>-PEG shows efficient and broad antioxidant activity toward all the kinds of free radicals. In addition, Ti<sub>3</sub>C<sub>2</sub>-PEG has a stable scavenging ability of ABTS<sup>•+</sup> at different pH values (Fig. S5†). As expected, the modification of PEG leads to considerable reduction of the antioxidant activity to varying degree for different free radicals. After PEG modification, the ABTS<sup>•+</sup> and DPPH<sup>•</sup> free radical scavenging rates reduced by ~20%, while the superoxide scavenging degree is comparable. The reduction of antioxidant activity of Ti<sub>3</sub>C<sub>2</sub> due to PEG

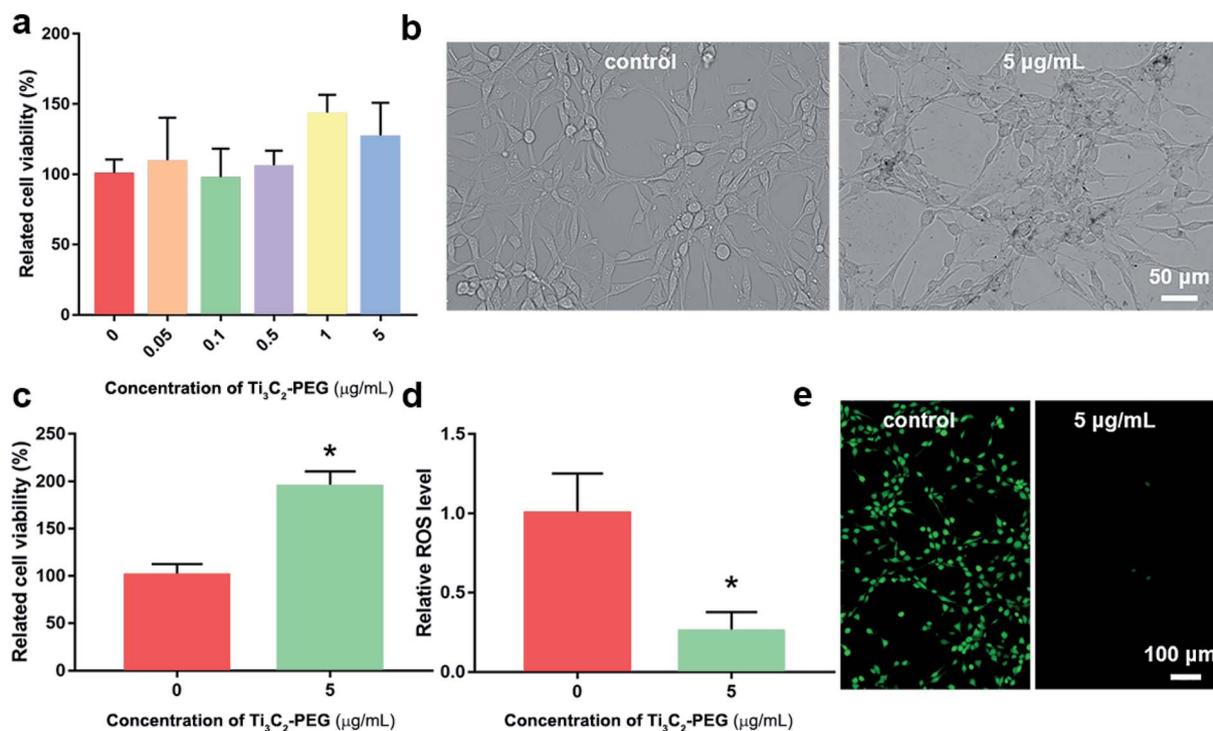


Fig. 4 Protective effect of Ti<sub>3</sub>C<sub>2</sub>-PEG NSs on NIH3T3 cell. (a) Viability of NIH3T3 cell exposed with different concentration of Ti<sub>3</sub>C<sub>2</sub>-PEG NSs from CCK-8 assay, (b) images of NIH3T3 cell in cell medium from optical microscope, (c) viability of NIH3T3 cell exposed to 25 μM H<sub>2</sub>O<sub>2</sub>, (d) the intracellular ROS level of NIH3T3 cell exposed to 25 μM H<sub>2</sub>O<sub>2</sub>, and (e) fluorescence images of NIH3T3 cell labeled with DCFH-DA; \**P* < 0.05 compared with the control group.



modification is inevitable, but it is more suitable for stable and biocompatible use in biological applications.

### 3.4 Cell protection against oxidative damage

In order to investigate the protective role against oxidative stress, the viability and ROS levels of NIH3T3 cell in the absent and present of  $\text{Ti}_3\text{C}_2$ -PEG for 24 h were measured (Fig. 4). Different concentration ranges of  $\text{Ti}_3\text{C}_2$ -PEG NSs were employed firstly to observe their biocompatibility. The cell viability of control group was set as a 100%. As shown in Fig. 4a,  $\text{Ti}_3\text{C}_2$ -PEG NSs did not affect the viability of NIH3T3 cell in the range of testing concentration, indicating their excellent biocompatibility. From the images with optical microscope (Fig. 4b), the dark  $\text{Ti}_3\text{C}_2$ -PEG were observed to exist in the cell medium without changing the morphology of cells, consistent with the good biocompatibility in Fig. 4a.  $\text{H}_2\text{O}_2$ , which was commonly used in inducing oxidative stress, was employed to determine the protective ability of  $\text{Ti}_3\text{C}_2$ -PEG NSs. In the presence of 25  $\mu\text{M}$   $\text{H}_2\text{O}_2$ , the relative cell viability of NIH3T3 cell increased from 100% to 194% after incubated with 5  $\mu\text{g mL}^{-1}$

$\text{Ti}_3\text{C}_2$ -PEG NSs (Fig. 4c), and the relative ROS level was also decreased to 1/4 of the control (Fig. 4d). In fluorescence images of DCFH-DA labeled NIH3T3 cells, the ROS level could be observed directly. As shown in Fig. 4e, the green fluorescence almost disappeared after incubated with  $\text{Ti}_3\text{C}_2$ -PEG NSs, which also confirmed that  $\text{Ti}_3\text{C}_2$ -PEG NSs have excellent elimination ability on ROS. These primary results give a promise of antioxidant  $\text{Ti}_3\text{C}_2$ -PEG NSs in cytoprotection and therapy of oxidative stress induced diseases.

### 3.5 Antioxidant mechanism of $\text{Ti}_3\text{C}_2$ NSs

As we known,  $\text{Ti}_3\text{C}_2$  is easily to be oxidized because of low valence of Ti with unoccupied surface, thus showing excellent reductive potential. For full understanding of the antioxidant mechanism of  $\text{Ti}_3\text{C}_2$  NSs, it is highly required to know how the changes in the surface and valence structure during oxidation. The  $\text{Ti}_3\text{C}_2$  NSs sample before and after oxidation by  $\text{ABTS}^{+\cdot}$  ( $\text{Ti}_3\text{C}_2$ -A) or  $\text{H}_2\text{O}_2$  ( $\text{Ti}_3\text{C}_2$ -B) were employed for comparative study. After oxidation by  $\text{ABTS}^{+\cdot}$ , TEM shows that small particles appeared on the edge of  $\text{Ti}_3\text{C}_2$  nanosheets (Fig. 5a). After

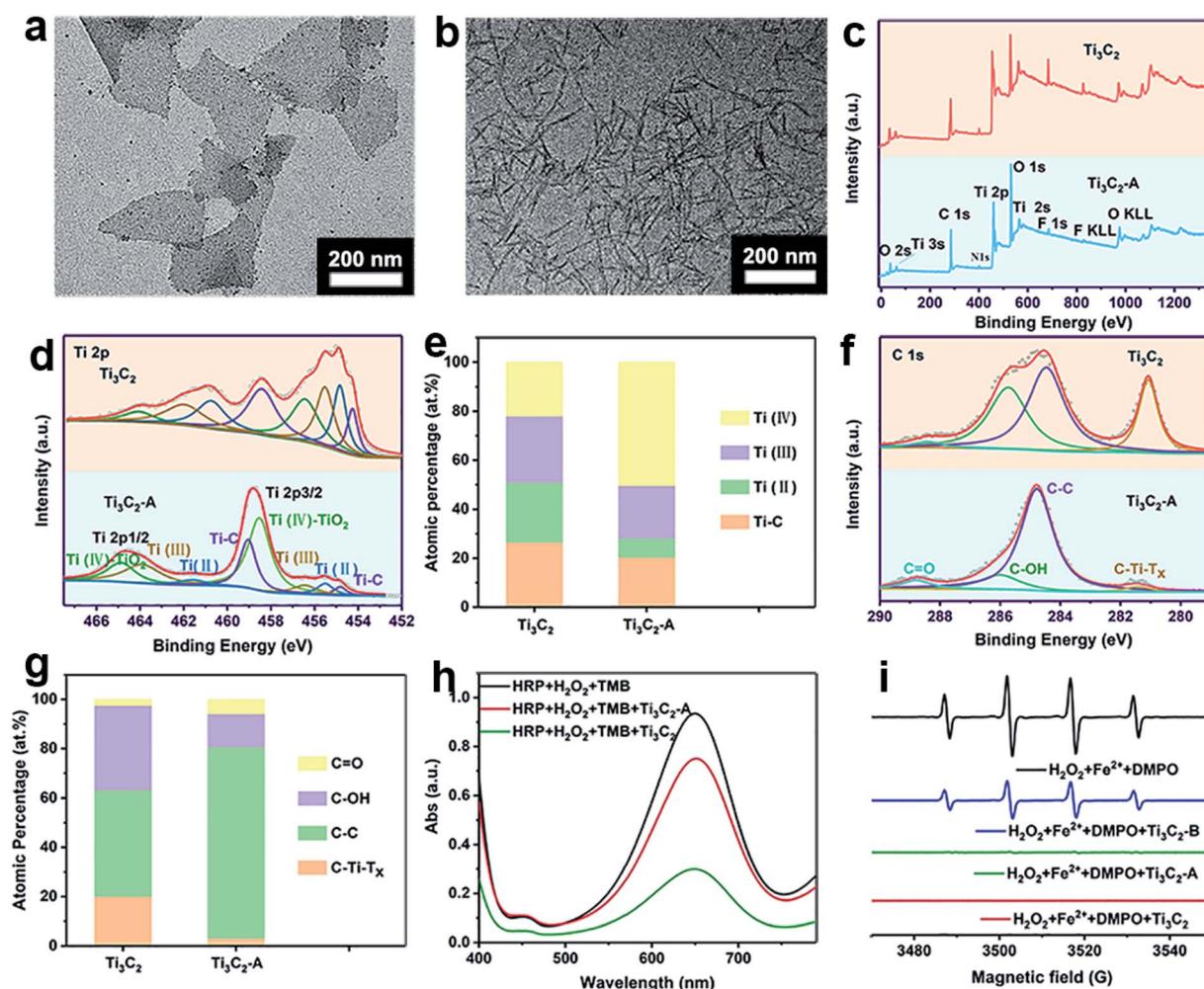


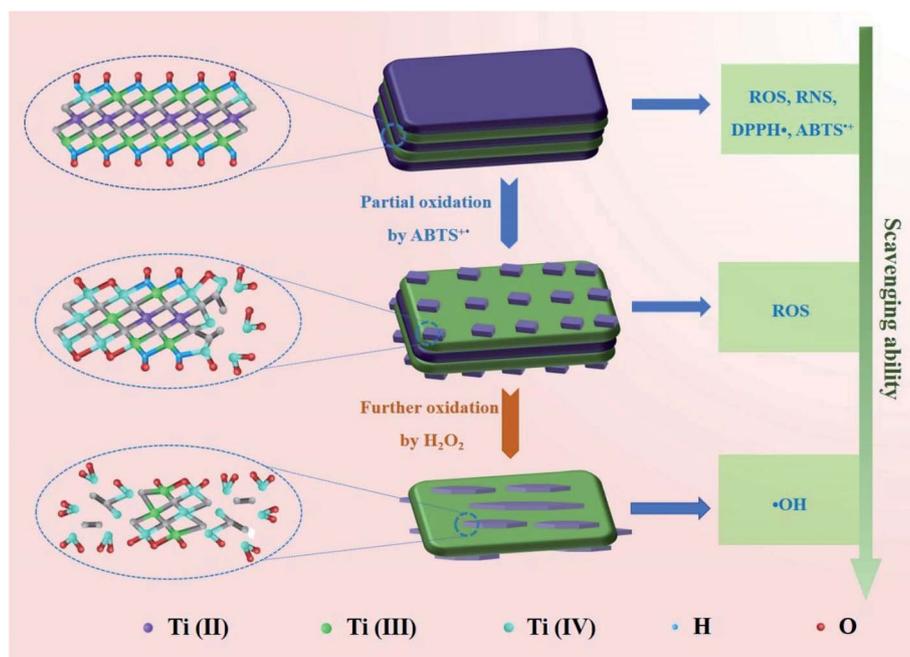
Fig. 5 Antioxidant mechanism of  $\text{Ti}_3\text{C}_2$  NSs. TEM image of  $\text{Ti}_3\text{C}_2$ -A (a) and  $\text{Ti}_3\text{C}_2$ -B (b), (c) XPS survey spectra, (d) high resolution XPS spectra of Ti 2p, (e) atomic percentages of different Ti species, (f) high resolution XPS spectra of C 1s, (g) atomic percentages of C=O, C-OH, C-C and C-Ti, (h) comparison of  $\text{H}_2\text{O}_2$  scavenging and (i)  $\cdot\text{OH}$  scavenging ability of  $\text{Ti}_3\text{C}_2$ ,  $\text{Ti}_3\text{C}_2$ -A and  $\text{Ti}_3\text{C}_2$ -B.



scavenging  $\text{H}_2\text{O}_2$ ,  $\text{Ti}_3\text{C}_2$  NSs have a great change in morphology, which seems to be caused by oxidation, leading to further fragmentation of layers (Fig. 5b). It can be seen from the XPS surveys of  $\text{Ti}_3\text{C}_2$  and  $\text{Ti}_3\text{C}_2\text{-A}$ , the relative intensity of O 1s signal was evidently increased, indicating the oxygen participated in the oxidation of  $\text{Ti}_3\text{C}_2$  during  $\text{ABTS}^{+\cdot}$  clearance (Fig. 5c). Fig. 5d fitted the XPS spectrum of Ti 2p signal for  $\text{Ti}_3\text{C}_2$  and  $\text{Ti}_3\text{C}_2\text{-A}$ , both of which shows the signal from Ti-C, Ti(II), Ti(III) and Ti(IV),<sup>52,53</sup> but exhibit different ratios. The percentage of Ti-C, Ti(II), Ti(III) and Ti(IV) was calculated to be 26.6%, 24.38%, 27.12%, 21.9% for original  $\text{Ti}_3\text{C}_2$ . In the case of  $\text{Ti}_3\text{C}_2\text{-A}$ , the corresponding percentage was 20.42%, 7.86%, 21.37% and 50.35%, respectively (Fig. 5e). Obviously, the proportion of Ti(II) reduced by 3 times while the proportion of Ti(IV) increased by more than 2 times. A slight reduction of the composition of Ti-C and Ti(III) was observed after oxidation. After oxidation with  $\text{H}_2\text{O}_2$  ( $\text{Ti}_3\text{C}_2\text{-B}$ ), the signals from Ti-C and Ti(II) disappeared, all the Ti species were converted into Ti(III) and Ti(IV) with percentage of 46.86% and 53.14%, respectively (Fig. S6†). The chemical bonding environment of atomic C was also analyzed by C 1s core level spectra (Fig. 5f and g). C 1s spectrum was fitted into five deconvoluted Gaussian peaks which corresponds to the C-Ti-T<sub>x</sub>, C-C, C-OH and C=O, respectively. It clearly shows that the signal intensity of peaks C-OH and C-Ti-T<sub>x</sub> decreased greatly after the  $\text{Ti}_3\text{C}_2$  was oxidized, while the peaks C-C and C-O increased significantly. The ratio of C-Ti-T<sub>x</sub>, C-C, C-OH and C=O in  $\text{Ti}_3\text{C}_2$  before and after oxidation was calculated and compared (Fig. 5g). The significant decrease of C-OH and C-Ti content and the increase of C-C may indicate that part of Ti-C bond was broken after oxidation. The increase of Ti-O and the decrease of Ti-O-H in O 1s spectra also verify the oxidation

process of  $\text{Ti}_3\text{C}_2$  and the reduction of surface OH groups (Fig. S7†). The XRD patterns indicate the obvious change in crystal structure of  $\text{Ti}_3\text{C}_2$  after oxidation with either  $\text{ABTS}^{+\cdot}$  or  $\text{H}_2\text{O}_2$  (Fig. S8†). Especially, the diffraction peaks characteristic to anatase in  $\text{Ti}_3\text{C}_2\text{-B}$  reflect the formation of  $\text{TiO}_2$  particles. These results prove not only the oxidation reaction occurring between  $\text{Ti}_3\text{C}_2$  and oxidative species, but also the structure change after oxidation (Fig. S9†).

The gradual transformation of  $\text{Ti}_3\text{C}_2$  nanosheets during oxidation may endow them intrinsic antioxidant capability. Thus, we test whether the  $\text{Ti}_3\text{C}_2$  nanosheets after scavenging  $\text{ABTS}^{+\cdot}$  can scavenge ROS. We collected  $\text{Ti}_3\text{C}_2$  ( $\text{Ti}_3\text{C}_2\text{-A}$ ) after complete oxidation with  $\text{ABTS}^{+\cdot}$ , and found that the  $\text{Ti}_3\text{C}_2\text{-A}$  cannot reduce  $\text{ABTS}^{+\cdot}$  but still can further eliminate  $\text{H}_2\text{O}_2$  and hydroxyl radical. Compared with pristine  $\text{Ti}_3\text{C}_2$ ,  $\text{Ti}_3\text{C}_2\text{-A}$  exhibit significantly decreased but considerable activity to reduce  $\text{H}_2\text{O}_2$  (Fig. 5h). However, it exhibits almost unreduced ability to scavenge hydroxyl radicals (Fig. 5i). The  $\text{Ti}_3\text{C}_2$  nanosheets after oxidation by  $\text{H}_2\text{O}_2$  ( $\text{Ti}_3\text{C}_2\text{-B}$ ), also show evident activity to reduce hydroxyl radicals but decreased considerably. This result indicate that the  $\text{Ti}_3\text{C}_2$  undergoes a gradient oxidation process when interact with oxidizing substances with distinct redox potential. The gradient oxidation event may be related to the abundant and mixed-valence Ti species in  $\text{Ti}_3\text{C}_2$ . When it reduces an oxidant, it still has the ability to remove free radicals with stronger oxidizing power. Scheme 1 illustrates the possible mechanism for gradient oxidation of  $\text{Ti}_3\text{C}_2$  and the corresponding changes in morphology, chemical valences and anti-oxidant ability to scavenge ROS, RNS and free radicals. Based on comprehensive analysis, it can be hypothesized that the anti-oxidant behavior of  $\text{Ti}_3\text{C}_2$  NSs is different from that of small



Scheme 1 Schematic diagram illustrating the mechanism for gradient oxidation of  $\text{Ti}_3\text{C}_2$  MXene and the corresponding changes in scavenging ability of ROS, RNS and free radicals.



molecules and enzymes, but it reflects the characteristics of both. Firstly, because of the Ti species with low valences ( $Ti^{2+}$  and  $Ti^{3+}$ ),  $Ti_3C_2$  NSs show very low redox potential and high reducing ability like natural antioxidant molecules. Secondly, the electron-rich quasi-metallic property induced strong electron transfer ability makes  $Ti_3C_2$  behave like antioxidant nanozymes that can scavenge free radicals. At last, the gradient oxidation behavior accounts for superior and controllable antioxidant capability.

## 4. Conclusions

The efficient and broad-spectrum antioxidant capability of 2D  $Ti_3C_2$  MXene nanosheets have been demonstrated systematically toward scavenging ROS ( $H_2O_2$ ,  $\cdot OH$ ,  $O_2^-$ ), RNS ( $\cdot NO$ ) and free radicals (ABTS $^{+}$ , DPPH $^{\cdot}$  and PTIO $^{\cdot}$ ). Regardless of the antioxidant activity or the types of free radicals that can be scavenged,  $Ti_3C_2$  nanosheets are more powerful than natural antioxidants (AA and TA). The Ti species with unsaturated valence state and the quasi-metallic property induced superior electron transfer ability endow  $Ti_3C_2$  NSs with efficient antioxidant activity. Benefiting from this superior antioxidant performance, we demonstrated *in vitro* that PEG modified  $Ti_3C_2$  with good biocompatibility and stability can effectively protect the normal cells against oxidative stress induced damages. The antioxidant mechanism based on the gradient oxidation of  $Ti_3C_2$  in interaction with oxidative species was revealed. This work demonstrates the application of antioxidant  $Ti_3C_2$  in prevention of ROS induced cell damage, exploring other MXenes with excellent antioxidant capacity and their potential applications in biological protection and treatment of oxidative stress related diseases will be the future direction.

## Conflicts of interest

The authors declare no competing financial interests.

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