

# Cross talk between poly(ADP-ribose) polymerase I methylation and oxidative stress involved in the toxic effect of anatase titanium dioxide nanoparticles

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**Abstract:** Given the tremendous growth in the application of titanium dioxide nanoparticles (TNPs), concerns about the potential health hazards of TNPs to humans have been raised. Poly(ADP-ribose) polymerase 1 (PARP-1), a highly conserved DNA-binding protein, is involved in many molecular and cellular processes. Limited data demonstrated that certain nanomaterials induced the aberrant hypermethylation of PARP-1. However, the mechanism involved in TNP-induced PARP-1 abnormal methylation has not been studied. A549 cells were incubated with anatase TNPs (22.1 nm) for 24 hours pretreatment with or without methyltransferase inhibitor 5-aza-2'-deoxycytidine and the reactive oxygen species (ROS) scavenger  $\alpha$ -lipoic acid to assess the possible role of methylation and ROS in the toxic effect of TNPs. After TNPs characterization, a battery of assays was performed to evaluate the toxic effect of TNPs, PARP-1 methylation status, and oxidative damage. Results showed that TNPs decreased the cell viability in a dose-dependent manner, in accordance with the increase of lactate dehydrogenase activity, which indicated membrane damage of cells. Similar to the high level of PARP-1 methylation, the generation of ROS was significantly increased after exposure to TNPs for 24 hours. Furthermore,  $\alpha$ -lipoic acid decreased TNP-induced ROS generation and then attenuated TNP-triggered PARP-1 hypermethylation. Meanwhile, 5-aza-2'-deoxycytidine simultaneously decreased the ROS generation induced by TNPs, resulting in the decline of PARP-1 methylation. In summary, TNPs triggered the aberrant hypermethylation of the PARP-1 promoter and there was a cross talk between oxidative stress and PARP-1 methylation in the toxic effect of TNPs.

**Keywords:** titanium dioxide nanoparticles, PARP-1, oxidative stress, DNA methylation

## Introduction

Titanium dioxide ( $\text{TiO}_2$ ) is a natural mineral used broadly in domestic and cosmetic products, including antifouling paints, coatings, ceramics, and additives in pharmaceuticals, food colorants, and sunscreen owing to its typical characteristics such as surface adsorption, photo-catalysis, and ultraviolet (UV) absorption. Titanium, either pure or in alloys, is also extensively used for implanted medical devices, such as dental implants, joint replacements, cardiovascular stents, and spinal fixation devices. Titanium dioxide nanoparticles (TNPs) are absorbed through inhalation, ingestion, and dermal penetration into the body, and distributed in important organs such as lungs,<sup>1,2</sup> lymph nodes,<sup>3</sup> brain, liver, and kidneys.<sup>1</sup>

There are growing concerns about the possible influence of TNPs on human health. It has been shown that ultrafine  $\text{TiO}_2$  particles could induce impairment of macrophage function, persistently high inflammatory reactions, and increased pulmonary retention,

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compared with fine  $\text{TiO}_2$ .<sup>4</sup> Particularly, TNPs were reported to induce reactive oxygen species (ROS) generation in many cell types, including rat neuronal cells<sup>5–8</sup> and human hepatoma cells.<sup>9</sup> Indeed, the capacity of TNPs to produce ROS is frequently proposed to account for their toxic effects.<sup>5,10,11</sup> Based on the evidences outlined earlier, it seems that TNPs can exert their toxic effects through the formation of ROS. Hence, TNPs were evaluated by the World Health Organization (WHO)/International Agency for Research on Cancer (IARC) as a Group 2B compound,<sup>12</sup> which is possibly carcinogenic to humans. However, much still remains to be elucidated concerning the possible health effects of TNPs.

DNA methylation, as an important regulatory factor of gene expression,<sup>13</sup> not only is crucial in the development of normal mammalian cells, but also plays a fundamental role in epigenetic silencing of cancer-related genes in tumorigenesis.<sup>14</sup> In general, a low level of DNA methylation is considered an activation of gene, while hypermethylation can contribute to inactivation.<sup>15</sup> The expression of several cancer-related genes has been reported to be silenced by DNA methylation of their promoter regions.

Poly(ADP-ribose) polymerase 1 (PARP-1), a highly conserved DNA-binding protein, is involved in many molecular and cellular processes, which include DNA repair, proliferation, and chromatin modification.<sup>16</sup> PARP-1 was also found in a complex with the DNA methyltransferase DNMT1, the histone H3K9 methyltransferase G9a, and the histone ubiquitin ligase Np95, indicative of a link between poly(ADP-ribosylation) and the epigenome,<sup>17</sup> and was described as a fundamental constituent of the transcription machinery that interacts with and modulates the activities of several transcription factors.<sup>18,19</sup> Chemical materials such as benzene and silica have been reported to cause the methylation of PARP-1 promoters.<sup>20,21</sup> A certain nanomaterial, nano- $\text{SiO}_2$ , was confirmed to induce the aberrant hypermethylation and inactivity of PARP-1.<sup>22</sup> However, the mechanisms responsible for TNP-induced PARP-1 abnormal methylation have not been elucidated.

It was reported that oxidative stress influenced the occurrence of disease through regulating the gene methylation status.<sup>23</sup> Tunc and Tremellen found that oxidative stress was associated with gene hypomethylation, on the contrary, and antioxidant supplements reduced oxidative damage and reverted the gene methylation level.<sup>24</sup> Therefore, this study focused on the regulating mechanisms between oxidative stress and PARP-1 methylation involved in the toxic effect of anatase TNPs. In this study, we analyzed for the first time the epigenetic regulatory mechanism of PARP-1 in TNP-treated cells. After TNP characterization, batteries of assays were performed to evaluate the toxic effect, PARP-1 methylation

status, and oxidative damage in TNP-exposed A549 cells. Results showed that TNPs triggered the generation of ROS and aberrant hypermethylation in the PARP-1 promoter region. We also used the inhibitor 5-aza-2'-deoxycytidine (5-aza) and  $\alpha$ -lipoic acid ( $\alpha$ -LA) to analyze the underlying mechanisms of the toxic effect of TNPs, suggesting that there was a cross talk between oxidative stress and PARP-1 methylation.

## Materials and methods

### Characterization of the TNPs

Commercial anatase TNPs, with a purity of at least 99.7%  $\text{TiO}_2$ , were purchased from Sigma-Aldrich Co. (St Louis, MO, USA). The primary particle size was  $<25$  nm with a specific surface area of 45–55  $\text{m}^2/\text{g}$ . The particle size and distribution of the TNPs were measured by transmission electron microscope (TEM) (JEOL, Tokyo, Japan) and Image J software. The hydrodynamic sizes and zeta potential of TNPs in ultrapure water as stock media and serum-free Roswell Park Memorial Institute 1640 (RPMI 1640) as culture media were examined by Zetasizer (Malvern Nano-ZS90; Malvern, UK). Suspensions of TNPs were dispersed by a sonicator (Bioruptor UCD-200; Diagenode, Liège, Belgium) at 300 W for 10 minutes before adding to dispersion media in order to minimize their aggregation.

### Cell culture and the TNPs exposure

The human alveolar epithelial cells (A549) were obtained from the National Institute of Occupational Health and Poisons Control, Chinese Center for Diseases Control and Prevention. The use of this cell line was approved by the Ethical Committee of Capital Medical University, Beijing, People's Republic of China. The cells were cultured in RPMI 1640 (HyClone, Logan, UT, USA) supplemented with 10% fetal bovine serum (Gibco, product line of Thermo Fisher Scientific, Waltham, MA, USA), 100 U/mL penicillin, and 100  $\mu\text{g}/\text{mL}$  streptomycin at 37°C in a humidified incubator with 5%  $\text{CO}_2$ . For experiments,  $3 \times 10^5$  A549 cells were seeded in a six-well culture plate for 24 hours. The cells were then washed twice with phosphate buffered saline and treated with TNPs suspended in RPMI 1640 of certain concentrations (6.25  $\mu\text{g}/\text{mL}$ , 12.5  $\mu\text{g}/\text{mL}$ , 25  $\mu\text{g}/\text{mL}$ , 50  $\mu\text{g}/\text{mL}$ , and 100  $\mu\text{g}/\text{mL}$ ) for another 24 hours. The suspension of TNPs was sterilized by an autoclave and dispersed by a sonicator (300 W, 10 minutes) at room temperature to minimize their aggregation. Cells maintained in RPMI 1640 without TNPs were used as the control group.

### Cell viability assay

The cytotoxicity of TNPs was determined using 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-*H*-tetrazolium bromide

(MTT). Briefly,  $1 \times 10^4$  cells were seeded into a 96-well plate in a volume of 100  $\mu$ L RPMI 1640 and incubated for 24 hours at 37°C. Cells were treated with varying concentrations of TNPs (6.25  $\mu$ g/mL, 12.5  $\mu$ g/mL, 25  $\mu$ g/mL, 50  $\mu$ g/mL, and 100  $\mu$ g/mL). After 24 hours incubation, 10  $\mu$ L MTT was added to each well at 5 mg/mL and further incubated for 4 hours. Next, 150  $\mu$ L of dimethylsulfoxide was added in and mixed thoroughly for 10 minutes. Optical density was then measured with a microplate reader (Thermo Multiskan MK3; Thermo Fisher Scientific) at 492 nm according to the instructions.

### Assessment of LDH activity

The lactate dehydrogenase (LDH) leakage assay, which is based on the measurement of LDH activity in the extracellular medium, was determined using a commercial LDH kit (Nanjing Jiancheng Bioengineering Institute, Nanjing, People's Republic of China) according to the manufacturer's protocols. The loss of intracellular LDH and its release into the culture medium is an indicator of irreversible cell death due to cell membrane damage. After cells were treated with different concentrations of TNPs for 24 hours, the supernatants were collected for LDH measurement. A cell medium of 100  $\mu$ L was used for LDH activity analysis and the absorbance at 440 nm was measured by a UV-visible spectrophotometer (SpectraMax M5; Molecular Devices, Sunnyvale, CA, USA).

### Intracellular ROS measurement

Intracellular ROS were measured by flow cytometry using 2,7-dichlorofluorescein diacetate (DCFH-DA) (Nanjing Jiancheng Bioengineering Institute) as a probe. Briefly,  $\sim 3 \times 10^5$  cells were seeded into a six-well plate for 24-hours pretreatment with or without methyltransferase inhibitor, 5-aza (5  $\mu$ M, 10  $\mu$ M, and 20  $\mu$ M), or ROS scavenger  $\alpha$ -LA (0.1  $\mu$ M, 10  $\mu$ M, and 100  $\mu$ M) for 1 hour, and A549 cells were incubated with TNPs (6.25  $\mu$ g/mL, 12.5  $\mu$ g/mL, 25  $\mu$ g/mL, 50  $\mu$ g/mL, and 100  $\mu$ g/mL) for 24 hours at 37°C. Then, cells were washed twice with phosphate buffered saline and co-incubated with serum-free RPMI 1640 containing 10  $\mu$ M DCFH-DA for 30 minutes at 37°C in dark, according to the manufacturer's instructions. The cells were trypsinized and intracellular ROS were measured using flow cytometry (BD LSRFortessa™, BD, Franklin Lakes, NJ, USA) at 488 nm excitation and 525 nm emission.

### DNA isolation and methylation-specific PCR

After A549 cells were pretreated with or without methyltransferase inhibitor, 5-aza (5  $\mu$ M, 10  $\mu$ M, and 20  $\mu$ M), or ROS scavenger  $\alpha$ -LA (0.1  $\mu$ M, 10  $\mu$ M, and 100  $\mu$ M) for 1 hour,

the cells were incubated with TNPs for another 24 hours at 37°C. Genomic DNA was extracted following the protocols of the DNA kit (Solarbio, Beijing, People's Republic of China). One microgram purified DNA was subjected to bisulfate modification, which was performed using CpGenome DNA Modification Kit (Merck Millipore, Billerica, MA, USA) according to the manufacturer's instructions. polymerase chain reaction (PCR) Magic Mix 3.0 (TIANDZ, Beijing, People's Republic of China) was used to detect the PARP-1 methylation under the following conditions: 94°C for 30 seconds; 35 cycles of 94°C for 20 seconds, 60°C for 45 seconds, and 72°C for 1 minute; 72°C for 3 minutes. These methylation-specific PCR (MSP) products were analyzed by 2% agarose gel electrophoresis staining with ethidium bromide and visualized under a UV illuminator. Distinct visible band of the amplicon with methylation-specific primers was considered positive. The density of each band was analyzed by image analysis software (Gel-Pro 4.5) for quantization. The sequences of primers were as follows:

For methylated DNA: MF-PARP-1 (5'-GAGATTAGT TTAGTTAATTGGCGA-3') and MR-PARP-1 (5'-ATCT AAAACTCCTAAACCCAACGAT-3'), a 139 bp fragment (−299 to −161 relative to transcription start site), was chosen to amplify by M primer; and for unmethylated DNA: UF-PARP-1 (5'-GAGATTAGTTTAGTTAATTGGTGA-3') and UR-PARP-1 (5'-ATCTAAACTCCTAAACCCAACA AT-3'), a 139 bp fragment (−299 to −161 relative to transcription start site), was chosen to amplify by U primer.

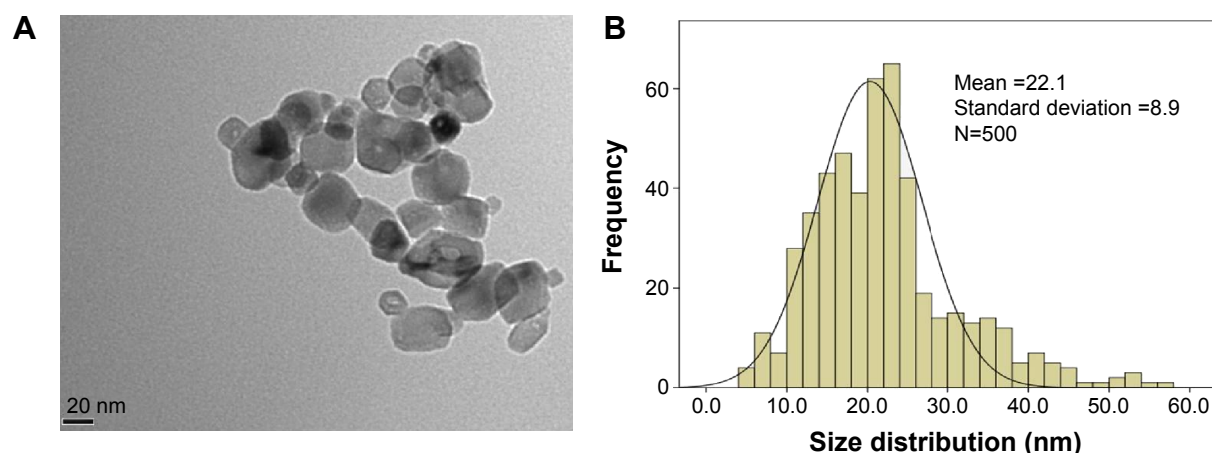
### Statistical analysis

Statistical analysis was performed using SPSS 16.0 software (SPSS Inc., Chicago, IL, USA). Results were expressed as mean  $\pm$  standard deviation of independent experiments. The least significant difference test was used to compare means of two samples, while one-way analysis of variance was used for group means. In all cases,  $P < 0.05$  was considered to be statistically significant.

## Results

### Characterization of TNPs

We first characterized the TNPs by TEM. As shown in Figure 1A, a near-spherical particle shape, with the mean pristine particle diameter of  $22.1 \pm 8.9$  nm (Figure 1B) was obtained by measuring 500 random particles using ImageJ. Hydrodynamic measurements of both ultrasonicated TNPs in ultrapure water and serum-free RPMI 1640 at 1 hour and a concentration of 100  $\mu$ g/mL by dynamic light scattering are shown in Table 1. TNPs in both dispersion mediums exhibited a larger hydrodynamic size than the original



**Figure 1** Characterization of TNPs by TEM.

**Notes:** Particle shape was analyzed by TEM (**A**) and the size distribution in the test media were evaluated by ImageJ software (**B**).

**Abbreviations:** TNPs, titanium dioxide nanoparticles; TEM, transmission electron microscope.

dimension, suggesting that TNPs formed aggregates in mediums due to the van der Waals force and hydrophobic interaction with surrounding media. The polydispersity index values acquired in our study further corroborate the observations made, with stable values being observed for particles dispersed in both ultrapure water and RPMI 1640. Zeta potentials provide quantitative information on the stability of the particles, confirming that the particles are more likely to remain dispersed if the absolute value of zeta potentials is higher than 30 mV.<sup>25</sup> Consequently, zeta potential values in RPMI 1640 and ultrapure water varied between  $-29.8$  mV and  $-11.9$  mV. Nonetheless, the larger aggregates represent the actual scenario that the cells were exposed to.

### Cytotoxicity induced by the TNPs

To evaluate the possible toxicity of TNPs on A549 cells, cell viability was determined after exposing to TNPs ( $0$   $\mu\text{g/mL}$ ,  $6.25$   $\mu\text{g/mL}$ ,  $12.5$   $\mu\text{g/mL}$ ,  $25$   $\mu\text{g/mL}$ ,  $50$   $\mu\text{g/mL}$ , and  $100$   $\mu\text{g/mL}$ ) for 24 hours. As shown in Figure 2, with the dosages increasing, viability of A549 cells induced by TNPs was significantly decreased than control. In addition, the MTT results were strongly in accordance with the increased membrane damage measured by LDH release (Figure 3). Our results suggested that TNPs induced cytotoxicity in a dose-dependent manner.

### Intracellular ROS generation induced by TNPs

To get a closer insight into the possible mechanism of TNP-induced cellular toxicity, the intracellular ROS levels were determined by using the DCFH-DA probe. From Figure 4A and B, we observed the gradually elevated ROS levels in A549 cells in a dose-dependent manner. The generation of ROS in all TNP-treated groups was significantly different with control group. Compared to the group of treatment with TNPs only, the fluorescence intensity of ROS in the group of pretreatment with methyltransferase inhibitor 5-aza was relatively weak (Figure 5A), in line with the effect of ROS scavenger  $\alpha$ -LA pretreatment (Figure 5B). It suggested that the methyltransferase inhibitor markedly decreased the generation of ROS induced by TNPs.

### PARP-1 methylation alteration induced by TNPs

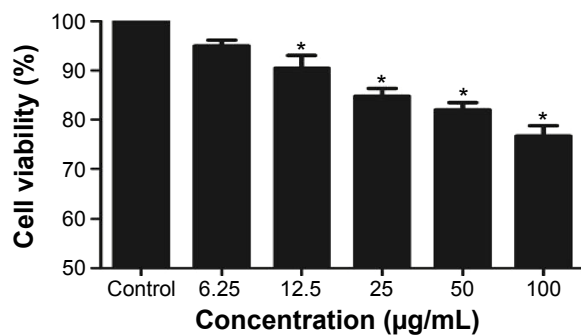
MSP, a common detection method of methylation, was performed to analyze the methylation status of PARP-1. Our results revealed that TNPs notably elevated the level of PARP-1 methylation in a dose-dependent way (Figure 6). However, pretreatment of both methyltransferase inhibitor 5-aza and ROS scavenger  $\alpha$ -LA inversely altered the TNP-induced hypermethylation of PARP-1 promoter

**Table 1** The hydrodynamic size and zeta potential of the TNPs in ultrapure water and RPMI 1640 culture media by DLS analysis

	Ultrapure water			RPMI 1640		
	Hydrodynamic size (nm)	Zeta potential (mV)	PDI	Hydrodynamic size (nm)	Zeta potential (mV)	PDI
1 hour	$276.9 \pm 9.3$	$-29.8 \pm 2.7$	$0.5 \pm 0.4$	$504.8 \pm 108.1$	$-11.9 \pm 1.2$	$0.5 \pm 0.1$

**Abbreviations:** TNPs, titanium dioxide nanoparticles; RPMI, Roswell Park Memorial Institute; DLS, dynamic light scattering; PDI, polydispersity index.

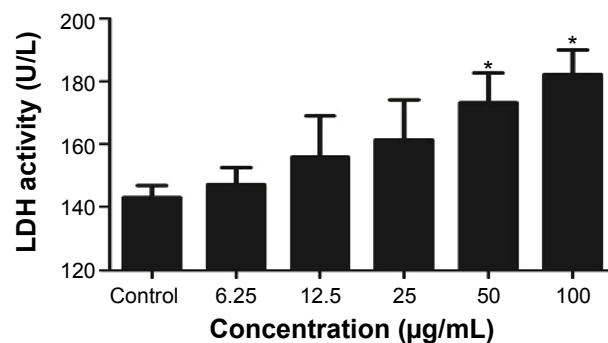




**Figure 2** Cell viability of A549 cells was measured by MTT assay after 24 hours TNP exposure.

**Notes:** Cell viability was significantly decreased in a dose-dependent manner after TNP treatment. \* $P < 0.05$ , compared with control group,  $n = 5$ .

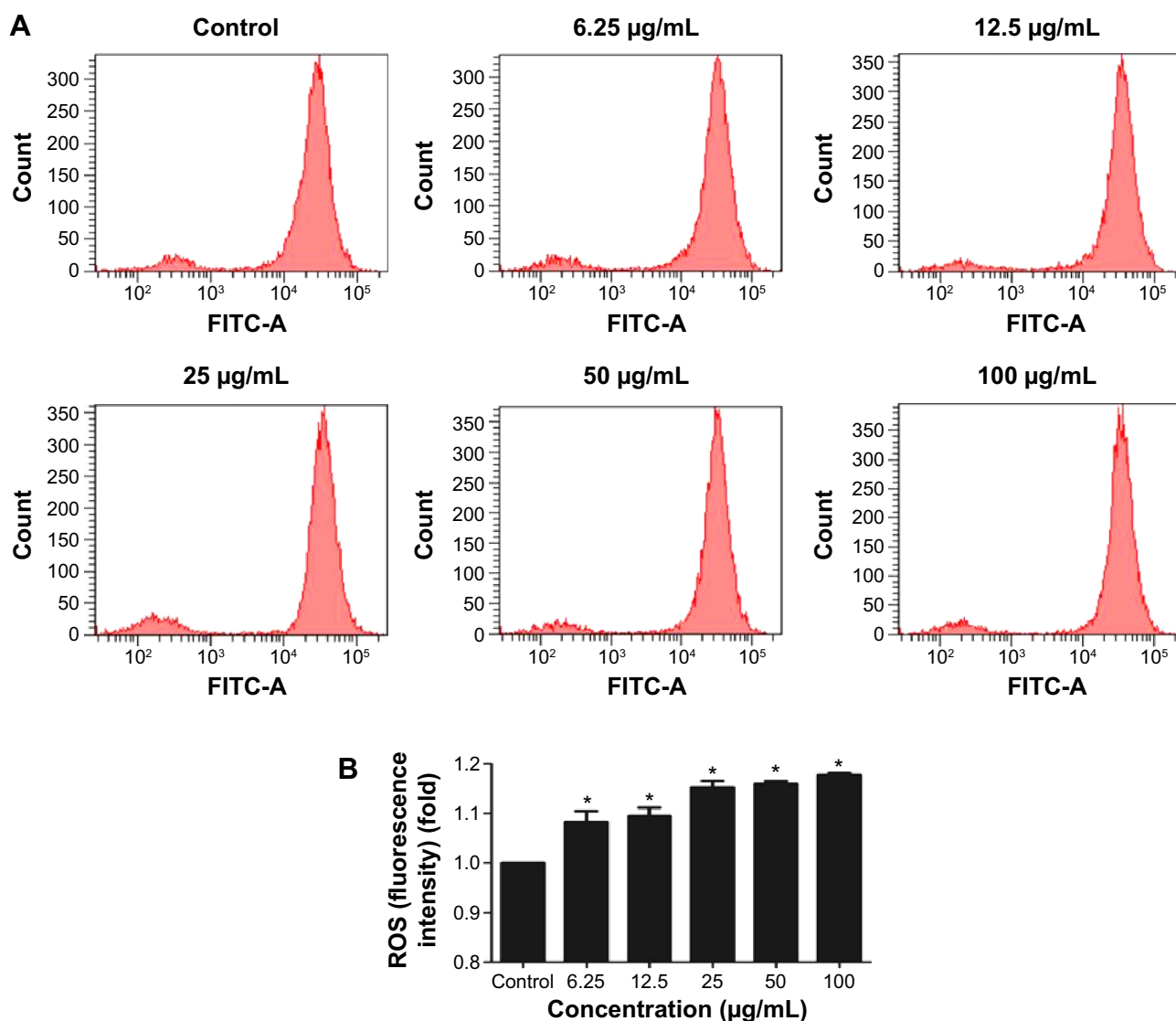
**Abbreviations:** MTT, 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide; TNP, titanium dioxide nanoparticle.



**Figure 3** LDH activity was detected in TNP-treated A549 cells.

**Notes:** After cells were exposed to TNPs (0 µg/mL, 6.25 µg/mL, 12.5 µg/mL, 25 µg/mL, 50 µg/mL, and 100 µg/mL) for 24 hours, the supernatants were determined using a commercial LDH kit according to the manufacturer's protocols. \* $P < 0.05$ , compared with control group,  $n = 5$ .

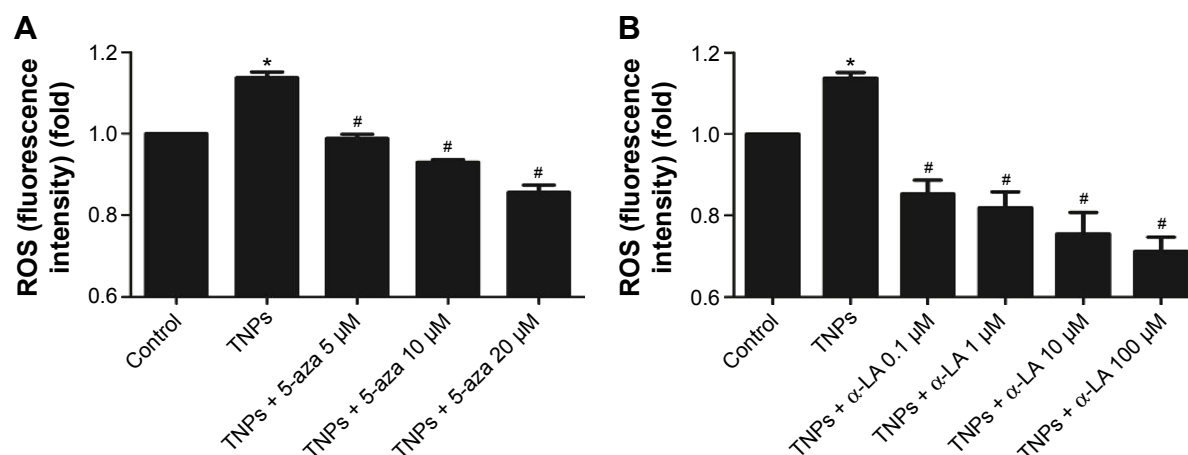
**Abbreviations:** LDH, lactate dehydrogenase; TNPs, titanium dioxide nanoparticles.



**Figure 4** Effects of TNPs on intracellular ROS generation in A549 cells.

**Notes:** The typical picture of ROS detected by flow cytometry (**A**); the intracellular ROS level significantly increased in a dose-dependent manner (**B**). Data are expressed as mean  $\pm$  SD.  $n = 5$ , \* $P < 0.05$ , compared with control.

**Abbreviations:** FITC, fluorescein isothiocyanate; TNPs, titanium dioxide nanoparticles; ROS, reactive oxygen species; SD, standard deviation.



**Figure 5** Effects of inhibitors, 5-aza or  $\alpha$ -LA, on TNP-induced ROS generation in A549 cells.

**Notes:** The methyltransferase inhibitor, 5-aza decreased TNPs-induced ROS generation (A); the antioxidant agent,  $\alpha$ -LA suppressed ROS generation triggered by TNPs (B). Data are expressed as mean  $\pm$  SD; n=5, \* $P$ <0.05, compared with control; # $P$ <0.05, compared with TNPs.

**Abbreviations:** 5-aza, 5-aza-2'-deoxycytidine;  $\alpha$ -LA,  $\alpha$ -lipoic acid; TNPs, titanium dioxide nanoparticles; ROS, reactive oxygen species; SD, standard deviation.

region (Figure 7A and B), implicating that the antioxidant agent  $\alpha$ -LA decreased TNP-induced ROS generation and simultaneously attenuated TNP-triggered PARP-1 hypermethylation.

## Discussion

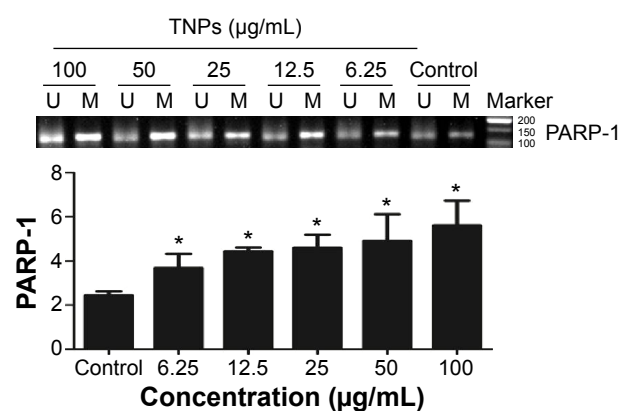
With the increasing TNP-related products, the concern of governments and the public about the health and environmental risks are gradually growing. However, the effects induced by TNPs on human health or environment and the mechanisms of toxicity are still poorly investigated.

TEM provides the direct measurement of particle size, distribution, and morphology by image analysis, while

dynamic light scattering, a common analysis for the size of nanoparticles and determining their aggregation in suspension,<sup>26</sup> often supplies a larger value than Brunauer Emmett Teller or TEM diameter. The Brunauer Emmett Teller size of TNPs claimed by its commercial supplier was <25 nm, which was considered potentially toxic according to previous reports.<sup>27–29</sup> In the TEM image, we obtained a near-spherical particle shape, with the mean pristine particle diameter at  $22.1 \pm 8.9$  nm. Furthermore, the mean hydrodynamic size of TNPs in RPMI 1640 was 504.8 nm, larger than that in ultrapure water (276.9 nm), suggesting that TNPs formed aggregates that were not thoroughly dispersed by ultrasonication. Therefore, the suspension was a mixture of nanoparticles and agglomerations/aggregations.

In our study, we evaluated the cytotoxicity of TNPs (6.25  $\mu$ g/mL, 12.5  $\mu$ g/mL, 25  $\mu$ g/mL, 50  $\mu$ g/mL, and 100  $\mu$ g/mL) by MTT assays and LDH release. Results revealed that TNPs significantly decreased the viability of A549 cells, with the enhancement of LDH activity in a dose-dependent manner. In line with the LDH release, the generation of ROS was dramatically triggered by TNPs, consistent with several published studies.<sup>30,31</sup> The cytotoxicity induced by TNPs has been demonstrated in various cell types such as WISH cells, human bronchial epithelial cell line, BEAS-2B, and mouse fibroblast cells.<sup>10,30,31</sup> Available toxic assays confirmed that TNPs (5 nm) can inhibit A549 cell proliferation, cause DNA damage, and induce apoptosis via a mechanism primarily involving the activation of intrinsic mitochondrial pathway.<sup>32</sup>

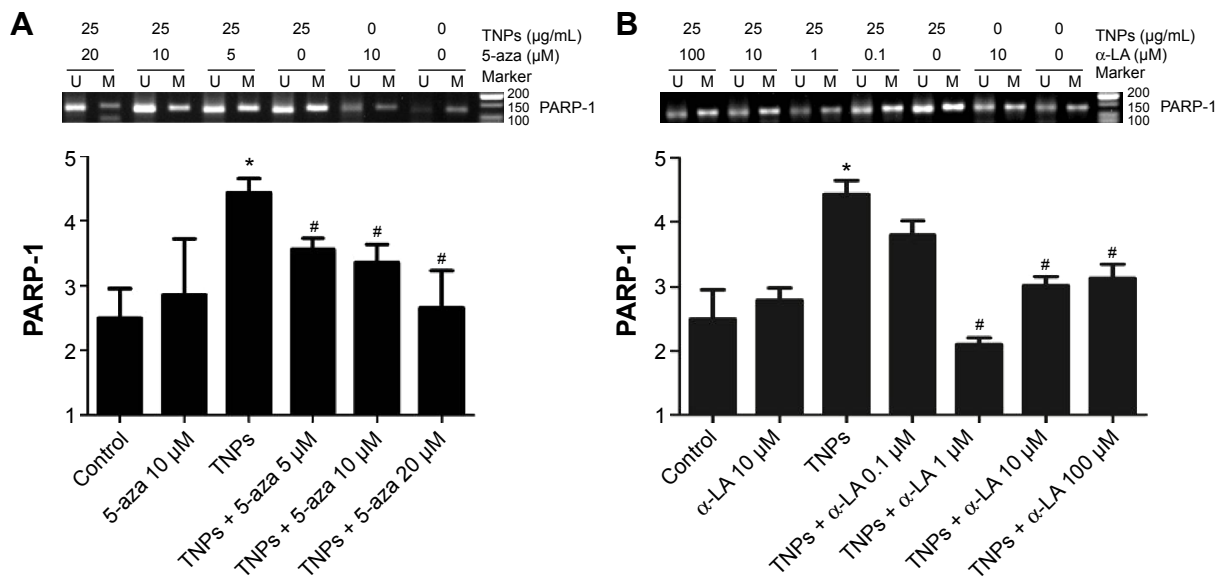
Generally, DNA methylation is the first step in epigenetic phenomena that modulate gene expression via the recruitment of transcription factors. However, the underlying



**Figure 6** TNPs induced the alternation of PARP-1 methylation status.

**Notes:** A549 cells were exposed to various concentrations (6.25  $\mu$ g/mL, 12.5  $\mu$ g/mL, 25  $\mu$ g/mL, 50  $\mu$ g/mL, and 100  $\mu$ g/mL) of TNPs for 24 hours, methylation-specific PCR was performed to examine the methylation status of PARP-1. U and M: primer sets specific to un-methylated (U) and methylated (M) DNA molecules. The data were typical examples of five independent experiments. \* $P$ <0.05, compared with control.

**Abbreviations:** TNPs, titanium dioxide nanoparticles; PARP-1, poly(ADP-ribose) polymerase I; PCR, polymerase chain reaction.



**Figure 7** Effects of inhibitors, 5-aza or  $\alpha$ -LA, on TNP-induced methylation of PARP-1 by MSP in A549 cells.

**Notes:** Treatment with methyltransferase inhibitor, 5-aza (**A**); treatment with antioxidant agent,  $\alpha$ -LA (**B**). U and M: primer sets specific to unmethylated (U) and methylated (M) DNA molecules. The data were typical examples of five independent experiments. \* $P < 0.05$ , compared with control; # $P < 0.05$ , compared with TNPs.

**Abbreviations:** 5-aza, 5-aza-2'-deoxycytidine;  $\alpha$ -LA,  $\alpha$ -lipoic acid; TNPs, titanium dioxide nanoparticles; PARP-1, poly(ADP-ribose) polymerase I; MSP, methylation-specific PCR; PCR, polymerase chain reaction.

mechanisms involved in TNP-induced epigenetic toxicity have not been clearly clarified. Limited data demonstrated that certain nanomaterials such as nano-SiO<sub>2</sub> caused aberrant PARP-1 hypermethylation and inactivity; yet the effects involved in TNP-related PARP-1 abnormal methylation have not been reported.

Our present studies focused for the first time on whether the epigenetic alteration of PARP-1 was responsible for TNP-induced cytotoxicity in A549 cells. Effects of the methyltransferase inhibitor 5-aza and antioxidant agent  $\alpha$ -LA on TNP-induced PARP-1 methylation expression were investigated. The data showed that TNPs triggered aberrant hypermethylation of PARP-1 promoter, and there was a cross talk between oxidative stress and PARP-1 methylation in TNP-induced cellular toxicity.

Numerous studies have defined the crucial role of oxidative stress in the toxicity induced by TNPs.<sup>31–36</sup> TNPs were demonstrated to have the potential to generate ROS and oxidative stress, leading to oxidative DNA damage and micronucleus formation in human epidermal cells (A431) subsequently.<sup>11</sup> The biouptake of TiO<sub>2</sub> in particulate form was the vital cause of ROS generation, which in turn was probably the cause of the DNA aberrations. Zhuang summarized that nano-SiO<sub>2</sub> exposure induced the decrease of genomic DNA methylation and global hypoacetylation in HaCaT cells.<sup>37,38</sup> Still, rare studies on nanomaterial triggered-epigenetic toxicity were published.<sup>39,40</sup>

## Conclusion

In conclusion, to fully understand the effects and interaction mechanisms between ROS and PARP-1 methylation triggered by TNPs, we focused on whether the epigenetic alteration of PARP-1 was responsible for TNP-induced toxicity in A549 cells for the first time. Effects of methyltransferase inhibitor 5-aza and antioxidant agent  $\alpha$ -LA respectively on TNP-induced ROS increasing and PARP-1 hypermethylation in promoter region suggested that there is a cross talk between oxidative stress and PARP-1 methylation in TNP-induced A549 cells toxicity. To our knowledge, this was the first study that showed TNPs can trigger PARP-1 hypermethylation and ROS generation may be involved in this process. These data further strengthened the notion that epigenetic alterations could play a significant role in TNP carcinogenesis.

This finding will not only be helpful to clarify the role of PARP-1 methylation in TNP-induced cytotoxicity but also offer a promising approach in prevention and therapy TNPs cytotoxicity using the methyltransferase inhibitor and antioxidant agent.

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## Disclosure

The authors report no conflicts of interest in this work.

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