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Introduction

In recent decades, titanium dioxide nanoparticles (TiO₂ NPs) have been mass-produced for their worldwide applications in food-related industries, materials for air pollution control, pharmaceuticals and personal care products.¹ The extensive production and use of TiO₂ NPs has increased the level of human exposure through multiple media and pathways. TiO₂ NPs can be delivered directly into the human body as nano-food or nanomedicine.² Due to their small particle size, industrially released TiO₂ NPs can also be inhaled as airborne particulate matter.³ Concerns are increasing about the possible health implications of exposure to TiO₂ NPs.^{4,5}

Increasing number of evidence has shown that TiO₂ NPs exert a variety of adverse health effects including liver function damage, nephrotoxicity and pulmonary toxicity.^{6–8} Both *in vitro*

Titanium dioxide nanoparticles induce sizedependent cytotoxicity and genomic DNA hypomethylation in human respiratory cells

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The widespread use of titanium dioxide nanoparticles (TiO₂ NPs) is gradually increasing the risk of exposure to these potentially hazardous materials. Although numerous health effects of TiO₂ NPs have been investigated, it remains unknown whether they could affect the respiratory cellular epigenome. We explored the viability, membrane integrity, intracellular ROS and genomic DNA methylation of human respiratory cells, as well as their expression of methylation-related genes, after treatment with TiO2 NPs with diameters of 25 nm (nanotube morphology) or 60 nm (anatase morphology). Two cell lines relevant to inhalation exposure, namely human bronchial epithelial cell line (16HBE) and human non-small cell lung cancer cells (A549), were tested, with treatment concentrations ranging from 0.1 to 100 μ g mL⁻¹. The TiO₂ NPs induced time- and concentration-dependent decreases in cell viability in both A549 and 16HBE cells. The reduction in cell viability was greater for the smaller particles (size 25 nm) of the nanotube type. Cellular membrane integrity assays revealed that 16HBE cells were less sensitive to TiO₂ NPs-25 nm (nanotube-type) than were A549 cells, as higher concentrations were required for cytotoxicity against the former. TiO₂ NPs-25 nm (nanotube-type) showed greater toxicity against both cell lines than TiO₂ NPs-60 nm (anatase-type). Intracellular ROS levels in both A549 and 16HBE cell were increased by TiO₂ NPs whereas pretreatment with the antioxidant N-acetyl-L-cysteine eliminated TiO₂ NPs-induced ROS accumulation and reduced cell death. Moreover, the anatase-type TiO₂ NPs resulted in decreased global DNA methylation and altered expression levels of methylation-related genes and proteins, suggesting that these NPs induce cellular epigenomic toxicity. These results allowed us to confirm the epigenetic mechanism by which TiO₂ NPs damage human respiratory cells.

and *in vivo* toxicological assays have characterised the harmful effects of TiO_2 NPs on organs and tissues, especially in the respiratory system. For example, a study on mice indicated that nano- TiO_2 could induce severe pulmonary emphysema, extensive disruption of alveolar septa and type II pneumocyte hyperplasia.⁸ Inhalation of nano- TiO_2 was found to provoke lung inflammation in mice *via* the biological activity of IL-1 α .⁹ Additionally, *in vitro* assays showed that TiO_2 NPs elicited distinct apoptotic pathways in bronchial epithelial cells through lysosomal membrane destabilisation and lipid peroxidation.¹⁰ Oxidative stress was also detected in human pulmonary epithelial cells after exposure to TiO_2 NPs.¹¹

To extend current knowledge, further study is required of the possible epigenetic effects of TiO₂ NPs on respiratory cells. At present, the genomic toxicity of TiO₂ NPs and their ability to affect the cellular epigenome of human respiratory cells remain largely unexplored. It is generally accepted that epigenetic factors regulate the interplay between genes and the environment, and thus affect human diseases.¹² Additionally, epigenetic alterations in airway cells have been found to be associated with respiratory diseases.¹³ Multiple studies have found evidence that DNA methylation plays a role in human

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respiratory diseases.^{14,15} Therefore, TiO₂ NPs may have epigenetic effects on human respiratory cells.

Epigenetic effects are heritable changes, caused by environmental factors, that regulate gene expression via alterations in chromatin proteins without changes in DNA sequences.16 DNA methylation is a major epigenetic modification that can regulate gene expression.17 Aberrant DNA methylation has been demonstrated in a variety of diseases.18 Genomic DNA methylation is catalysed by the activities of methylation enzymes, including DNA methyltransferase 1 (DNMT1), DNMT3a and DNMT3b. Ten-eleven translocation (TET) proteins catalyse the hydroxymethylation step in the DNA demethylation pathway.19 Methyl-CpG-binding domain protein 2 (MBD2), with its ability to bind to methylated DNA, appears to function as a mediator of the biological consequences of the methylation signal.²⁰ Numerous studies have indicated that exposure to nanoparticles can compromise the DNA methylome. For example, exposure of human small-airway epithelial cells to engineered nanoparticles emitted by laser printers was found to result in alterations in both global DNA methylation patterns.²¹ Short-term exposure to engineered nanomaterials was also found to affect the epigenome of macrophages and airway epithelial cells.²² Another study found that multi-walled carbon nanotubes could induce DNA hyper-methylation.23 Taken together, these studies confirm the epigenetic effects of nanoparticles and highlight the importance of DNA methylation in the study of nanoparticle toxicity. In this context, TiO₂ NPs of different diameters and crystal forms may represent a new set of tools to study the influence of nanoscale geometry on cell behaviour.24,25

In this study, the biological responses of human respiratory cells exposed to TiO_2 NPs were evaluated across a wide range of exposure doses. TiO_2 NPs with two different diameters (25 nm and 60 nm) and of nanotube or anatase type were used to investigate the influence of particle size and crystal form. The methylation status of genomic DNA and the expression of *Dnmt3b*, *TETs* and *Mbd2* in the treated cells were also assessed.

Materials and methods

1. Chemicals

Titanium dioxide nanotubes of 25 nm average diameter (TiO₂-N25) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Anatase TiO₂ of 60 nm average diameter (TiO₂-A60) was purchased from Aladdin (Shanghai, China). A stock suspension of TiO₂ NPs at a concentration of 10 mg mL⁻¹ was prepared in cell culture media and ultra-sonicated for 10 min. The TiO₂ NPs were further diluted in cell culture media to 0.1, 1, 10 and 100 μ g mL⁻¹ and sonicated within 5 min before the treatment.

Dimethylsulphoxide (DMSO) and *N*-acetyl-L-cysteine (NAC) were also obtained from Sigma-Aldrich (St. Louis, MO, USA). NAC was used at 3 mmol L^{-1} and added 1 h before TiO₂ NPs-treatment. Trypsin and a penicillin–streptomycin mixture for cell culture were purchased from Hyclone Laboratories, Inc. (Logan, UT, USA). Roswell Park Memorial Institute (RPMI) 1640 medium, minimum essential Eagle's medium (MEM) and foetal bovine serum (FBS) were purchased from Gibco (Carlsbad, CA, USA). A PrimeScript 1st Strand cDNA Synthesis Kit was purchased from Takara Biotechnology (Dalian, China).

2. TiO₂ NP characterisation

The TiO_2 NP stock suspensions were diluted in water and sonicated for 60 s at 60 W (Bioblock ultrasonic processor 75038, Bioblock Scientific, Illkirch, France) for preparation. The size and shape of the TiO_2 NPs were determined using a scanning electron micrographs (SEM, S-4800, Hitachi, Tokyo, Japan). The hydrodynamic diameter and zeta potential of the TiO_2 NPs were determined by dynamic light scattering (Brookhaven 90 Plus, Brookhaven Instruments Co, NY, USA). Measurements were run in triplicate for each sample.

3. Cell culture

The human bronchial epithelial cell line 16HBE was kindly provided by Prof. D. C. Gruenert (University of California, CA, USA). The human non-small cell lung cancer cell line A549 was obtained from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). The 16HBE cells were cultured in fresh MEM supplemented with 10% (v/v) FBS, 100 μ g mL⁻¹ penicillin and 100 μ g mL⁻¹ streptomycin in a cell culture flask in an atmosphere containing 5% CO₂ at 37 °C in an incubator. The A549 cells were cultured in fresh RPMI 1640 medium containing 10% (v/v) FBS, 100 μ g mL⁻¹ penicillin and 100 μ g mL⁻¹ penicillin ang penicillin ang penicillin ang penic

The culture medium was changed every 48 h, and the cells were subcultured every 3 to 4 days at approximately 80% confluence. When the cultured cells had grown to about 80% confluence, the cultures were treated with TiO_2 -N25 or TiO_2 -A60 at different concentrations (0.1, 1, 10 and 100 µg mL⁻¹) for 48 h, and the control cells were cultured with regular culture medium without TiO_2 NPs.

4. Cell viability assay

The cell viability of the 16HBE and A549 cells after treatment with TiO₂ NPs was tested by 3-(4,5-dimethylthiazol-2-yl)-2,5diphenyltetrazolium bromide (MTT) assay. The cells were plated on 96-well plates at a density of 4×10^4 cells per mL and then incubated for 24 h. Different concentrations of TiO2-N25 or TiO₂-A60 (in the range 0.1–100 μ g mL⁻¹) were added into the cell culture and incubation was continued for 24-72 h. Wells containing culture medium but no cells were used as the blank, and wells containing culture medium without TiO₂ NPs treatment were used as the control. After adding 20 µL of MTT (5 mg mL⁻¹) into each well and incubating for 4 h at 37 °C, the MTT medium was discarded and the cells were lysed in 100 μ L of DMSO. The optical density (OD) at 490 nm was measured by a multiwell-plate reader (Bio-Tek EL 808, Bio-Tek Instruments Inc, Colmar, France). Cell viability values (%), expressed as the percentage of absorbance values at each dose compared to the vehicle control, were calculated by (OD_{TiO2 NPs} - OD_{blank})/ $(OD_{control} - OD_{blank}) \times 100\%$.

5. Cellular membrane integrity assay

Cellular membrane integrity was evaluated using the CytoTox-One Homogenous Membrane Integrity Assay (Promega, Madison, WI, USA) following exposure to different concentrations of TiO₂-N25 or TiO₂-A60 (0.1–100 μ g mL⁻¹) for 24, 48 and 72 h. This assay was performed to estimate the number of non-viable cells present after exposure to TiO₂ NPs by measuring the activity of lactate dehydrogenase (LDH) leaked from the cells.

6. Determination of the intracellular reactive oxygen species (ROS) content

The ROS content generated in the 16HBE and A549 cells was measured by the oxidant-sensitive probe DCFH-DA according to the manufacturer's instructions (Beyotime, Shanghai, China). Briefly, the cells were treated with different concentrations of TiO₂ NPs (N25 or A60 in the range 0.1–100 μ g mL⁻¹) or co-treatment with NAC for 48 h and then stained with DCFH-DA 10 μ M at 37 °C for 20 min. The cells were rinsed and imaged by using the fluorescence microscope (Olympus, Tokyo, Japan) at 488 nm for excitation. The fluorescence intensities of DCFH-DA in the cells were analyzed using Image-Pro Plus software (Media Cybernetics, Silver Spring, MD, USA).

7. Analysis of genomic DNA methylation

Genomic DNA from the 16HBE and A549 cells was extracted using a Wizard Genomic DNA Purification Kit (Promega, Madison, WI, USA). Genomic DNA methylation was determined by a Methyl-Flash Methylated DNA Quantification Kit (Epigentek, Brooklyn, NY, USA) according to the manufacturer's instructions.

8. RNA isolation and quantitative real-time PCR (Q-PCR)

Total cellular RNA was isolated from the cells using Trizol reagent (Invitrogen, Carlsbad, CA, USA). Reverse transcription for cDNA synthesis was performed using Revert Aid First Strand cDNA Synthesis Kits (Fermentas, Hanover, MD, USA). Quantifications of gene transcripts were performed by Q-PCR on the 7900HT fast real-time PCR system (Applied Biosystems, Foster City, CA, USA) using SYBR Green PCR Master Mix reagents (Applied Biosystems, Framingham, MA, USA). Samples were analysed in triplicate. The Q-PCR cycle conditions were 95 °C for 2 min, followed by 40 cycles of denaturation at 95 °C for 15 s and annealing and extension at 60 °C for 1 min. The relative gene expression values were calculated by the $2^{-\Delta\Delta C_t}$ method and normalised to values obtained from the housekeeping gene GAPDH. Three independent experiments were performed for each target. Primer sequences for Q-PCR are listed in Table 1.

9. Western blot analysis

The total proteins in the 16HBE and A549 cells were extracted using radio-immunoprecipitation assay (RIPA) lysis buffer (Beyotime, Shanghai, China) containing 1 mM phenylmethanesulfonyl fluoride (Sigma-Aldrich, St. Louis, MO, USA). The concentrations of total protein were determined by BCA protein assay kits (Beyotime, Jiangsu, China). Total proteins from each group were separated by a 12% denaturing polyacrylamide gel. The blotted membranes were blocked using 5% (w/v) non-fat milk in Tris-buffered saline (TBS) and incubated with anti- β -actin, anti-DNMT3B or MBD2 (1 : 1000, Abcam, Cambridge, UK) for 2 h at 37 °C. The membranes were Table 1 Primer sequences for real-time PCR

Gene	Primer sequence (5'-3')	Accession number			
Gapdh	Forward: AACGACCCCTTCATTGAC	NM_001256799.2			
1	Reverse: TCCACGACATACTCAGCAC				
Dnmt3b	Forward: CCGCTTCCTCGCAGCAG	NM_001207055.1			
	Reverse: TGGGCTTTCTGAACGAGTCC				
Mbd2	Forward: GGGAAGAGGATGGATTGCCC	NM_003927.4			
	Reverse: AGCTGACGTGGCTGTTCATT				
Tet1	Forward: CCAAGTCATGCAGCCCTACC	XM_011540204.1			
	Reverse: CACAAGGTTTTTGGTCGCTGG				
Tet2	Forward: CCCGCTGAGTGATGAGAACA	NM_001127208.2			
	Reverse: TGTGCTGCTGAATGTTTGCC				
Tet3	Forward: ACCTGCCAGGCCTTTATGAC	NM_001287491.1			
	Reverse: ACCACACCGTTTCCGTTTCT				

incubated with secondary antibody (LI-COR Bioscience, Lincoln, NE, USA) and the protein bands were visualized using an enhanced chemiluminescence method, and then quantified by Image J.

10. Statistical analysis

Data are expressed as means \pm SD. Statistical analysis was performed using SPSS 13.0 (SPSS Inc., Chicago, IL, USA). Comparisons between all cellular parameters after exposure were performed using one-way analysis of variance and the Tukey correction for multiple-comparison statistical significance. A *P* value <0.05 was considered to be statistically significant. All experiments were independent and conducted in triplicate or more.

Results

1. TiO₂ NP characterization

The TiO_2 NPs (of the nanotube type with particle size of 25 nm, or of the anatase type with particle size of 60 nm) were characterized for their size, shape, hydrodynamic diameter and zeta potential before cell culture experimentation. The SEM images of TiO2 NPs were shown in Fig. 1 which confirmed the size and shape described by Sigma-Aldrich and Aladdin. Physicochemical characteristics (diameter, zeta potentials and hydrodynamic diameters of the suspended particles in water and culture mediums) of the TiO2 NPs are given in Table 2. Both types of TiO2 NPs showed negative zeta potentials in all media. In both MEM and RPMI 1640 culture mediums, the TiO₂-A60 particles showed less negative surface charge than TiO₂-N25. The analysis of the hydrodynamic diameters showed that both types of TiO₂ NPs formed small aggregates in water, RPMI 1640 and MEM suspension. TiO2-N25 presents a population of aggregates with a mean particle size of 125 nm in water, 372 nm in RPMI 1640 and 254 nm in MEM. In water, the TiO₂-A60 particles formed one population of smaller aggregates (51 nm) and another population of larger ones (320 nm). TiO₂-A60 also formed both smaller and larger aggregates in RPMI 1640 and MEM, with size distributions of 86 and 620 nm, and 101 and 620 nm, respectively.



Fig. 1 (A) SEM images of TiO₂-N25. (B) SEM images of TiO₂-A60

Table 2 Physico-chemical characteristics of titanium dioxide nanoparticles

		Average SEM size $(nm \pm SD)$	Zeta potential (mV)		Hydrodynamic diameter (nm)			IV	
	Diameter		Water	RPMI 1640	MEM	Water	RPMI 1640	MEM	structure
TiO ₂ -N25 TiO ₂ -A60	25 nm 60 nm	$\begin{array}{c} 20.9 \pm 1.5 \\ 51.9 \pm 8.7 \end{array}$	-22.3 -13.1	$-12.5 \\ -7.6$	$-10.8 \\ -6.4$	125 51 and 320	372 86 and 572	254 101 and 620	Nanotube Anatase

2. Effects of TiO₂ NP exposure on cell viability

To investigate whether the TiO₂ NPs exerted cytotoxic effects on the A549 and 16HBE cells, the cell viabilities were determined by MTT after 24, 48 and 72 h of treatment. We observed distinct cellular responses to the two types of TiO₂ NP. As shown in Fig. 2A, the cell viabilities of the A549 cells significantly decreased (P < 0.05) after exposure to either TiO₂-N25 or TiO₂-A60 at concentrations of 1–100 μ g mL⁻¹ for 24 h. For the 16HBE cells, TiO₂-N25 inhibited the cell viabilities (compared with the control) at all concentrations in the range 1–100 $\mu g m L^{-1}$ after 24 h treatment (Fig. 2B, P < 0.05). However, 24 h treatment with TiO₂-A60 caused a significant decrease in the cell viabilities of the 16HBE cells only at concentrations of 10 and 100 μ g mL⁻¹. For longer treatment times (48 and 72 h), the decrease in cell viability increased with each increment of the treatment time. The cell viability of both cell lines decreased more strongly with higher treatment concentrations, longer treatment times and smaller particle sizes.

3. Effects of TiO₂ NP exposure on cellular membrane integrity

The cellular membrane integrity of the 16HBE and A549 cells was tested after treatment with TiO₂-N25 or TiO₂-A60 at concentrations of 0, 0.1, 1, 10 and 100 µg mL⁻¹ for 24, 48 and 72 h. Fig. 3A and B illustrates the results of the LDH assay, showing each treatment at various administered doses and times. The A549 cells experienced a significant increase of cell death after exposure to TiO₂-N25 at concentrations of 1–100 µg mL⁻¹ for 24 h (Fig. 3A, P < 0.05). The effects of TiO₂-N25 on the A549 cells were more prominent after 48 h and 72 h than after

24 h (Fig. 3A, P < 0.01). Compared with the control, treatment with TiO₂-A60 significantly increased the cytotoxicity against A549 cells at 0.1–100 µg mL⁻¹ after 48 h and 72 h of exposure (P < 0.01). For the 16HBE cell line, the TiO₂ NPs were cytotoxic only at higher concentrations and longer treatment times, as shown in Fig. 3B. It is evident from Fig. 3B that the 16HBE cells were less sensitive to TiO₂-N25 than the A549 cells were, as higher concentrations were required for cytotoxic effects after 24 h treatment. Nevertheless, the TiO₂-N25 particles were more toxic against both cell lines than TiO₂-A60. In the subsequent experiments, treatments were performed with TiO₂ NPs at 0, 0.1, 1, 10 and 100 µg mL⁻¹ for 48 h.

4. Effects of TiO₂ NP exposure on ROS synthesis

After exposure to TiO₂ NP or co-treatment with NAC, the levels of intracellular ROS in the A549 and 16HBE cells were determined using the fluorescent ROS indicator DCFH-DA. The levels of intracellular ROS in the A549 cells were found markedly increased at 10–100 μ g mL⁻¹ compared with the control (Fig. 4A and B, *P* < 0.01). Quantification data demonstrated that the intensity of DCFH-DA fluorescence (representing the intracellular ROS level) in the 16HBE cells were significantly increased at 1–100 μ g mL⁻¹ compared with the control (Fig. 4A and C). These results indicated that TiO₂ NPs induced ROS accumulation in the A549 and 16HBE cells. Statistical analysis of the DCFH-DA fluorescence levels of cells co-treatment with NAC demonstrated that NAC significantly reduced the TiO₂ NPinduced ROS accumulation (Fig. 4A–C).

Besides, the anti-oxidant NAC was used to test whether TiO_2 NPs increased cell death *via* inducing ROS accumulation. As shown in Fig. 4D and E, co-treatment with NAC abolished TiO_2



Fig. 2 Cell viability was assessed by MTT analysis after the 16HBE and A549 cells were treated with TiO₂ NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$), or culture medium as the control, for 24, 48 or 72 h. (A) Cell viability of A549 cells after treated with TiO₂-N25 and TiO₂-A60. (B) Cell viability of 16HBE cells after treated with TiO₂-N25 and TiO₂-A60. Results are expressed as cell viability relative to control. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls. #, statistically different between each other P < 0.05.



Fig. 3 Cytotoxicity in 16HBE and A549 cells relative to control determined using LDH assay following exposure to TiO₂ NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \,\mu\text{g mL}^{-1}$) for 24, 48 or 72 h. (A) Cytotoxicity in A549 cells after treated with TiO₂-N25 and TiO₂-A60. (B) Cytotoxicity in 16HBE cells after treated with TiO₂-N25 and TiO₂-A60. Results are expressed as cytotoxicity relative to control. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls. #, statistically different between each other P < 0.05.



Fig. 4 ROS accumulation in 16HBE and A549 cells relative to control after exposure to TiO₂ NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$) or co-treatment with NAC for 48 h. (A) ROS accumulations in A549 and 16HBE cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC were photographed by a fluorescence microscope at 488 nm. (B) Relative ROS levels in A549 cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC were photographed by a fluorescence microscope at 488 nm. (B) Relative ROS levels in A549 cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC for 48 h. (C) Relative ROS levels in 16HBE cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC for 48 h. Results from (B) and (C) are expressed as ROS level relative to control. (D) Cell viability of A549 cells after exposure to TiO₂ NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$) or co-treatment with NAC for 48 h. (E) Cell viability of 16HBE cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC for 48 h. (E) Cell viability of 16HBE cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC for 48 h. (E) Cell viability of 16HBE cells after treated with TiO₂ NPs (N25 or A60) or co-treatment with NAC for 48 h. (E) Cell viability relative to control. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls. # or Δ , statistically different between each other P < 0.05, $\Delta \Delta$, P < 0.01 compared with controls.

NP-induced cell death as determined by MTT which was consistent with the effect of NAC of reducing the levels of TiO_2 NPs-induced ROS. Taken together, this study suggested that TiO_2 NPs induced cell death mainly *via* a ROS-dependent mechanism.

5. Effects of TiO_2 NP exposure on genomic DNA methylation

The degree of genomic DNA methylation of the A549 and 16HBE cells was determined after treatment with different concentrations of TiO_2 -N25 or TiO_2 -A60 for 48 h. As shown in Fig. 5A, the genomic DNA methylation levels of the A549 cells were reduced

after exposure to TiO₂-N25 at 0.1–100 μ g mL⁻¹ compared with the control (P < 0.01). After exposure to TiO₂-A60, at all concentrations in the range 0.1–100 μ g mL⁻¹, the extent of DNA methylation in the A549 cells decreased significantly (hypomethylation) compared with the control (Fig. 5A, P < 0.05). The obtained results thus indicated that TiO₂ NPs exposure induced a significant decrease of genomic DNA methylation in A549 cells. Fig. 5B shows that for the 16HBE cells, the global DNA methylation levels decreased after exposure to TiO₂-N25 at 100 μ g mL⁻¹, compared with the control (P < 0.01). Moreover, for treatment with TiO₂-A60, the genomic DNA methylation levels



Fig. 5 Genomic DNA methylation levels of A549 and 16HBE cells after treatment with different concentrations of TiO₂ NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$) for 48 h. (A) Genomic DNA methylation levels in A549 after treated with TiO₂-N25 and TiO₂-A60. (B) Genomic DNA methylation levels in 16HBE cells after treated with TiO₂-N25 and TiO₂-A60. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls. #, statistically different between each other P < 0.05.

of the 16HBE cells were significantly decreased by all concentrations in the range 0.1–100 μ g mL⁻¹ (Fig. 5B, *P* < 0.05).

6. Effects of TiO_2 NP exposure on expression of *Dnmt3b*, *Mbd2* and *TETs* in A549 and 16HBE cells

To investigate the possible mechanisms of the genomic DNA hypomethylation observed after exposure to TiO_2 NPs, we further examined the expression levels of *Dnmt3b*, *Mbd2* and *TETs* in the A549 and 16HBE cells.

As shown in Fig. 6A, C and J, the mRNA expression levels of *Dnmt3b*, *Mbd2* and *Tet3* decreased in the A549 cells after exposure to TiO₂-N25 at 0.1–100 µg mL⁻¹ (P < 0.01). The mRNA expression levels of *Tet1* and *Tet2* decreased in the A549 cells after exposure to TiO₂-N25 at 1–100 µg mL⁻¹ (Fig. 6E and G, P < 0.01). TiO₂-A60 reduced the mRNA expression levels of *Dnmt3b*, *Mbd2*, *Tet2* and *Tet3* in the A549 cells at 0.1–100 µg mL⁻¹ (Fig. 6B, D, H and K, P < 0.01). The mRNA expression levels of *Tet1* decreased in the A549 cells after exposure to TiO₂-A60 at 1–100 µg mL⁻¹ (Fig. 6F, P < 0.01).

Interestingly, in the 16HBE cells, exposure to $\text{TiO}_2\text{-N25}$ or $\text{TiO}_2\text{-A60}$ (0.1–100 µg mL⁻¹) had different (mostly opposite) effects on the expression of *Dnmt3b*, *Mbd2*, *Tet1*, *Tet2* and *Tet3* compared with the A549 cells (Fig. 7A–K). Exposure to $\text{TiO}_2\text{-N25}$ caused an increase in the expression of *Dnmt3b*, *Tet1*, *Tet2* and *Tet3* in the 16HBE cells relative to the untreated cells (Fig. 7A, E, G and J). Only the expression of *Mbd2* was reduced in the 16HBE cells after exposure to $\text{TiO}_2\text{-N25}$, and only at relatively high concentrations (10–100 µg mL⁻¹), as shown in Fig. 7C. Likewise, in the TiO₂-A60 exposure groups, the expression of *Mbd2*, *Tet1*, *Tet2* and *K*).

Based on these results, we hypothesised that the changes in expression levels of Dnmt3b, Mbd2 and TETs may be associated with TiO₂ NPs-induced genomic hypomethylation in the A549 and 16HBE cells.

7. Effects of TiO₂ NP exposure on the expression levels of DNMT3B and MBD2 proteins

As shown in Fig. 8A and C, treatment of TiO_2 NPs significantly reduced the levels of DNMT3B in A549 cells in a dose-dependent manner when compared with control (P < 0.05). The expression levels of MBD2 proteins were significantly decreased after TiO₂-N25-treatment at concentrations of 0.1, 1, and 100 µg mL⁻¹ and at concentrations of 10–100 µg mL⁻¹ in the TiO₂-A60 group (P < 0.05, Fig. 8A and D).

In the 16HBE cells, exposure to TiO₂-N25 caused an increase in the expression of DNMT3B at relatively high concentrations (10– 100 µg mL⁻¹) which was opposite to the A549 cells (P < 0.05, Fig. 8B and E). And in the TiO₂-A60 groups, the MBD2 proteins were decreased at concentrations of 10–100 µg mL⁻¹ (P < 0.01, Fig. 8B and E). However, the expression of MBD2 proteins were decreased after exposure to TiO₂-N25 only at the highest concentration (100 µg mL⁻¹) and were increased in the TiO₂-A60 groups at concentrations of 1–100 µg mL⁻¹, as shown in Fig. 8B and F.

Discussion

This study evaluated the potential toxicity of varying doses of TiO_2 NPs on human respiratory cells. The alveolar epithelium and bronchial epithelial cells make direct contact with inhaled nanoparticles, so we chose two cell lines as *in vitro* models of this contact.^{11,25} The viability, membrane integrity, intracellular ROS and genomic DNA methylation of the cell lines, as well as their expression of methylation-related genes and proteins, were measured after exposure to two types of TiO_2 NPs: those with particle sizes of 25 nm (nanotube crystalline phase) or 60 nm (anatase crystalline phase).

Both types of TiO_2 NPs were characterised physicochemically before the exposure experiments. In MEM and RPMI 1640, the surface charge on the TiO_2 NPs was less negative than in water. The decrease in charge was caused by a partial compensation of the negative charges by inorganic cations attracted to the surfaces of the TiO_2 NPs in the cell culture media.²⁶ The TiO_2 -N25 particles retained more negative surface charge than TiO_2 -A60. The relatively low zeta potentials of the particles reduced the stability of their nanodispersions, causing them to aggregate. The particles had greater hydrodynamic diameters in cell culture media than in water due to this enhanced aggregation. However, it has been reported that even



Fig. 6 Effects of exposure to TiO_2 NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$) for 48 h on mRNA expression of *Dnmt3b*, *Mbd2* and *Tet1*-3 after exposure to TiO_2 -N25 (A, C, E, G and J) and TiO_2 -A60 (B, D, F, H and K). Results are expressed as cytotoxicity relative to control. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls.



Fig. 7 Effects of exposure to TiO_2 NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$) for 48 h on mRNA expression of *Dnmt3b*, *Mbd2* and *Tet1*-3 after exposure to TiO_2 -N25 (A, C, E, G and J) and TiO_2 -A60 (B, D, F, H and K). Results are expressed as cytotoxicity relative to control. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls.

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Fig. 8 Effects of exposure to TiO₂ NPs (particle sizes of 25 nm and 60 nm, concentrations of $0.1-100 \ \mu g \ mL^{-1}$) for 48 h on expression of DNMT3B and MBD2 proteins in A549 and 16HBE cells. (A), (C) and (D) Relative expression level of DNMT3B and MBD2 in A549 cells. B, E and F Relative expression level of DNMT3B and MBD2 in 16HBE cells. Results are expressed as cytotoxicity relative to control. Data from three independent experiments are expressed as means \pm SD (n = 3). Error bars indicate standard deviation of three samples. *, P < 0.05 and **, P < 0.01 compared with controls.

in the form of aggregates, the cytotoxic effects of nanoparticles depend on the primary particle size and surface area.²⁷

The viability and membrane integrity of both cell lines were negatively affected by treatment with TiO₂-N25 or TiO₂-A60. For both cell lines, the decrease in cell viability after exposure to TiO₂ NPs was stronger for higher treatment concentrations, longer treatment times and smaller particle sizes. A previous study demonstrated that TiO₂ NPs with diameters of 5 nm inhibited the proliferation of A549 cells in a dose- and timedependent manner.²⁸ In addition, the cytotoxicity results in this study suggested that A549 cells were more sensitive to exposure to TiO₂-N25 than were 16HBE cells. This agrees with an earlier study showing that although TiO₂ NPs with diameters of 21 nm were cytotoxic for both A549 and 16HBE cell lines at relatively high concentrations, A549 cells were more sensitive than 16HBE cells.¹¹ The influence of size on the toxicity of TiO_2 NPs has been further reported elsewhere.^{29,30} The cytotoxic effects of TiO_2 NPs exposure on A549 and 16HBE cells evidently impair their normal functioning. More detailed mechanistic studies are needed to clarify the cellular responses to TiO_2 NPs exposure.

 ${\rm TiO}_2$ NPs possesses toxicity has been proofed, but the underlying mechanisms of which remain elusive. Disturbances in oxidative stress state of cells can cause toxic effects that damage all components of the cell, including proteins, lipids, and DNA. Oxidative stress causes base damage, which is mostly caused by ROS generated. Increased intracellular ROS accumulation has been shown to induce cell death. As demonstrated in a previous study, TiO₂ (15 nm) NPs induces increased intracellular ROS in bronchial epithelial cells (16HBE14o-cell line and primary cells).¹⁰ In this study, the levels of intracellular ROS in the A549 and 16HBE cells were increased after exposure to TiO₂ NP. Furthermore, the anti-oxidant NAC completely abolished the TiO₂ NPs-induced ROS accumulation and cell death in both A549 and 16HBE cells. This study supports the concept that ROS accumulation plays a role the cytotoxicity of TiO₂ NPs.

Although the potential adverse effects of TiO₂ NPs have become more apparent in recent years,⁴ the epigenetic mechanisms responsible for the toxic effects of TiO₂ NPs remain largely unknown. This work represents the first genome-wide study of DNA methylation in human respiratory cells treated with TiO2 NPs. We found that both TiO2-N25 and TiO2-A60 significantly reduced global DNA methylation levels in A549 and 16HBE cells. Moreover, the A549 cell line proved more sensitive to genomic DNA methylation by TiO₂ NPs than did 16HBE cells. For the latter cell line, the epigenetic effects of TiO₂-A60 were more significant than those of TiO₂-N25. This observation is in line with previous studies reporting that anatase TiO₂ NPs exhibited more pronounced genotoxicity and deleterious effects, due to the photocatalytic properties of anatase TiO2.25,30 Recently, exposure to various sources of particulate matter has been reported to affect DNA methylation.31-33 Consistent with our results, an in vitro study indicated that TiO2 particles of diameter 21 nm modestly affected DNA methylation in human small-airway epithelial cells (SAECs).22 Preliminary evidence has also been found that exposure to TiO₂ nanotubes can regulate methylation levels in gene promoters.34 Global hypomethylation occurs early in tumorigenesis and predisposes cells to genomic instability and further genetic changes.35 DNA hypomethylation is associated with opening of the chromatin configuration and transcriptional activation, leading to chromosomal instability and aberrant expression of genes.³⁶ Additionally, changes in global DNA methylation patterns have been shown to play a role in airway diseases.^{14,15} Thus, we conclude that these alterations in global DNA methylation were associated with the cytotoxic effects of TiO₂ NPs and might further contribute to the higher risk of respiratory diseases. In addition, the variation in the effects of different TiO2 NPs on different cell lines suggests that both the cell line sensibility and the nanoscale particle geometry affect the outcome.

As previously reported, analysis of global DNA methylation may mask the redistribution of hypomethylation and hypermethylation, which may result in cumulatively unchanged levels of global DNA methylation.³³ Therefore, to further investigate the mechanism of global hypomethylation induced by TiO₂ NPs, the expression levels of *Dnmt3b*, *Mbd2* and *TETs* were determined in this study. In addition to changes in genomic DNA methylation, changes in the expression of DNA methylation-related genes were evident in both cell lines posttreatment with TiO₂ NPs. In the A549 cells, the expression levels of *Dnmt3b*, *Mbd2* and *TETs* decreased after exposure to TiO₂ NPs. However, in the 16HBE cell line, increased expression levels of *Dnmt3b* and *TETs* were observed in the TiO₂-N25 groups, and of *Mbd2* and *TETs* in the TiO₂-A60 groups. Evidence indicates that DNMT3B can methylate hemimethylated and functions.37,38 MBD2, with its ability to bind to methylated DNA, can suppress transcription from methylated target gene promoters. MBD2 is also reported to function as a demethylase to activate transcription.³⁹ 5-Methylcytosine (5-mC) can be converted to 5-hydroxymethylcytosine (5-hmC) by the TET proteins during DNA demethylation.19 Down-regulation of methyltransferases and Mbd2 at mRNA levels was previously reported after exposure to SiO₂ nanoparticles, which were also associated with global hypomethylation.40 The expression levels of DNA methyltransferases and Tet1 were reduced in alveolar macrophages after treatment with laser printer-emitted engineered nanoparticles.41 Moreover, significant losses in the expression of Tet1 were observed following CuO exposure in alveolar macrophages and SAEC cells.^{22,41} Consequently, the observed loss of Dnmt3b and Mbd2 expression may be the cause of the global hypomethylation in this study, implying that hypomethylation was the result of inhibitory effects on *de novo* DNA methylation by TiO₂ NPs. Indeed, there is evidence that increased expression of Dnmt3b and Mbd2 acts as regulatory feedback in response to genomic hypomethylation.42,43 The alterations in TETs expression suggest the diminished ability of TETs to convert 5-mC into 5-hmC during demethylation. As shown by the western blot analyses, the expression levels of these proteins were consistent with the expression level of their mRNAs.

unmethylated CpG sites and also possesses maintenance

In summary, exposure to TiO₂ NPs has the potential to trigger an unfavourable biological response in two cell lines relevant to the respiratory system. The negative effects of TiO2 NPs include a decrease in cell viability, a rise in cell death, genomic hypomethylation and altered expression of methylation-related genes, and may lead to increased risk of respiratory diseases in individuals exposed to TiO₂ NPs. The different levels of cytotoxicity suggest a cell-specific sensitivity in addition to the influence of nanoparticle characteristics. While TiO₂ NPs are weak mutagens,⁴⁴ they may have detrimental effects on the expression of methylation-related genes, leading to alterations in the extent of DNA methylation. Importantly, we have shown that genomic hypomethylation and altered expression of DNA methylation-related genes may occur after exposure to low doses $(0.1 \ \mu g \ mL^{-1})$ of TiO₂ NPs. Thus, our results indicated that both types of TiO₂ NPs inhibited cell viability in a dose- and time-dependent manner. Moreover, the ability of TiO₂ NPs to affect the cellular epigenome was demonstrated. The subsequent consequences of these altered epigenetic states remain unclear. Further investigations are needed to clarify the epigenomic effects of TiO2 NPs in cells and more importantly in vivo. In addition, a broader range of diameters for each particle type of TiO2 NPs remains to be investigated in future research work. This may enable us to identify key factors to predict the toxicity of nanomaterials.

Conclusion

Our data indicate that TiO_2 NPs can elicit unfavourable biological responses *in vitro*. Exposure to TiO_2 NPs led to significant changes in cell viability, cell death, intracellular ROS, cell genomic DNA methylation and methylation-related gene and protein expression. Moreover, the observed dysfunction of DNA methylation suggests that TiO_2 NPs exert effects on the cellular epigenome. To further investigate the mechanism of toxicity in detail, a study on murine responses to TiO_2 NP exposure *via* intratracheal instillation and whole-body inhalation is currently in progress.

Author contributions

Y. M. and Y. G. performed the experiments involving cell culture, cell viability, cell membrane integrity and intracellular ROS. Y. G. analysed the DNA methylation, gene and protein expressions in the cultured cells. Y. M., Y. G., S. W., Z. L. and Z. Q. carried out the analyses of data and the statistical tests. Y. M. designed the experiments and Y. G. drafted the manuscript. All authors approved the final version of the manuscript to be published.

Conflict of interest

None of the authors has any potential conflict of interest or financial interests to disclose.

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