

Effects of titanium nanoparticles on adhesion, migration, proliferation, and differentiation of mesenchymal stem cells

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Background: The purpose of this study was to investigate the influences of nanoscale wear particles derived from titanium/titanium alloy-based implants on integration of bone. Here we report the potential impact of titanium oxide (TiO₂) nanoparticles on adhesion, migration, proliferation, and differentiation of mesenchymal stem cells (MSC) from the cellular level to the molecular level in the Wistar rat.

Methods: A series of TiO₂ nanoparticles (14 nm, 108 nm, and 196 nm) were synthesized and characterized by scanning electron microscopy and transmission electron microscopy, respectively.

Results: The TiO₂ nanoparticles had negative effects on cell viability, proliferation, and the cell cycle of MSC in a dose-dependent and size-dependent manner. Confocal laser scanning microscopy was used to investigate the effects of particle internalization on adhesion, spreading, and morphology of MSC. The integrity of the cell membrane, cytoskeleton, and vinculin of MSC were negatively influenced by large TiO₂ nanoparticles.

Conclusion: The Transwell migration assay and a wound healing model suggested that TiO₂ nanoparticles had a strong adverse impact on cell migration as particle size increased ($P < 0.01$). Furthermore, alkaline phosphatase, gene expression of osteocalcin (OC) and osteopontin (OPN), and mineralization measurements indicate that the size of the TiO₂ nanoparticles negatively affected osteogenic differentiation of MSC.

Keywords: mesenchymal stem cells, titanium dioxide, nanoparticles, cytotoxicity, adhesion, migration

Introduction

Titanium and its alloys are widely used as bone implants in the clinical setting, mainly because of their good biocompatibility and mechanical properties.^{1,2} Nevertheless, owing to their inherent surface inertness, titanium-based implants can only integrate with the surrounding bone tissue via mechanical interlocking, and the ability to do this is closely related to surface topography at the microscale or nanoscale.^{3,4} Titanium substrates with a nanorough surface have been shown to influence cell adhesion, proliferation, and differentiation of osteoblasts.⁵ Guida et al confirmed that microscale or nanoscale modification of titanium substrates enhanced proliferation and synthesis of the extracellular matrix in human bone marrow-derived mesenchymal stem cells (MSC) and expression of osteoprotegerin.⁶ Annunziata et al also demonstrated that nanostructured titanium surfaces enhance adhesion and differentiation of human bone marrow-derived MSC.⁷ Thus, fabrication of titanium substrates with desirable topography has been a focus in the development of high quality titanium implants in

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recent decades. With the rapid development of nanotechnology, surface engineering of titanium using nanotopography to regulate cell behavior is now attracting much research attention.^{8,9} In our previous research, we confirmed that hierarchically microstructured or nanostructured surfaces¹⁰ and nanostructured titanium substrates¹¹ could efficiently regulate the proliferation and differentiation of MSC.

However, in long-term clinical application (eg, titanium-based hip joint implants), microscale and/or nanoscale wear particles can be generated at the implant/bone interface due to corrosion, aseptic loosening, and prosthesis friction.^{12,13} On the other hand, composite bone prostheses containing titanium nanoparticles may be released directly at the implant/bone interface.¹⁴ Previous studies have confirmed that microscale wear debris in the form of titanium particles released from a titanium implant occur *in vivo*, resulting in adverse effects, ie, significant attenuation of osteoblast function, increased osteoclast recruitment, and decreased bone integration.^{15,16}

Given the worldwide concerns about nanotoxicity in biomaterials, the potential cytotoxicity of titanium nanoparticles has also been investigated in different cell lines. Previous studies have demonstrated that titanium nanoparticles have negative effects on cells, including inflammatory reactions in endothelial cells,¹⁷ damage to the DNA of human lymphoblastoid cells,¹⁸ and induction of micronuclei and apoptosis in Syrian hamster embryo fibroblasts.¹⁹ Other reports have shown that titanium dioxide (TiO₂) nanoparticles delivered to the abdominal cavity and liver in mice cause brain injury and liver damage, respectively.^{20,21} Wang et al also confirmed that TiO₂ nanoparticles are potentially toxic to major organs and cause damage to the knee joints in rabbits due to their ability to catalyze oxidative DNA damage.²²

To be able to confirm further the feasibility of development of surface nanostructured titanium-based bone implants, there is an urgent need for increased knowledge about how titanium nanoparticles affect biological function in cells involved in bone formation. In a previous study, we established a correlation between the cytotoxicity of TiO₂ nanoparticles and different particle sizes on a sub-200 nm scale in respect to osteoblasts.²³ In this study, we investigated the interaction between TiO₂ nanoparticles and MSC. Although a previous study investigated the interaction between nanoparticles and neural stem cells,²⁴ the effects of titanium nanoparticles on the biological behavior of bone marrow-derived MSC have not yet been reported. The rationale for choosing bone marrow-derived MSC was that the head of a titanium-based hip joint must be inserted into

the medullary cavity of a long bone for fixation in clinical application, and the medullary cavity contains plentiful MSC. At this site, titanium implants can interact with MSC directly. In our previous study,²³ we investigated the influence of different-sized titanium nanoparticles on the biological behavior of osteoblasts.

The objective of this study was to use titanium nanoparticles of different sizes mimicking the wear debris potentially loosening from a joint implant and to investigate how titanium nanoparticles influence the biological behavior of MSC, including cell adhesion, migration, proliferation, and differentiation.

Materials and methods

Materials

Reagents used in this study, including fluorescein isothiocyanate (FITC), alkaline phosphatase (ALP), a bicinchoninic acid assay kit, Hoechst 33258, dexamethasone, L-ascorbate-2-phosphate, 1,25-dihydroxy vitamin D3, b-glycerol phosphate, and alizarin red-S, were purchased from Sigma-Aldrich Chemical Co (St Louis, MO, USA). Rhodamine-phalloidin was purchased from Invitrogen Co (Carlsbad, CA, USA). A CCK-8 assay kit was provided by Dojindo Laboratories (Kumamoto, Japan). Vinculin antibody was purchased from Santa Cruz Biotechnology Co (Santa Cruz, CA, USA). DyLight 649 goat anti-mouse secondary antibody was supplied by ZSGB-Bio Co (Beijing, People's Republic of China). Other reagents were sourced from Oriental Chemical Co (Chongqing, People's Republic of China).

Synthesis and characterization of TiO₂ nanoparticles

A series of TiO₂ nanoparticles was synthesized with a hydrothermal method.²³ The morphological features of the TiO₂ nanoparticles were characterized by field emission scanning electron microscopy (FEI Nova 400 Nano SEM, Philips, Eindhoven, The Netherlands) and transmission electron microscopy (TECNAI-10, Phillips), respectively.

Cell culture

MSC were isolated from the bone marrow of the femur and tibia of a Wistar rat (weight 150–200 g) using a technique reported previously.²⁵ The MSC were cultured in a 75 cm² flask containing 5 mL of Dulbecco's Modified Eagle's Medium supplemented with 10% bovine serum (Gibco, Grand Island, NY, USA) at 37°C in a 5% CO₂ atmosphere. The cell culture medium was changed on the first day and thereafter every 2 days. After reaching confluence, the cells were detached with 0.25% trypsin in 1 mM

tetrasodium EDTA, centrifuged, and resuspended in complete medium for reseeding in new culture flasks. The complete medium contained osteogenic supplements, ie, L-ascorbate-2-phosphate 50 mg/mL, dexamethasone 0.1 mM, β -glycerol phosphate 10 mM, and 1,25-dihydroxy vitamin D3 10 nM. Cells at the third passage were used for our experiments.

Cell viability assay

The viability of MSC was examined using the CCK-8 assay kit. Briefly, the cells were cultured in a 24-well plate with or without (control) TiO₂ particles at an initial seeding density of 2×10^4 cells/cm². For investigation of dose-dependent effects, MSC were incubated with different concentrations of TiO₂ nanoparticles (0.05, 0.10, and 0.20 mg/mL) for 48 hours. For investigation of time-dependent effects, MSC were incubated with TiO₂ 0.10 mg/mL nanoparticles for 3, 7, and 14 days. CCK-8 kit reagents were added to each well according to the manufacturer's instructions. Absorbance was measured using a microplate reader (model 550, Bio-Rad Laboratories, Hercules, CA, USA) at a wavelength of 450 nm ($n = 6$).

Cell cycle assay using flow cytometry

The cells were cultured in a 24-well plate at an initial cell seeding density of 1×10^5 cells/cm². TiO₂ nanoparticles were added into each well when cell confluence reached around 70%, and the final concentration of nanoparticles was adjusted to 0.10 mg/mL. After culture for one, two, and three days, the cells were washed twice with phosphate-buffered solution and centrifuged. The cells were then fixed with 70% ethanol at 4°C for one hour and washed with phosphate-buffered solution. Next, the treated cells were resuspended with propidium iodide solution 0.05 mg/mL containing RNase and incubated in the dark at room temperature for 30 minutes. The DNA content of the cells was then analyzed using a flow cytometer (EPICS XL-MCL, Beckman Coulter, Fullerton, CA, USA).

Particle internalization assay

FITC-TiO₂ nanoparticles were constructed in a manner reported elsewhere²⁶ and their translocation was investigated. Briefly, 15 mg of TiO₂ nanoparticles were suspended in 1 mL of distilled water and treated with ultrasonication at room temperature for 45 minutes. The sample was then centrifuged to remove the aqueous solution. Next, 0.4 mL of dopamine 1 mM solution was added with stirring for 10 minutes. After removal of the free dopamine by centrifugation, the resulting dopamine-modified TiO₂ nanoparticles were further reacted with FITC to yield FITC-labeled TiO₂ nanoparticles.

Free FITC molecules were removed by centrifugation. FITC-labeled TiO₂ nanoparticles were resuspended in phosphate-buffered solution for further experiments. Cells were cocultured with or without (control) FITC-TiO₂ nanoparticles for 24 hours. Internalization of the TiO₂ nanoparticles was investigated by confocal laser scanning microscopy (TCS SP5, Leica Microsystems, Wetzlar, Germany).

Cell adhesion

MSC were cultured with 0.10 mg/mL of FITC-TiO₂ nanoparticles for 12 and 24 hours, respectively. The cells were washed with phosphate-buffered solution three times and fixed with 4% paraformaldehyde at 4°C for 30 minutes. The treated cells were then permeabilized with 0.25% Triton X-100 for 10 minutes and washed with phosphate-buffered solution three times. Next, the cells were incubated with 5% bovine serum albumin/phosphate-buffered solution at 37°C for one hour. Subsequently, mouse monoclonal antibody against vinculin (1:200) was added and incubated at 4°C overnight. Next, DyLight 649 goat anti-mouse secondary antibody (1:100) was added and incubated at 37°C for a further hour. Subsequently, the cells were stained with 5 U/mL of rhodamine-phalloidin at 4°C overnight and counterstained with 10 μ g/mL of Hoechst 33258 at room temperature for 5 minutes. Finally, the cells were mounted with 95% glycerinum and visualized by confocal laser scanning microscopy.

Wound healing

Wound healing experiments were performed for quantitative investigation of the migration of MSC when stimulated by TiO₂ nanoparticles of different sizes. MSC were incubated without (control) or with the different TiO₂ nanoparticles (0.10 mg/mL) at 37°C for 4 hours. The initial cell seeding density was 1×10^5 cells/cm². A linear wound was made by scratching with a cell scraper. The cells were washed twice with phosphate-buffered solution and incubated with fresh medium for another 12 and 16 hours, respectively. Cells were then visualized by confocal laser scanning microscopy.

Transwell assay

A further cell migration assay was performed as reported elsewhere.²⁷ MSC were cultured with TiO₂ nanoparticles (0.10 mg/mL) for 8 hours. After detaching with trypsin, the treated cells were seeded into a Millicell (Greiner Bio-One, Frickenhausen, Germany) containing pores 8 μ m in diameter at an initial seeding density of 1×10^5 cells/cm² suspended in Dulbecco's Modified Eagle's Medium containing 1%

fetal bovine serum. Next, the Millicell was put into the well of a 24-well plate containing 20% fetal bovine serum in Dulbecco's Modified Eagle's Medium. The cells were allowed to migrate at 37°C for another 8 hours. After staining with crystal violet (0.1 wt%), the migrated cell clones were recorded using an optical microscope. The cell number was counted at 12 different areas, and the data were averaged from three parallel experiments and normalized by data from the control group.

Total intracellular protein content

MSC were cultured without or with TiO₂ nanoparticles (0.10 mg/mL) in Dulbecco's Modified Eagle's Medium with 10% fetal bovine serum for 7 and 14 days, respectively. The initial cell seeding density was 2×10^4 cells/cm². At prescribed time intervals, the cells were lysed using 1% Triton X-100 with three freeze-thaw cycles. Total protein content in the cell lysates was determined spectrophotometrically using a commercially available kit (bicinchoninic acid, Sigma-Aldrich). The light absorbance of the samples was measured at 570 nm with the Bio-Rad 550 spectrophotometric microplate reader. Total intracellular protein (expressed as mg) synthesized by the cells was determined from a standard curve of absorbance versus the known concentration of albumin run in parallel experiments.

Alkaline phosphatase activity

Cells were incubated with 0.10 mg/mL TiO₂ particles for 7 and 14 days, respectively. The initial seeding density was 2×10^4 cells/cm². Cells were then collected and lysed as mentioned above. The supernatant was used for determining ALP activity, with paranitrophenyl phosphate as a substrate. Absorbance at 405 nm was measured using a spectrophotometer in a 96-well microplate reader (Bio-Rad 550). ALP activity (expressed as nmol of p-nitrophenol converted per minute) was normalized by total intracellular protein synthesis and expressed as mol p-nitrophenol per minute per mg protein. The ALP activity of MSC cultured without TiO₂ particles was used as the control.

Mineralization

Cell mineralization was quantitatively measured after alizarin red-S staining elution as described elsewhere.²⁸ Briefly, MSC were incubated with TiO₂ particles (0.10 mg/mL) for 15 days at an initial seeding density of 5×10^4 cells/cm². The cells were washed with phosphate-buffered solution and fixed with 2% glutaraldehyde at 4°C for 20 minutes. The samples were then washed three times in distilled water and stained

with 40 mM alizarin red-S (pH 4.1) with shaking at room temperature for 20 minutes. Next, 400 µL of 10% acetic acid was added to each well and incubated for another 30 minutes with shaking. The cell monolayer was then scraped with a cell scraper and transferred into a 1.5 mL tube with 10% acetic acid. Samples were heated up to 85°C for 10 minutes and centrifuged at 20,000 g for 15 minutes. Subsequently, 300 µL of the supernatant was transferred into a new tube and neutralized with 200 µL of 10% ammonium hydroxide. The absorbance of supernatant (100 µL) was measured at a wavelength of 405 nm using a spectrophotometric microplate reader (Bio-Rad 680). Cell mineralization was finally normalized for protein content.

Reverse transcription polymerase chain reaction

Cells were collected after culture for 14 days. The total RNA in the cells was extracted according to the instructions on an RNA extract kit (Bio-Tek Instruments Inc., Winooski, VT, USA). Aliquots of total RNA (5 mL) were reverse transcribed, and a 5 mL cDNA sample was amplified. The following oligonucleotide primer pairs related to the targeted RNA were used in this study: β -actin, 5'-ATATCGCTGCGCTGGTCGTC-3' and 5'-AGGATGGCGTGAGGGAGAGC-3'; osteocalcin, 5'-TGGCCCTGACTGCATTCTGC-3' and 5'-GCTGTGCCGTCCATACTTTCG-3'; and osteopontin, 5'-CCTCTGAAGAAACGGAT GACT-3' and 5'-CTGTGTGTTCCACGCTT-3'. The polymerase chain reaction conditions were as follows: first cycle at 94°C for 5 minutes, followed by 35 cycles at 94°C for 30 seconds, 56°C for 30 seconds, and 72°C for 30 seconds, and the final cycle at 72°C for 10 minutes. The amplified cDNA was analyzed by gel electrophoresis with 1.5% agarose gel.

Statistical analysis

All data are expressed as the mean \pm standard deviation. The statistical analysis was performed using the Student's *t*-test and one-way analysis of variance at confidence levels of 95% and 99% (OriginPro, version 7.5).

Results

Synthesis and characterization of TiO₂ nanoparticles

TiO₂ nanoparticles of different sizes were home-synthesized by a hydrothermal method,²⁹ and characterized by scanning and transmission electron microscopy, respectively. Three types of TiO₂ nanoparticles with average particle sizes of

14 nm, 108 nm, and 196 nm were synthesized by simply changing the ratios of peptizing agents, titanium precursor solution (titanium butoxide), and reaction time. Scanning and transmission electron microscopic images (Figure 1) show that the TiO₂ nanoparticles had a relatively uniform morphology with a narrow size distribution.

Cell viability and proliferation assay

A CCK-8 kit was used to investigate the influence of TiO₂ nanoparticles on the biological behavior of MSC, ie, their

potential cytotoxicity. First, we investigated the dose-dependent effects of TiO₂ nanoparticles on the viability of MSC. The results show that high concentrations of TiO₂ nanoparticles (0.10 and 0.20 mg/mL) had strong adverse effects on the viability of MSC (Figure 2).

Second, we investigated the effects of TiO₂ nanoparticle size on cell proliferation (Figure 3). MSC were cultured with TiO₂ nanoparticles (0.10 mg/mL) for 3, 7, and 14 days. The results show that the negative effect increased significantly with increasing particle size. After culture for 3 days with TiO₂ nanoparticles 196 nm in size, relative cell viability decreased to 51.73% compared with the control group. Relative cell viability decreased further to 36.39% and 34.92% after culture for 7 and 14 days, respectively. Over all time intervals, the TiO₂ nanoparticles showed a clear tendency for size-dependent inhibition of growth ($P < 0.01$).

Third, we investigated the influence of different-sized TiO₂ nanoparticles on cell division by flow cytometry to show the cell cycle during proliferation. Previous research has suggested that metal nanoparticles might potentially induce oxidative stress, leading to intracellular changes and alterations in the cell cycle.³⁰ In this study, we measured the change in percent proliferation index (percent synthesis phase of DNA [%S] + percent post-synthesis phase of DNA [%G₂]) in the cell cycle to indicate the effects of the TiO₂ nanoparticles on MSC.

The percentage of TiO₂ nanoparticle-treated MSC in G1 phase increased when compared with that of the control after culture for one, two, and three days, respectively. Accordingly, the percent proliferation index (%S + %G₂) of MSC treated with TiO₂ nanoparticles decreased after culture for one, two, and three days (Table 1). Further, inhibition of

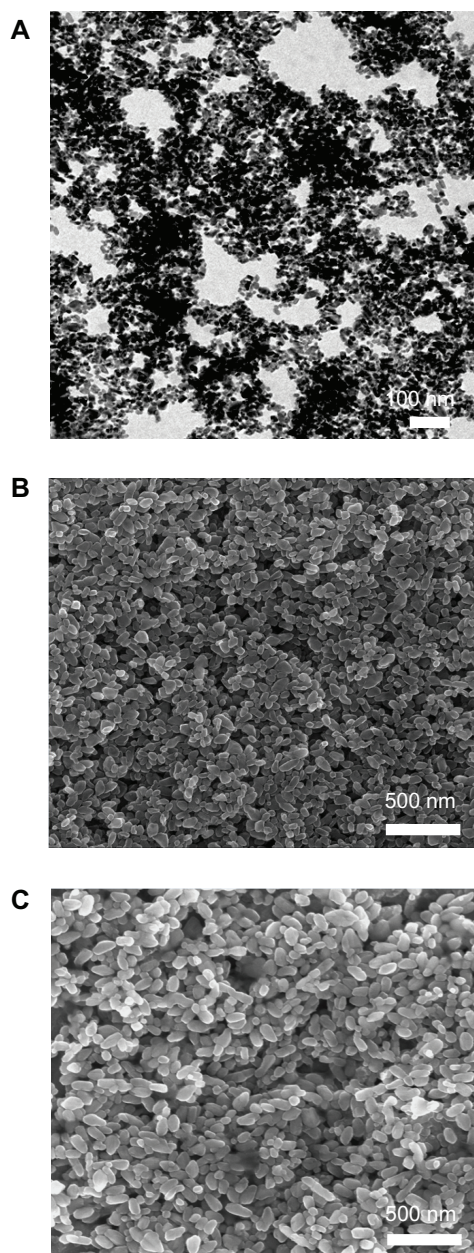


Figure 1 Characterization of TiO₂ nanoparticles with different sizes from scanning electron microscopy observation (except for A from transmission electron microscopy observation). (A) 14 nm, (B) 108 nm, and (C) 196 nm.

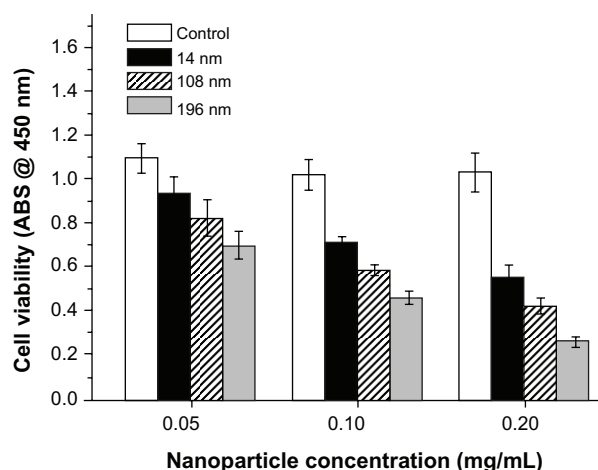


Figure 2 Dose-dependent effects of various nanoparticles on cell viability after culture for 2 days. The initial cell seeding density was 2×10^4 cells/cm². **Abbreviation:** ABS, absorbance.

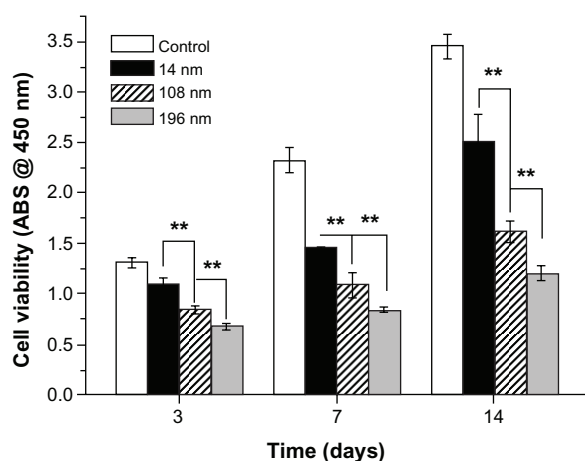


Figure 3 Viability of mesenchymal stem cells incubated with various TiO_2 nanoparticles for different time periods.

Notes: The initial cell seeding density was 2×10^4 cells/cm². Error bars represent the mean \pm standard deviation for $n = 6$, $**P < 0.01$.

Abbreviation: ABS, absorbance.

proliferation of the TiO_2 nanoparticles was also dependent on their particle size at each time interval. In particular, the percent proliferation index of MSC treated with 196 nm TiO_2 nanoparticles decreased 2.51-fold compared with controls after culture for 3 days. This result indicates that the change in percent proliferation index for the MSC was mediated mainly by the size of the TiO_2 nanoparticles. This finding was consistent with a previous study showing that polystyrene nanoparticle size and charge had a pronounced effect on mitotic spindle formation and chromosomes during cell division.³¹ Moreover, other studies have demonstrated that DNA strand breaks were related to cell cycle arrest and cell death in epithelial cells treated with TiO_2 nanoparticles and copper dioxide nanoparticles.^{32,33}

Table I Cell cycle analysis. Cells were incubated without (control) or with various titanium dioxide nanoparticles for one, two, and three days, respectively

Time (days)	Samples	G ₁ %	S%	G ₂ %	PI (S + G ₂)%
1	Control	71.30	18.60	10.10	28.70
	14 nm	72.30	12.30	15.40	27.70
	108 nm	77.24	5.76	17.00	22.76
	196 nm	82.08	3.72	14.20	17.92
2	Control	69.80	11.60	18.60	30.20
	14 nm	73.51	5.79	20.70	26.49
	108 nm	79.41	8.79	11.80	20.59
	196 nm	84.23	8.07	7.70	15.77
3	Control	67.39	12.20	20.40	32.60
	14 nm	75.85	8.050	16.10	24.15
	108 nm	78.05	13.00	8.95	21.95
	196 nm	87.00	5.21	7.79	13.00

Notes: Initial cell seeding density was 1×10^5 cells/cm². DNA content was analyzed by flow cytometry. G₁, S, and G₂ indicate cell cycle phases.

Abbreviation: PI, proliferation index; DNA, deoxyribonucleic acid.

Particle internalization and cell adhesion

To identify the location of TiO_2 nanoparticles within cells and the effect of particle size on adhesion of MSC, FITC- TiO_2 particles (green), vinculin (focal adhesion protein, cyan), actin (cytoskeleton) filaments (red), and cell nuclei (blue) were visualized by confocal laser scanning microscopy. MSC cultured without (control) or with FITC- TiO_2 nanoparticles of different sizes were observed after culture for 12 and 24 hours (Figure 4A), respectively. Large amounts of FITC- TiO_2 nanoparticles (108 nm and 196 nm) were observed to accumulate in MSC (green). Moreover, the digested nanoparticles were only distributed in the cytoplasm rather than the nuclei of MSC. The same phenomenon was also observed in previous studies of neural stem cells²⁴ and fibroblasts.³⁴

Adhesion is the first response of the cell when it comes into contact with a biomaterial, and plays an essential role in regulation of the subsequent biological behavior of cells. Previous research indicates that the adaptor protein of vinculin is a key regulator of focal adhesion, which is activated by binding of talin.³⁵ Compared with the controls, a distinct difference was observed regarding cell adhesion (vinculin spots, arrows) and the cytoskeletons of MSC treated with TiO_2 nanoparticles. The larger the size of the TiO_2 nanoparticles, the lower the amount of vinculin that was present (Figure 4A). A similar tendency was also observed for cell spreading, indicated by distribution of actin within cells.

The amount of vinculin in MSC treated with 14 nm TiO_2 nanoparticles was significantly greater ($P < 0.01$) than that in MSC treated with 108 nm TiO_2 nanoparticles, and yet greater ($P < 0.01$) in MSC treated with 196 nm TiO_2 nanoparticles after culture for 24 hours (Figure 4B). The average amount of vinculin in MSC treated with 196 nm TiO_2 nanoparticles decreased to 63.4% of control levels after culture for 12 hours, and decreased further to 51.3% of control levels after culture for 24 hours. These results suggest that TiO_2 nanoparticle size strongly affects adhesion of MSC, which is consistent with previous reports.^{16,36}

Cell migration

Focal adhesion of cells generally has a close relationship with cell migration.³⁷ To investigate the effect of TiO_2 nanoparticles on cell migration, a wound healing (scrape migration) assay was performed, as described elsewhere.³⁸ After culture for 12 hours, many cells treated with 14 nm TiO_2 nanoparticles migrated to the center of the wound field, whereas few cells treated with 108 nm TiO_2 nanoparticles and even fewer cells treated with 196 nm TiO_2 nanoparticles did so (Figure 5).

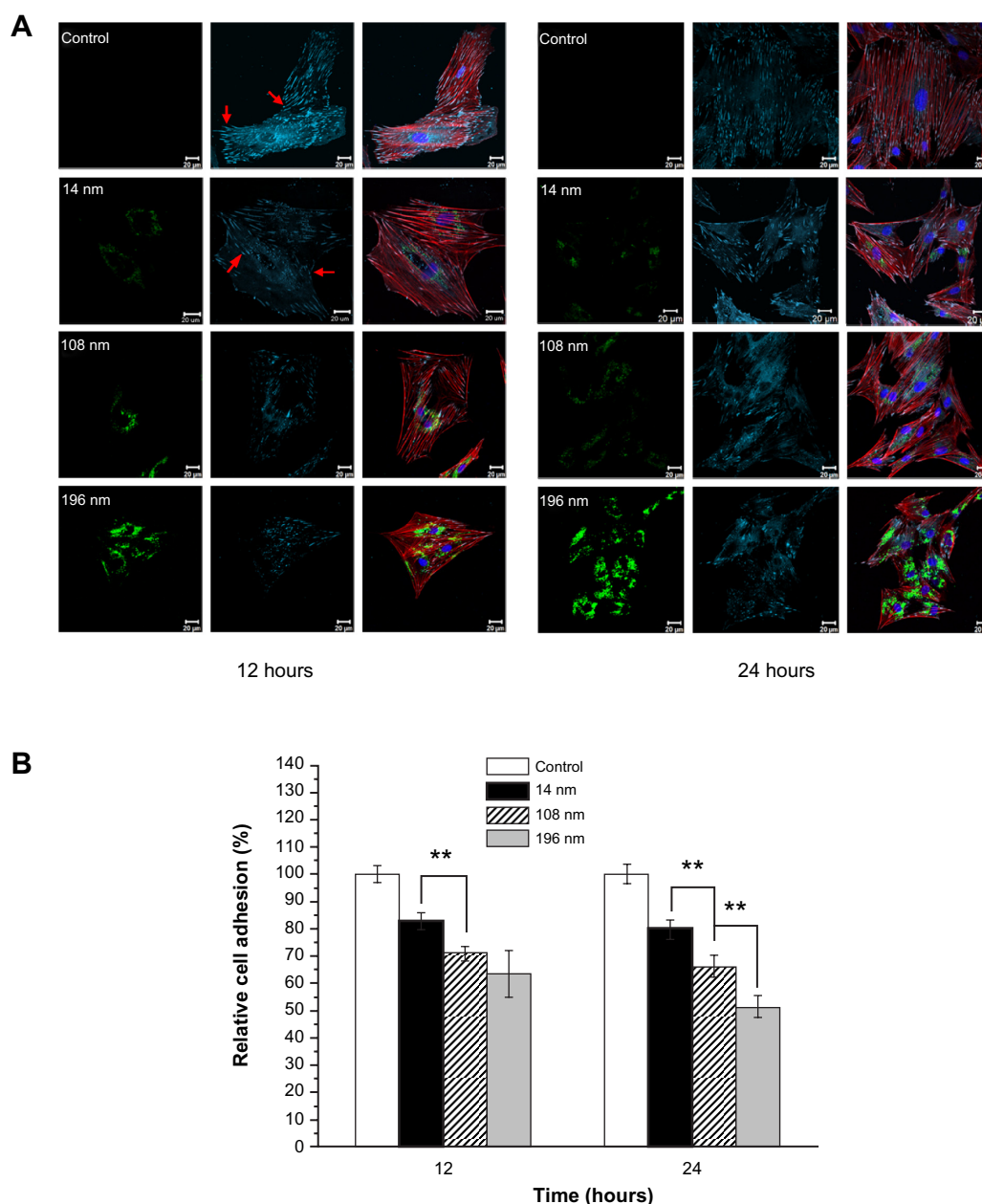


Figure 4 (A) Confocal microscopic images of mesenchymal stem cells incubated without (control) or with FITC-TiO₂ particles (0.10 mg/mL) for 12 and 24 hours, respectively. FITC-TiO₂ particles (green), vinculin (cyan), actin filaments (red), and cell nuclei (blue). Scale bar = 20 μm. **(B)** Relative fluorescence intensity analysis of vinculin. Average fluorescence intensity per cell was normalized to the control sample.

Notes: Error bars represent the mean ± standard deviation for $n = 50$, ** $P < 0.01$.

Abbreviation: FITC, fluorescein isothiocyanate.

To investigate further the effects of TiO₂ nanoparticles on cell migration, a matrigel invasion assay was performed using Transwell chambers. Figure 6A shows representative images for control MSC and those treated with TiO₂ nanoparticles. Quantitative analysis showed that treatment with TiO₂ nanoparticles significantly decreased migration of MSC. Moreover, the relative migration of MSC treated with 14 nm TiO₂ nanoparticles was significantly greater ($P < 0.01$) than that of MSC treated with 108 nm TiO₂ nanoparticles, and

yet higher ($P < 0.01$) than migration of MSC treated with 196 nm TiO₂ nanoparticles (Figure 6B). Our results suggest that TiO₂ nanoparticle size strongly affects migration of MSC, and are consistent with those of a previous study of hydroxyapatite nanoparticles.³⁹

Cell differentiation

To investigate the effects of TiO₂ nanoparticles on osteogenic differentiation of MSC, we firstly measured ALP activity

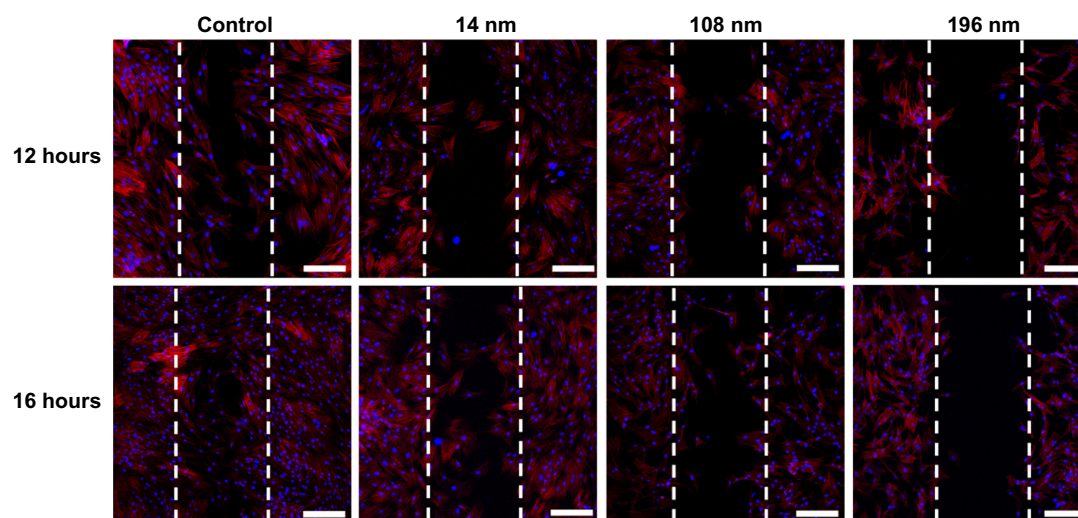


Figure 5 Wound healing in mesenchymal stem cells incubated without (control) or with different TiO_2 particles (0.10 mg/mL) shown by confocal laser scanning microscopy. **Notes:** Scale bar = 100 μm . The initial cell seeding density was 1×10^5 cells/ cm^2 .

in MSC treated with different-sized TiO_2 nanoparticles (Figure 7). ALP is an early mineralization-related protein marker for osteogenesis of osteoblasts.⁴⁰ MSC cultured with different-sized TiO_2 nanoparticles showed significantly weaker ($P < 0.01$) ALP activity than the control group at all time points. After culture for 7 days, MSC treated with 14 nm and 108 nm TiO_2 nanoparticles demonstrated significantly higher ($P < 0.01$) ALP activity than those treated with 196 nm TiO_2 nanoparticles. After culture for 14 days, MSC treated with 14 nm TiO_2 nanoparticles showed significantly more ($P < 0.01$) ALP activity than those treated with 108 nm and 196 nm TiO_2 nanoparticles. Next, we measured mineralization ability in MSC treated with TiO_2 nanoparticles of different sizes. The normalized mineralization demonstrated a trend similar to that for ALP activity after culture for 15 days (Figure 8).

To investigate further the impact of TiO_2 nanoparticles on differentiation of MSC, we characterized messenger RNA (mRNA) expression of osteocalcin and osteopontin by reverse transcription polymerase chain reaction. Osteocalcin and osteopontin are noncollagenous matrix proteins, which are regarded as mature osteoblast markers in MSC at a late stage.⁴¹ We found that MSC treated with 196 nm TiO_2 nanoparticles produced significantly less ($P < 0.05$ and $P < 0.01$) osteocalcin and osteopontin, respectively, than MSC treated with 14 nm and 108 nm TiO_2 nanoparticles (Figure 9). This result is consistent with the findings of a previous study showing that calcium phosphate-based particles of increasing size impaired osteogenic differentiation and bone matrix mineralization in MSC.⁴² However, other research has confirmed that a high dose of zinc oxide nanoparticles is lethal

to proliferating pluripotent MSC but has negligible toxicity in osteogenically differentiated MSC.⁴³ Nanohydroxyapatite and nanohydroxyapatite-poly(lactic-co-glycolic acid) (PLGA) composites promote osteogenic differentiation of human MSC.⁴⁴ In another study, iron oxide nanoparticles of different sizes and surface charge were found to regulate the differentiation of MSC.⁴⁵

Discussion

In recent years, much effort has been made to improve the properties of nanostructured titanium implants.^{3,7,8,10} Nevertheless, nanoscale wear debris could potentially be generated during long-term implantation in the clinical setting, mainly because of abrasion and invasion of various enzymes and chemicals in an environment containing biological fluids. Therefore, investigation of the potential cytotoxicity of nanoscale debris released from a titanium implant is essential.

The head of a titanium joint implant not only comes into contact with osteoblasts derived from the surrounding bone tissue but also interacts with bone marrow stem cells in the long bone cavity. In a previous study,²³ we highlighted the size-dependent negative effects of different titanium nanoparticles on viability of osteoblasts, liver cells (L-02), and kidney cells (HEK 293), and investigated the size-dependent cellular uptake pathways and activation of matrix metalloproteinase 9. In this study, we used titanium nanoparticles (ie, a debris model) to investigate how titanium nanoparticles of different sizes affect adhesion and proliferation (ie, DNA synthesis), osteocalcin and osteopontin gene expression, and migration of MSC. Investigating the migration of MSC surrounding a titanium implant is of particular importance for

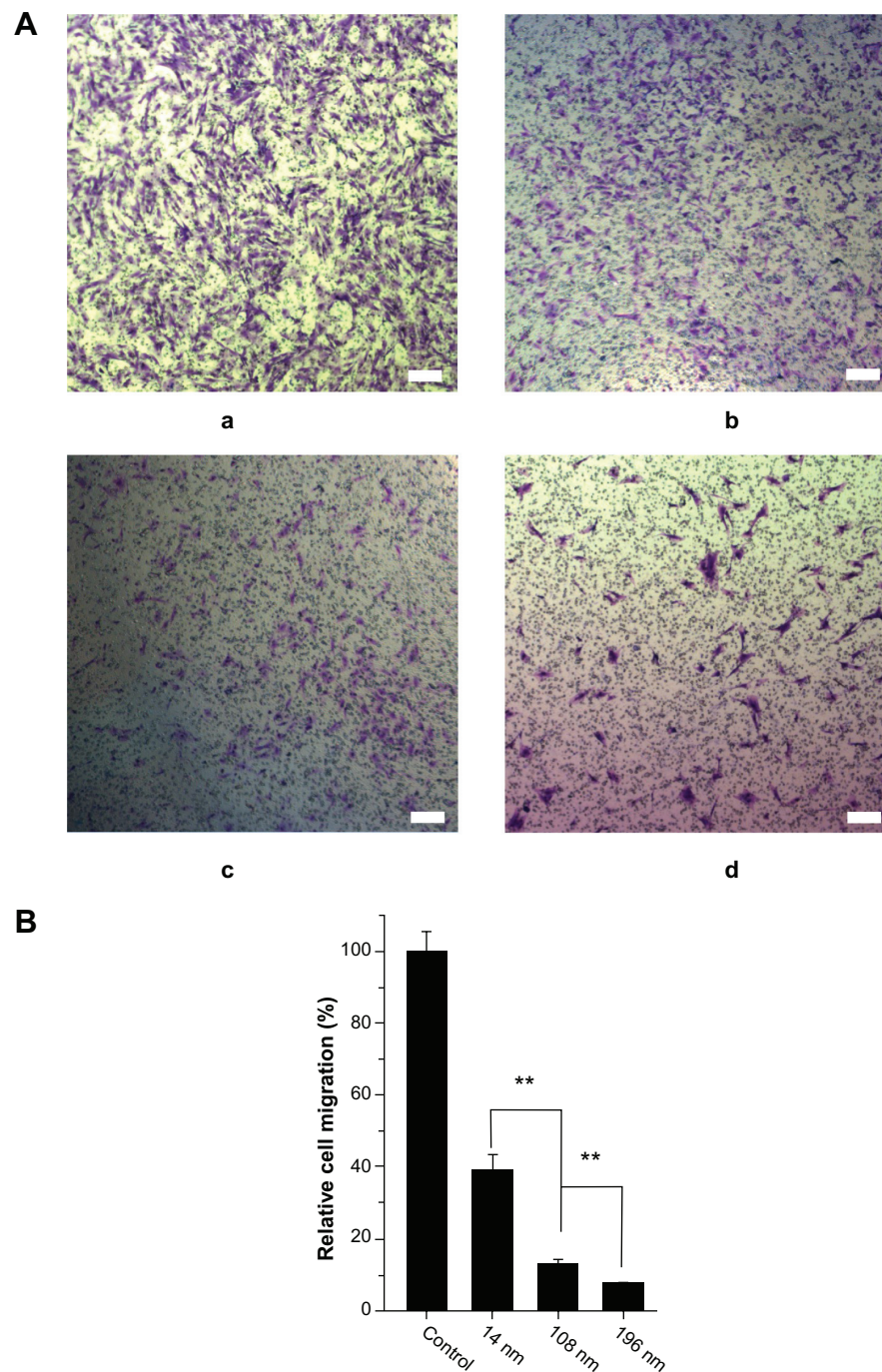


Figure 6 Characterization of cell migration. (A) Qualitative cell migration assay shown by Transwell assay after culture for 8 hours. a: control; b: 14 nm; c: 108 nm; d: 196 nm; and (B) quantitative analysis of relative cell migration from A (n = 100).

Notes: The initial cell seeding density was 1×10^5 cells/cm². Scale bar = 100 μ m. ** $P < 0.01$.

understanding how MSC potentially contribute to osseous integration in implant/bone tissue.

Once wear debris (ie, TiO₂ nanoparticle model) detaches from a titanium implant, it can affect the biological function of the surrounding cells in two ways, firstly by affecting protein adsorption around cells and secondly by being taken up directly by cells. The extent of this in both cases depends

on particle size and nanotopography. However, the exact contribution of each factor needs to be investigated in more depth. Our interest was focused on the latter in this study.

Our results confirmed that the synthesized TiO₂ nanoparticles (14 nm, 108 nm, and 196 nm) had a cytotoxic effect on MSC depending on their concentration (Figure 2). This phenomenon could be interpreted as highly concentrated TiO₂

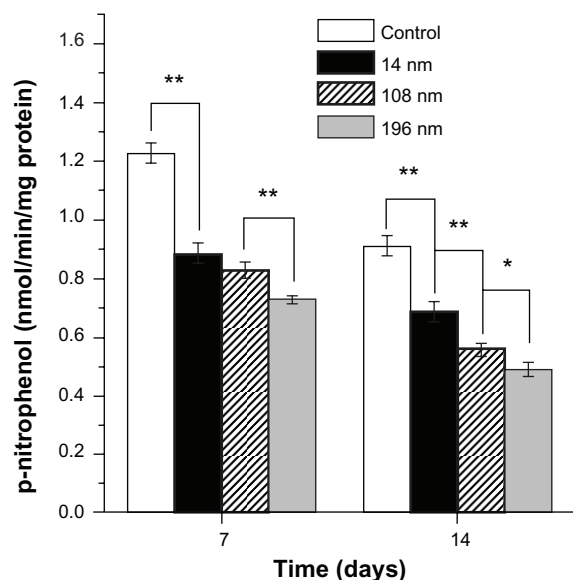


Figure 7 Alkaline phosphatase activity of mesenchymal stem cells cultured with different-sized TiO_2 nanoparticles (0.10 mg/mL) for 7 and 14 days, respectively. **Notes:** The initial cell seeding density was 2×10^4 cells/cm². Error bars represent the mean \pm standard deviation for $n = 6$, * $P < 0.05$, ** $P < 0.01$.

nanoparticles producing oxidative stress as well as reactive oxygen species, leading to cell death.³⁴ Furthermore, we confirmed that growth and division of MSC were negatively affected in a size-dependent manner by the TiO_2 nanoparticles (Figure 3 and Table 1) potentially via a mechanism causing intracellular acidity, thus inhibiting cell proliferation.⁴⁶

It is well known that uptake of nanoparticles can occur via different pathways in cells, depending on the nanoparticle dimensions. In a previous study, we confirmed that large TiO_2 nanoparticles (74 nm and 196 nm) were taken up by osteoblasts mainly via clathrin-mediated and caveolae-mediated

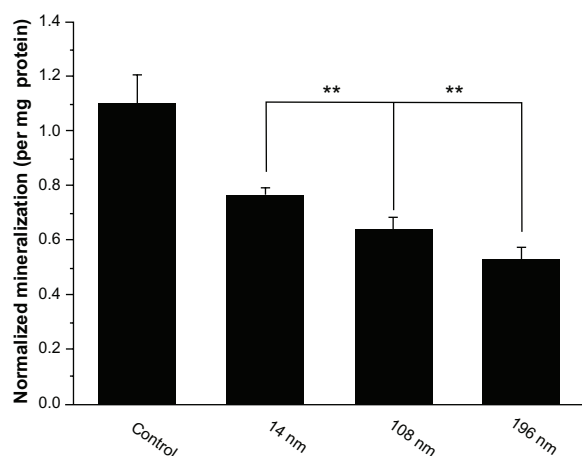


Figure 8 Quantification of cell mineralization. Mesenchymal stem cells were cultured without (control) or with different TiO_2 particles (0.10 mg/mL) for 15 days. **Notes:** The initial cell seeding density was 5×10^4 cells/cm². Error bars represent the mean \pm standard deviation for $n = 6$, ** $P < 0.01$.

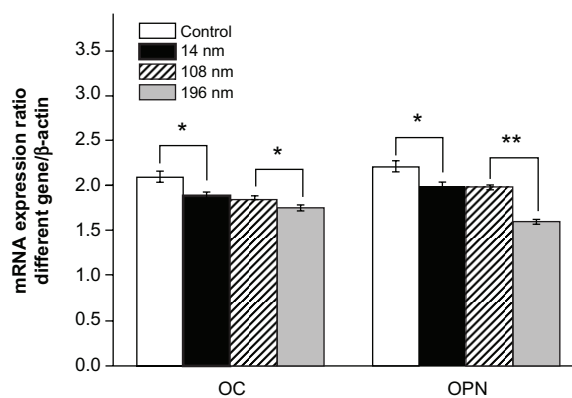


Figure 9 Relative mRNA expression of osteocalcin and osteopontin analyzed by real-time polymerase chain reaction. The mesenchymal stem cells were cultured without or with different TiO_2 particles (0.10 mg/mL) for 14 days. **Notes:** The initial cell seeding density was 5×10^4 cells/cm². Error bars represent the mean \pm standard deviation for $n = 5$, * $P < 0.05$, ** $P < 0.01$. **Abbreviations:** mRNA, messenger RNA; OC, osteocalcin; OPN, osteopontin.

pathways, whereas small TiO_2 nanoparticles (14 nm) were digested by osteoblasts through a caveolae-mediated pathway.²³ We assume that TiO_2 nanoparticles were taken up by MSC in a similar manner. In our study, we found that greater amounts of 196 nm TiO_2 nanoparticles accumulated within MSC than did 108 nm and 14 nm TiO_2 nanoparticles (Figure 4A). Previously, Win et al demonstrated that Caco-2 cells took up much fewer 50 nm PLGA nanoparticles than particles with sizes of 200 and 500 nm.⁴⁷ Unlike the small (14 nm) TiO_2 nanoparticles, high accumulation of large (196 nm) TiO_2 nanoparticles within MSC led directly to more cytotoxicity, in turn affecting biological functions, such as migration, differentiation, and gene expression (Figures 4–9).

The relatively low accumulation of small-sized (14 nm) TiO_2 nanoparticles might be interpreted as meaning that small nanoparticles could escape more easily from endocytic vesicles in cells, thus resulting in limited cytotoxicity to MSC. Pan et al confirmed that 1.2 nm gold nanoparticles were markedly less cytotoxic than 1.4 nm gold nanoparticles, despite a three-fold higher concentration.⁴⁸ Another example was internalization of FITC-mesoporous silica nanoparticles that did not affect cell viability, proliferation, or differentiation potential of MSC.⁴⁹ An explanation for this is that FITC-mesoporous silica nanoparticles could escape endosomal vesicles in MSC and retain their architectonic integrity after internalization.

On the other hand, the physicochemical properties (chemistry and surface charge) of a nanoparticle could affect its escape mechanism and potential cytotoxicity.⁵⁰ A previous study showed that poly-L-lysine-functionalized PLGA nanoparticles could escape more rapidly and efficiently

from endosomes than native PLGA nanoparticles and result in 10-fold higher intracellular delivery of an encapsulated model protein.⁵¹

Although the positive effects of the nanotopography of biomaterials on cell function are well documented, for instance, cell density and cell adhesion are greatly enhanced by surface-nanostructured biomaterials,^{52,53} the present study implies that loss of such a nanoscale structure may have negative effects on cell and tissue function. Thus, ensuring the stability of a nanostructured entity is one of the major concerns when nanostructuring a biomaterial or exploiting a new nanostructured biomaterial, in particular inorganic materials.

Conclusion

In this study, the cytotoxicity of a series of TiO₂ nanoparticles was systematically investigated from the cellular to the molecular level. We confirmed that TiO₂ nanoparticles of different sizes have a negative impact on viability, adhesion, migration, proliferation, and differentiation of MSC in a size-dependent and dose-dependent manner. Accumulation of such knowledge might provide new insights into the development of novel nanostructured titanium implants.

Acknowledgments

This work was financially supported by the China Ministry of Science and Technology (973 Project 2009CB930000), Natural Science Foundation of Chongqing Municipal Government (CSTC, 2011JJQ10004), Natural Science Foundation of China (11032012, 31170923, 51173216), National Key Technology R&D Program of the Ministry of Science and Technology (2012BAI18B04), Fok Ying Tung Education Foundation (121035), and the “111” Project (B06023).

Disclosure

The authors report no conflicts of interest in this work.

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