In vitro apoptotic and DNA damaging potential of nanobarium oxide

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Abstract: Barium oxide nanoparticles (BaONPs) are an important industrial compound and are widely used in polymers and paints. In this study, apoptotic and genotoxic effects of BaONPs in mouse embryonic fibroblast (L929) cells were determined by using single-cell gel test. In vitro cytotoxicity assays were performed to assess BaONPs’ toxicity in L929 cells. Mild cytotoxicity was observed in L929 cells due to BaONPs. BaONPs increased lipid peroxidation, catalase, and superoxide dismutase levels and lowered glutathione levels in L929 cells. This was accompanied by concomitant generation of reactive oxygen species and activation of caspase-3 in BaONPs-treated L929 cells. On the other hand, when we exposed L929 cells to BaONPs for 24 and 48 hours (comet assay), there was a duration- and dose-dependent increase in DNA impairment detected in the single-cell gel test. Thus, BaONPs exhibit genotoxic and apoptotic effects in L929 cells, most likely due to initiation of oxidative damage.

Keywords: nanobarium oxide, L929 cells, DNA damage, apoptosis

Introduction
Engineered metal nanoparticles have been widely used in cosmetics, painting, textiles, and water treatments. Barium oxide nanoparticles (BaONPs) have been used in diagnostic imaging, orthopedic medicines, and other applications. Because of their increased use in human applications and their release into the environment, it has become important to study and understand the impact of nanoparticles on animal health. Hiraoka et al reported that barium induces automaticity in ventricular muscle. Nayler and Grinwald reported that marked contractures develop in the heart of rat, with depletion of tissue ATP and creatinine phosphate after perfusion with barium (1.3 mM/L) in place of calcium ions. 

Even though there are existing studies on the toxicity of barium nanoparticles, the underlying machinery inducing toxicity remains unexplored till now. Moreover none of the studies, so far, have explored the adverse effect of BaONPs in mouse embryonic fibroblast cells. Ostrovsky et al reported that oxidative stress and reactive oxygen species (ROS) cause adverse effects such as apoptosis and DNA fragmentation. The toxic potential of BaONPs is of specific concern because the changes of the genetic materials have potential to cause cell death, cancer development, and reproductive adverse effects. ROS has been explained as a concerning factor in BaONPs toxicity. We have already found glutathione (GSH) reduced as an oxidative biomarker and ROS as a group marker of $O_2^-$, OH radicals, $H_2O_2$ and malondialdehyde (MDA) in reaction to BaONPs exposure.

So, this work was planned to measure the toxicity and genotoxic effect of BaONPs in L929 cells.
Materials and methods

Ethics
Ethics approval was not sought, as our University does not require it for in vitro studies.

Chemicals
BaONPs, 2, 7-dichlorofluorescin diacetate, DTNB, glutathione, and propidium iodide were acquired from Sigma-Aldrich, St Louis, MO, USA. Dulbecco’s Modified Eagle’s Medium (DMEM)/F-12 medium, were bought from Invitrogen Co. (Carlsbad, CA, USA).

Characterization of BaONPs
A suspension of BaONPs was prepared in DMEM (1 mg/mL) and sonicated in a sonicator with 40 W capacity for 15 minutes at room temperature to form a uniform suspension. The size of BaONPs was characterized by field-emission transmission electron microscopy (FETEM). The hydrodynamic size of BaONPs were checked by dynamic light scattering (DLS; Nano-Zeta Sizer-HT, Malvern Instruments, Malvern, UK).

L929 cells
L929 cells were procured from ATCC (Rockville, MD, USA).

Morphological analysis of cells
The shape and size of L929 cells were observed by microscope (Leica DMIL, Leica Microsystems, Wetzlar, Germany) after exposure to BaONPs for 24 and 48 hours.

MTT assay
The cytotoxicity of L929 cells due to exposure to BaONPs (0, 50, 100, 150, and 300 μg/mL) for 24 and 48 hours was measured by MTT assay.

LDH assay
Leakage of cytoplasmic lactate dehydrogenase (LDH) enzyme in L929 cells due to treatment with BaONPs (0, 50, 100, 150, and 300 μg/mL) for 24 and 48 hours was determined.

Quantitation of ROS
ROS production was evaluated in L929 cells after treatment with BaONPs (0, 50, 100, 150, and 300 μg/mL) by using DCFH-DA dye. To confirm the role of ROS, the cells were exposed to BaONPs (300 μg/mL) with and without 5 mM N-acetylcysteine (NAC) for 24 hours.

Estimation of oxidative parameters
L929 cells, (~5×10⁶) seeded in a 75 cm² flask, were treated with BaONPs (0, 50, 100, 150, and 300 μg/mL) for 24 and 48 hours. After treatment, the cells were scraped and washed with ice-cold phosphate-buffered saline (PBS). Scraped cells were broken down in lysing. These cells were centrifuged for 15 minutes at 12,000× g, 4°C, and cell lysate was kept at −20°C for testing oxidative stress parameters. The amount of protein in cell lysate was quantified by Bradford’s method using bovine serum albumin as the standard.

Lipid peroxidation was evaluated by estimating the production of MDA by Ohkawa et al’s method. Superoxide dismutase (SOD) and catalase (CAT) levels were measured by Alarifi et al’s method. Glutathione level was estimated by using Ellman’s reagent.

Chromosome condensation
Condensation of chromosomes in L929 cells due to BaONPs exposure (0, 50, 100, 150, and 300 μg/mL) was observed using Hoechst 33342 dye. Treated and untreated cells were stained with Hoechst 33342 for 30 minutes at 37°C in dark. Photomicrographs of nuclei were taken by fluorescent microscope.

Cysteine-aspartic acid protease-3 activity
Cells (5×10⁶ cells/well) were spread on a culture plate (96 well) and incubated at 37°C. Cells were exposed to different concentrations of BaONPs (0, 50, 100, 150, and 300 μg/mL) for 48 hours. Caspase-3 level was assessed by Alarifi and Ali’s method. To confirm the role of ROS in caspase-3 activity, L929 cells were exposed to BaONPs (150 μg/mL) with and without 5 mM NAC for 24 hours.

Analysis of apoptosis through fluorescence-activated cell sorting
Apoptosis was detected by using the Annexin V–FITC (fluorescein isothiocyanate) kit (Cayman Chemicals, Ann Arbor, MI, USA) according to manufacturer’s information. After exposure to BaONPs (150 μg/mL) for 24 and 48 hours, cells were trypsinized and washed with PBS and then incubated with binding buffer (400 μL) containing propidium iodide (20 μL) and Annexin V FITC (10 μL) for 30 minutes in the dark at room temperature. The density of Annexin V–FITC-positive L929 cells was scrutinized by using fluorescence-activated cell sorting (FACS) (BD Biosciences, San Jose, CA, USA). Cells in the initial phase of apoptosis were found in Q₁ quadrant, while those in the delayed phase were found in Q₄. The percentage of late apoptotic cells was plotted.
Comet assay
Single-cell gel test was completed by Ali et al’s\textsuperscript{14} method.

Analysis of data by statistics
A minimum of three independent experiments in two replicates were done for each result. Results are presented as average (± standard error) and investigated by analysis of variance. \(P\)-value less than 0.05 reflected statistical significance.

Results
Changes in L929 cell lines
Figure 1 unveils morphological changes in control cells and L929 cells treated with BaONPs for 48 hours. BaONPs (300 μg/mL)-treated L929 cells changed into round shape and separated from the outer layers of the culture flask (Figure 1B and C).

L929 cells toxicity
BaONPs-induced cytotoxicity in L929 cells was observed as MTT declined, releasing lactate dehydrogenase enzymes. MTT results indicated a dose- and time-dependent cell toxicity in the wake of L929 cells were treated with BaONPs (Figure 2A). LDH discharge, a biomarker of cellular membrane damage, was estimated in L929 cells after exposure to BaONPs for 48 hours.

Oxidative stress
Induction of oxidative stress by BaONPs was evaluated by estimating the glutathione, lipid peroxidation, ROS, catalase, and superoxide dismutase levels in L929 cells. BaONPs incited ROS production in a dose- and duration-dependent manner (Figure 3). BaONPs-induced oxidative stress was further evidenced by depletion of glutathione and raise of lipid peroxide, superoxide dismutase, and catalase due to BaONPs exposure (Figure 4).

Cysteine-aspartic acid protease-3 enzyme and apoptosis
Caspase-3 enzyme shows main activity in apoptotic cells, which was caused due to BaONPs (Figure 5). After being exposed to BaONPs for 48 hours, the role of Cysteine-aspartic acid protease-3 increased in a dose- and duration-dependent manner.

Staining of Annexin V-FITC and PI were applied to detect apoptotic and necrotic stimulation in L929 cells following BaONPs (150 μg/mL) exposure for 24 and 48 hours (Figure 6). Treatment with BaONPs indicated a significant

![Figure 1](image_url) Shape and size of L929 cells.

Notes: (A) Untreated cells. (B) Cells with BaONPs (150 μg/mL) for 24 hours. (C) Cells with BaONPs (150 μg/mL) for 48 hours. Scale bar (——) 50 μm. The arrows indicate damaged cells.

Abbreviation: BaONPs, barium oxide nanoparticles.
Figure 2 Cytotoxicity of BaONPs in mouse fibroblast cells over 48 hours, as assessed by (A) MTT and (B) LDH assays.

Notes: Data represent the average ± SE of three tests. Statistical differences with respect to the controls are shown (*P<0.05).

Abbreviations: BaONPs, barium oxide nanoparticles; LDH, lactate dehydrogenase; SE, standard error.

(P<0.05) increase in late apoptotic cells in a duration-dependent manner (Figure 6B–D).

Impairment of DNA

DNA impairment was assessed as % tail DNA in untreated and BaONPs-treated L929 cells. L929 cells treated with BaONPs showed higher DNA damage than control. Notable damage of DNA was observed in BaONPs (300 μg/mL)-treated L929 cells (Figure 7).

Discussion

This study suggested that BaONPs are slightly lethal to L929 cells, and this cell toxicity turns significant with dose and duration rise. Murdock et al\(^1\) reported that nanoparticles can induce adverse effects due to their small size and unique physiochemical characters. Nel et al\(^1\) reported that the key parameters of nanoparticles are structure, shape, purity, and hydrodynamical agglomeration, which control the biological activity of BaONPs. Specific physical–chemical characters of BaONPs are due to surface area to volume ratio. Nanoparticles have a high percentage of atoms on their surface compared to macroparticles, studying the genotoxic effect of BaONPs, we had characterized their size and zeta potential of the BaONPs by FETEM and DLS, respectively. However, the hydrodynamic dimension obtained by DLS was greater than quantified size of FETEM.

L929 cells are commonly used to test the cytotoxic potential of nanoparticles. BaONPs induced caspase-3 activity and ROS in L929 cells. In this study, we observed that BaONPs increased cytotoxicity and apoptosis in L929 cells. Nel et al\(^1\) reported that the toxic effect of nanoparticles was due to oxidative stress. Oxidative stress provokes a long range of cellular and physiological events, eg, DNA damage, inflammation, apoptosis, and stress.\(^1\)

Experimental evidence has shown that barium sulfate nanoparticles cause lung injury and inflammation in rats.\(^1\) During oxidative stress, the antioxidant mechanisms are
unable to counter ROS production. ROS, e.g., $O_2^-$, $H_2O_2$, and OH radicals, have a higher chemical activity than $O_2$. Oberdorster et al.\textsuperscript{19} reported that the reduction of GSH and stimulation of LPO and ROS induce oxidative damage of cell components. ROS, lipid peroxidation, and SOD activity was increased, whereas glutathione level was less in BaONPs-treated L929 cells. Glutathione was implicated in the first-line cellular defense mechanism against oxidative injury in cells.\textsuperscript{20}

In this study, we found a correlation between the production of ROS and glutathione decline, suggesting that higher generation of ROS induces proliferated lipid peroxidation accompanying decline in GSH. Kang et al.\textsuperscript{21} reported that oxidative stress and lipid peroxidation lead to apoptosis and DNA damage. During apoptosis, a chain of biochemical reactions occurs, inducing an alteration of L929 cell shape, and ultimately resulting in cell death.

Kroemer et al.\textsuperscript{22} reported some biomarkers that characterize apoptosis, such as DNA fragmentation, cell membrane blebbing, condensation, shrinking, and disintegration of cell organelles. In this study, Hoechst 33342 staining of BaONPs-treated L929 cells gave rise to nuclear fragmentation and condensation. Chen et al.\textsuperscript{23} demonstrated that nanomaterials interact with DNA due to their small size and shape. Nanoparticles can have an indirect effect on DNA production of ROS.\textsuperscript{24} The DNA damage can either induce
Figure 4 (A) Levels of lipid peroxidation; (B) glutathione; (C) SOD; and (D) catalase in L929 cells after exposure to BaONPs for 48 hours.

Notes: Data represent the average ± SE of three tests. Statistical differences with respect to the controls are shown (*P<0.05).

Abbreviations: BaONPs, barium oxide nanoparticles; GSH, glutathione; SOD, superoxide dismutase; SE, standard error.
Figure 5 Chromatin condensation and stimulation of caspase-3 activity in L929 cells after exposure to BaONPs (A). Control (B) at 150 μg/mL of BaONPs for 48 hours of BaONPs. (C) Caspase-3 activity. (D) Effect of NAC (5 mM) on BaONPs (150 μg/mL) induced caspase-3 activity in L929 cells over 24 hours.

Notes: Data represent the average ± SE of three tests. Statistical differences with respect to the controls are shown (*P < 0.05). Scale bar (——) 50 μm. The white arrows indicate the condensed chromosomes.

Abbreviations: BaONPs, barium oxide nanoparticles; NAC, N-acetylcysteine; SE, standard error.

Figure 6 (A–C) Flow cytometric analysis of Annexin V–FITC/PI stained cells. (D) Percentage of Annexin V–FITC/PI-positive cells in control and L929 cells exposed to BaONPs (150 μg/mL) for 24 and 48 hours.

Notes: Each value represents the average ± SE of two experiments. Statistical differences with respect to the controls are shown (*P < 0.05).

Abbreviations: BaONPs, barium oxide nanoparticles; FITC, fluorescein isothiocyanate; PI, propidium iodide; SE, standard error.
cell death or carcinogenesis, consequently disrupting normal cell functions.

We found that DNA damaging effects of BaONPs in L929 cells by using comet assay, which is a technique used to identify double- and single-strand breaks and alkali labile sites even at low levels of DNA damage.25 Martinez et al26 reported that ROS reacts with DNA molecules, inducing damage to both bases (pyrimidine and purine) and the DNA backbone. This seems to be the underlying mechanism of BaONPs-mediated cell toxicity and DNA damage in L929 cells.

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Disclosure

The authors report no conflicts of interest in this work.

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