

Aneuploidogenic effects and DNA oxidation induced in vitro by differently sized gold nanoparticles

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Abstract: Gold nanoparticles (Au NPs) are used in many fields, including biomedical applications; however, no conclusive information on their potential cytotoxicity and genotoxicity mechanisms is available. For this reason, experiments in human primary lymphocytes and murine macrophages (Raw264.7) were performed exposing cells to spherical citrate-capped Au NPs with two different nominal diameters (5 nm and 15 nm). The proliferative activity, mitotic, apoptotic, and necrotic markers, as well as chromosomal damage were assessed by the cytokinesis-block micronucleus cytome assay. Fluorescence in situ hybridization with human and murine pancentromeric probes was applied to distinguish between clastogenic and aneuploidogenic effects. Our results indicate that 5 nm and 15 nm Au NPs are able to inhibit cell proliferation by apoptosis and to induce chromosomal damage, in particular chromosome mis-segregation. DNA strand breaks were detected by comet assay, and the modified protocol using endonuclease-III and formamidopyrimidine-DNA glycosylase restriction enzymes showed that pyrimidines and purines were oxidatively damaged by Au NPs. Moreover, we show a size-independent correlation between the cytotoxicity of Au NPs and their tested mass concentration or absolute number, and genotoxic effects which were more severe for Au NP 15 nm compared to Au NP 5 nm. Results indicate that apoptosis, aneuploidy, and DNA oxidation play a pivotal role in the cytotoxicity and genotoxicity exerted by Au NPs in our cell models.

Keywords: Au nanoparticles, cytotoxicity, aneuploidy, oxidative DNA damage, micronuclei, particle size

Introduction

Evidence has emerged for the important role of physicochemical properties in controlling the cellular uptake and toxicity of nanoparticles in a biological system. In vitro uptake of gold nanoparticles (Au NPs) depends on the particle physical dimensions, suggesting that size and shape can mediate particle-cell interactions, receptor recycling rates and exocytosis.¹ Au NPs enter the cells via a clathrin-mediated endocytosis and are exocytosed by a size-dependent relationship.² It is also known that 10–100 nm gold particles can enter or exit cells via wrapping, even in the absence of clathrin- or caveolin-coated vesicles.³ However, the toxicity of Au NPs, as well as of others such as silver NPs, might not correlate with the level of uptake but cell-type and the size can play a role.^{4,5} In fact, 20 nm citrated Au NPs resulted as more toxic than 14 nm ones in ovarian hamster's cells but not in bronchial and kidney derived cell cultures.⁴ However, even if it is recognized that the cell type plays a role in the uptake of Au NPs, the size is not involved: differently sized Au NPs displayed a similar uptake in epithelial (A549 and NCIH441) and in endothelial (HDMEC and hCMEC) cells, but endothelial cells were shown to internalize a higher amount of Au NPs than epithelial ones.^{6,7} Despite these

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indications, it remains unclear how the physicochemical properties of NPs change when the exposure occurs in a biological system. In fact, it is known that different serum proteins form protein coronas, which significantly modulate NPs behavior.⁸ The cellular environment contains a variety of proteins rapidly adsorbing onto the surface of NPs and forming a protein corona which defines the NPs biological identity, altering size, aggregation state, surface chemistry, uptake and ion release.⁹ The alteration of these physicochemical properties can result in increasing or protecting from NPs toxicity. Since Au NPs are used in medical applications, such as photothermal therapy,¹⁰ and are often administered intravenously, the adsorption of blood proteins onto their surface is one of the most relevant and studied parameters. Blood is a complex human-derived proteome and contains other tissue subsets of proteins,¹¹ which potentially interact with NPs changing their physicochemical characteristics and their biological identity. For example, NPs size and surface chemistry can mediate plasma and serum protein adsorption to Au NPs and their biological responses and/or uptake by macrophages.^{12,13} From the different proteins accumulation patterns, observed as a result of differently sized and surface-charged Au NPs administered intravenously to rats, Hirn et al concluded that the alterations of the Au NPs surface area, determined by the size of the particles, are the main mechanism determining Au NPs accumulation in various organs and tissues.¹⁴ Size-dependent tissue distribution of Au NPs was observed upon intravenous administration in mice and size-dependent alterations occurred in vivo in the hepatic tissue of exposed rats.^{15,16} In contrast, monodispersed Au NPs did not exert a higher acute effect compared to their agglomerates, when using pulmonary inflammation as a marker for toxicity after intratracheal instillation in rats.¹⁷ Furthermore, the model organism *Drosophila melanogaster* showed DNA fragmentation and significant concentration-dependent and size-independent modulation of expression levels of the genes involved in stress stimuli (*hsp70* and *hsp83*), DNA damage (*p53*) and apoptosis pathway (*Dark*, *Dronc*, and *Dredd*) upon ingestion of Au NPs.¹⁸

To study the toxic mechanisms of Au NPs, we assessed the cytotoxic and genotoxic responses induced by Au NPs in human peripheral blood lymphocytes (PBL) and murine macrophages Raw264.7, by differentiating clastogenic and aneuploidogenic effects, DNA strand breaks and oxidative DNA damage. Cell viability was determined by methylthiazol tetrazolium (MTT) assay, and the proliferative activity, cytotoxicity, mitotic, apoptotic and necrotic markers, as well as the chromosomal damage induced by Au NPs, were

assessed by the cytokinesis-block micronucleus cytome assay. To distinguish between clastogenic and aneuploidogenic effects, we performed the fluorescence in situ hybridization (FISH) with human and murine pancentromeric probes. Single cell DNA strand breaks were detected by comet assay, and the oxidative DNA damage induced by Au NPs was investigated applying the modified comet protocol with endonuclease-III (EndoIII) and formamidopyrimidine-DNA glycosylase (Fpg) enzymes. In addition, the data were further analyzed taking into account the mass concentrations and the theoretically calculated number of Au NPs used. To verify whether size is the factor that regulates the toxic responses of Au NPs, we also exposed Raw264.7 cells to Au NPs with different nominal diameters (5 nm and 15 nm) at increasing exposure times and increasing absolute numbers of Au NPs.

Material and methods

Au NPs synthesis and characterization

Au NPs of two different sizes (5 nm and 15 nm) were prepared and characterized at the Joint Research Centre of the European Commission (Ispra, Italy) as a suspension of 98.5 mg/L (total Au content) in citrate buffer, as previously reported.^{19,20} Particles were characterized by three different techniques, dynamic light scattering, centrifugal particle sedimentation (CPS), and scanning transmission electron microscopy (STEM), and the zeta potential was determined by Zetasizer ZS-Nano (Malvern Instruments, Worcestershire, UK). To assess if NPs aggregate, their size distribution in the as-synthesized state and following dispersion in complete culture medium (9.85 µg/mL) was measured using the DC24000UHR centrifuge (CPS Instruments, Oosterhout, the Netherlands). The size and shape of the Au NPs was also evaluated using STEM images obtained using a FEI Nova nanolab microscope (FEI, Eindhoven, the Netherlands). The inductively coupled plasma-mass spectrometry technique (ICP-MS, 7700 series, Agilent Technologies Inc., Santa Clara, CA, USA) was used to assess the presence of dissolved ions in the Au NPs suspensions after removal of all solid particulate material by ultrafiltration (10 kDa Amicon filters, 20 minutes, room temperature, 1,500× g).

Cell culture

We drew 4–6 mL of blood from two healthy volunteers by venipuncture in Li-heparin vials, according to standard procedure. The donors were chosen according to the following criteria: young age, nonsmokers, without pharmacological treatments for at least 3 weeks before donation and without

any radiological examination performed within the previous 3 months. The PBL were cultured as previously described.²¹

Raw264.7 cells were purchased from the Istituto Zooprofilattico of Brescia (Italy). The cells were cultured in MEM (minimal essential medium) (CELBIO, Milan, Italy) supplemented with 10% fetal bovine serum (CELBIO), 1% penicillin/streptomycin (CELBIO), and 1% L-glutamine (CELBIO). The cell cultures were maintained in a humidified atmosphere (5% CO₂) at 37°C.

Au NP exposure

After 24 hours of seeding, the cell cultures were treated either with different mass concentrations (0.1, 1, 10, and 100 µg/mL) or with increasing absolute numbers (1×10⁹, 1×10¹⁰, 1×10¹¹) of 5 nm and 15 nm Au NPs. The tested samples were prepared by diluting suitable aliquots of Au NPs stock solutions in complete culture medium. The number of Au NPs within each mass concentration dose and the tested number of Au NPs were determined by the following formula:

$$N = \frac{V_{NP}}{v_{NP}} = \frac{6 \cdot M \cdot m \cdot V_{sample}}{\rho \cdot \pi \cdot d^3} = \frac{6 \cdot V_{sample}}{\rho \cdot \pi} \cdot \frac{M \cdot m}{d^3} \quad (1)$$

where, v_{NP} = volume of a nanoparticle = $4/3 \cdot r^3 \cdot \pi = 1/6 \cdot d^3 \cdot \pi$ (r = nanoparticle radius; d = nanoparticle diameter), V_{NP} = total volume of nanoparticles in the sample = $(M_{NP}/\rho) = (M \cdot m \cdot V_{sample})/\rho$ and M_{NP} = mass of nanoparticles in the sample volume = $V_{sample} \cdot M \cdot m$ (ρ = specific mass or density expressed as g/cm³; m = molar mass expressed as g/mol; M = molarity; V_{sample} = sample volume). For Au NPs nanoparticles: $m=197$ g/mol and $\rho=19.3$ g/cm³.

Cell viability

Cytotoxicity was assessed by MTT assay. Subconfluent Raw264.7 cells were grown in 96-well plates and exposed to Au NPs suspensions for 2 hours and 24 hours. PBL were treated at the same conditions of adherent Raw264.7 cells. PBL were collected in sterile tubes and resuspended in 0.5 mL complete MEM; erythrocyte lysis buffer (155 mM NH₄Cl, 10 mM KHCO₃, 1 mM Na₂EDTA, pH 7.4; 3.5 mL/tube) was added, and after 90 seconds cells were collected by centrifuging (3,000 rpm, 5 minutes). PBL were washed in 3.5 mL of complete medium and then pelleted (3,000 rpm, 5 minutes). The pellet was resuspended in 3.5 mL complete MEM and 100 µL of the cell suspension was pipetted in each well of a 96-well plate. Hydrogen peroxide (10 µM H₂O₂) was used as positive control.

After 2 hours and 24 hours of exposure of Raw264.7 and PBL to Au NPs, 10 µL of a 5 mg/mL MTT solution was added to each well (nine wells for each condition). After 3 hours at 37°C, Raw264.7 medium was replaced by 100 µL of dimethyl sulfoxide (DMSO) and mixed thoroughly to dissolve the formazan crystals; to PBL in suspension, 100 µL of DMSO was added without discarding the medium. To limit potential interactions due to the presence of residual NPs that could interfere with the assay, blanks without cells were used to detect the absorption of Au NPs themselves. Absorbance was measured at 570 nm (630 nm background) and cell viability was determined as the percentage of the blanks subtracted to the corresponding percentage of the exposed cells.

Cytokinesis-block micronucleus cytome assay

The cytokinesis-block micronucleus cytome (CBMN Cyt) assay after whole blood treatments was performed according to the procedure described by Migliore et al,²¹ while the CBMN Cyt assay on Raw264.7 was performed according to Migliore et al.²² The exposure to Au NPs lasted for 48 hours. Mitomycin C (0.1 µg/mL; MMC, Kyowa Hakko Kogyo Co, Chiyoda, Tokyo, Japan) was used as positive control. As described by Fenech, 500 cells were scored to evaluate the percentage of mono-, bi- and multinucleated cells, and the cytokinesis block proliferation index (CBPI) was calculated as an index of cytotoxicity by comparing the distribution of mono-, bi-, and multinucleated cells in the treated and control cells.²³ The CBPI indicates the average number of cell cycles per cell during the period of exposure to cytoB, and is used to calculate cell proliferation. The number of apoptotic, necrotic, and mitotic cells per 500 cells was also evaluated. The genotoxic potential of Au NPs was evaluated by scoring the binucleated/micronucleated cells frequency as number of 1,000 binucleated cells containing one or more micronuclei (MN). Finally, on these 1,000 binucleated cells other parameters such as nucleoplasmic bridges (NPB), a biomarker of DNA misrepair and/or telomere end-fusions, and nuclear buds (NBUD), a biomarker of elimination of amplified DNA and/or DNA repair complexes, were also scored.

For PBL, data were reported as mean of four independent experiments (from two donors) ± standard error of the mean (SEM); for Raw264.7 cells, results were plotted as mean of three independent experiments (two replicates each) ± SEM.

FISH

Cells to perform FISH were prepared and spotted onto CBMN Cyt slides as previously described.²⁴ Samples

exposed for 48 hours to 0.1 $\mu\text{g/mL}$, 1 $\mu\text{g/mL}$, and 10 $\mu\text{g/mL}$ of Au NPs, and to positive (0.1 $\mu\text{g/mL}$ MMC, clastogen) and negative controls, were classified as centromere positive or centromere negative based on the presence or on the absence of centromeres in MN. Data were reported as mean of four independent experiments (from two donors) \pm SEM; for Raw264.7 cells, results were plotted as mean of three independent experiments (two replicates each) \pm SEM.

Comet assay and oxidative DNA damage

DNA damage was measured using the alkaline comet assay as described by Di Bucchianico et al.²⁴ The percentage of total DNA fluorescence in tail in a total of 100 randomly selected cells per sample (two replicates, each with 50 cells/slide) was used as a measure of the amount of primary DNA damage. Ten micromolar H_2O_2 was used as positive control. Three independent experiments were performed for both 2- and 24-hour treatments. Analysis was carried out by using a Comet Image Analysis System, version 5.5 (Kinetic Imaging, Nottingham, UK).

To determine the presence of oxidized pyrimidine and purine bases, EndoIII and Fpg were used as previously described.²⁴ Three independent experiments were performed for both 2- and 24-hour treatments. To determine the number of enzyme-sensitive sites, the difference between the value of the percent DNA fluorescence in tail obtained after digestion with each enzyme and with the buffer only was calculated.

Results

Characterization of Au NPs

The characterization of Au NPs was previously reported by Coradeghini et al and is summarized in Table 1.²⁰ UV-vis spectrophotometry of Au NPs showed different absorption peaks at 508 nm and 520 nm wavelengths, consistent with their different diameters. A good agreement was found between the results of electron microscopy and CPS measurements. The CPS analysis was also used to measure

the size distribution of 5 nm and 15 nm Au NPs in culture medium. Results showed that $>70\%$ of the 5 nm and 15 nm Au NPs are monodispersed, even after being resuspended in complete MEM for 72 hours at 37°C . The concentration of ions released from the Au NPs suspension, measured by ICP-MS up to 72 hours in complete medium, was for both sizes of Au NPs below the limit of detection of the instrument (<1 ppb).

Cell viability

Cell viability was evaluated in PBL and Raw264.7 by MTT assay, a standard in vitro test already optimized to study NP toxicity.²⁵ In PBL, 100 $\mu\text{g/mL}$ of 5 nm Au NPs showed more severe cytotoxicity after 2 hours of exposure (Figure 1), while 15 nm Au NPs exerted no signs of cytotoxicity. The 24-hour treatments showed a significant decrease in cell viability for both NPs, and in the presence of 15 nm Au NPs the effect was dose dependent.

In Raw264.7 cells (Figure 1), the cell viability decreased in a dose-dependent manner after 2 hours of exposure to 5 nm Au NPs, and the same was observed after 24 hours of exposure to both 5 nm and 15 nm Au NPs. Interestingly, at 2 hours of exposure, the smallest NPs showed higher cytotoxicity in both macrophages and lymphocytes.

Cytotoxicity and cytostasis

PBL proliferation, measured as CBPI, decreased in a dose-dependent manner following exposure to Au NPs (Table 2); 5 nm Au NPs showed a more effective antiproliferative activity than 15 nm Au NPs, which in contrast was the most effective at the highest tested dose (100 $\mu\text{g/mL}$). In PBL, significant antimitotic and apoptotic activities were observed after exposure to both Au NPs, but the most severe apoptotic effect was seen after exposure to 10 $\mu\text{g/mL}$ and 100 $\mu\text{g/mL}$ 15 nm Au NPs. The necrotic index increased for both Au NPs in a dose-dependent manner; the exposure to 5 nm Au NPs showed higher necrotic effect than

Table 1 Au NP characterization

Au NP nominal size	STEM	DLS in native solution			CPS in native solution			CPS in MEM		
	Mean \pm SD (nm)	Mean \pm SD (nm)	PDI	Z-pot (mV)	Mean (nm)	Half-width (nm)	PDI	Mean (nm)	Half-width (nm)	PDI
5 nm	3.3 \pm 1.7	7.1 \pm 2.1	0.097	-26 \pm 11	6.6	2.2	1.37	3.4	1.5	5
15 nm	12.0 \pm 1.0	17.0 \pm 4.7	0.028	-30 \pm 12	12.9	1.7	1.04	7.4	1.3	2.3

Notes: Mean size distribution \pm SD of 5 nm and 15 nm Au NPs measured in native solution by STEM, DLS, and CPS. PDI and Z-potential values are also included. Values are expressed as mean of three measurements for DLS and CPS, and for STEM analysis of at least 300 NPs. The mean size distribution of 5 nm and 15 nm Au NPs obtained by CPS was measured in MEM after 72 hours of incubation at 37°C . Z-potential measurements of the as-synthesized Au NPs were performed at the physiological pH range of 6.5–7.5.

Abbreviations: Au NP, gold nanoparticle; CPS, centrifugal particle sedimentation; DLS, dynamic light scattering; MEM, minimal essential medium; PDI, polydispersity index; SD, standard deviation; STEM, scanning transmission electron microscopy; Z-pot, Z-potential.

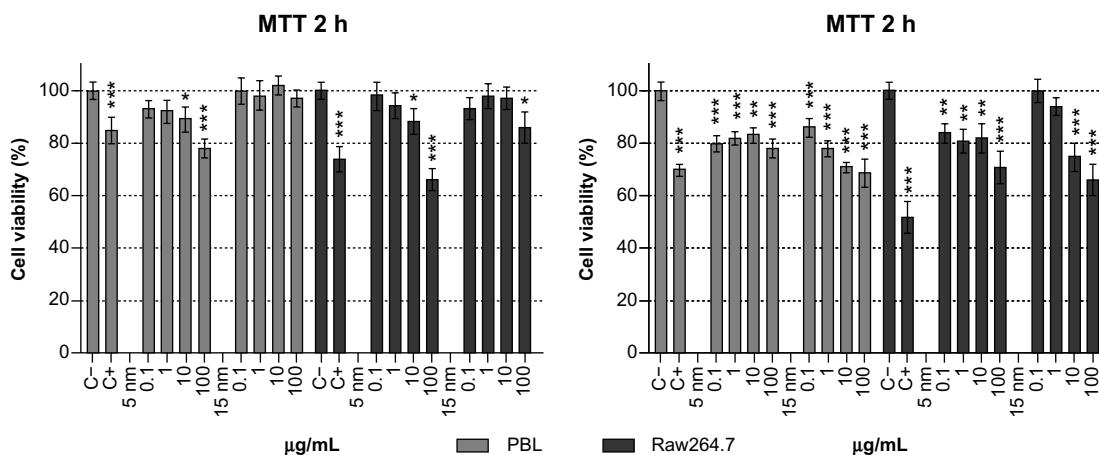


Figure 1 Cell viability, evaluated by MTT assay, in PBL and Raw264.7 after 2 hours and 24 hours of exposure to 5 nm and 15 nm Au NPs. **Notes:** PBL following 24 hours treatment with Au NP 15 nm showed a dose-effect relationship ($P=0.01$). Raw264.7 following 2 hours treatment with 5 nm Au NPs showed a dose-effect relationship ($P=0.04$) as well as after 24 hours treatment to 5 nm and 15 nm Au NPs ($P=0.03$ and $P=0.016$, respectively). Data show the percent of viable cells normalized to untreated control (\pm SEM, $n=9$). Statistically significant differences from the C- were determined by Student's *t*-test (* $P<0.05$; ** $P<0.01$; *** $P<0.001$). **Abbreviations:** Au NP, gold nanoparticle; C+, positive control (10 μ M hydrogen peroxide); C-, negative (untreated) control; h, hours; MTT, methylthiazol tetrazolium; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

15 nm Au NPs, but 15 nm Au NPs was the most effective at 100 μ g/mL.

In Raw264.7 cells, proliferation decreased after exposure to both Au NPs, and the effect was dose dependent in the presence of 5 nm Au NPs. The clearest low-dose effect was seen following exposure to 5 nm Au NPs. Although no significant dose-response was observed, the highest dose (100 μ g/mL) of 15 nm Au NPs resulted in illegible slides because of the lack of morphologically undamaged

cells (Table 2). Concentration-dependent decrease in the mitotic index was noticed, and the inhibition of cell division occurred more significantly in macrophages exposed to 15 nm Au NPs. The apoptotic index was significantly increased at the highest 5 nm and 15 nm Au NPs concentrations, and in a dose-dependent manner with 15 nm Au NPs. While 15 nm Au NPs induced a dose-dependent effect, 5 nm Au NPs increased the necrotic index at all tested concentrations.

Table 2 Cytostasis and cytotoxicity evaluated by CBMN Cyt assay

	CBPI		Mitotic index		Apoptotic index		Necrotic index	
	PBL	Raw264.7	PBL	Raw264.7	PBL	Raw264.7	PBL	Raw264.7
C-	1.88±0.02	1.85±0.04	6.5±0.1	5.6±0.2	0.7±0.1	1.7±0.2	0.2±0.2	0.9±0.3
C+	1.17±0.13***	1.41±0.06***	0.3±0.2***	2.3±0.6***	4.7±0.3***	4.6±0.5***	2.5±0.4***	4.0±0.4
Au NP 5 nm								
0.1 μ g/mL	1.55±0.02***	1.62±0.06**	3.9±0.2***	4.0±0.4***	1.2±0.2*	2.0±0.5	1.6±0.3***	3.4±0.4***
1 μ g/mL	1.54±0.05***	1.62±0.05**	4.6±0.6**	3.9±0.4***	1.9±0.5**	1.2±0.3	2.4±0.3***	2.1±0.5**
10 μ g/mL	1.47±0.02***	1.57±0.05**	4.4±0.3***	3.6±0.3***	1.6±0.2**	1.8±0.5	2.6±0.2***	2.9±0.2***
100 μ g/mL	1.36±0.05***	1.51±0.07***	2.8±0.4***	3.1±0.4***	1.8±0.4*	2.6±0.2***	3.0±0.6***	4.3±0.6***
Au NP 15 nm								
0.1 μ g/mL	1.70±0.03*	1.68±0.04**	4.9±0.5**	3.7±0.3***	1.5±0.1**	1.7±0.1	0.6±0.2*	1.4±0.2*
1 μ g/mL	1.57±0.02***	1.61±0.01***	4.0±0.6***	2.8±0.2***	1.9±0.2**	2.3±0.3*	0.9±0.1**	1.7±0.3**
10 μ g/mL	1.56±0.05***	1.60±0.04***	3.5±0.3***	2.5±0.2***	3.5±0.3***	2.8±0.4**	1.6±0.2***	2.5±0.3***
100 μ g/mL	1.34±0.04***	Toxic	1.1±0.6***	Toxic	9.1±0.2***	Toxic	3.7±0.6***	Toxic

Notes: CBPI and mitotic (cytostasis), apoptotic, and necrotic (cytotoxicity) indices scored by CBMN Cyt assay. CBPI of PBL exposed to 5 nm and 15 nm Au NPs decreased with a dose-dependent relationship ($P=0.027$ and $P=0.02$, respectively) as well as for Raw264.7 exposed to 5 nm Au NPs ($P=0.04$). Dose-dependent decrease in the mitotic index was observed in PBL exposed to 15 nm Au NPs ($P=0.029$) and in Raw264.7 exposed to 5 nm and 15 nm Au NPs ($P=0.033$ and $P=0.05$, respectively). Both PBL and Raw264.7 exposed to 15 nm Au NPs showed dose-dependent apoptotic index increase ($P=0.049$ and $P=0.05$, respectively); the necrotic index induced a dose-dependent effect in PBL exposed to 5 nm and 15 nm Au NPs ($P=0.016$ and $P=0.03$, respectively) and in Raw264.7 exposed to 15 nm Au NPs ($P=0.017$). 100 μ g/mL 15 nm Au NPs resulted in illegible slides. Statistically significant differences from the control were determined by the Student's *t*-test (* $P<0.05$; ** $P<0.01$; *** $P<0.001$). Data represent the mean \pm SEM. PBL: $n=4$; Raw264.7: $n=6$.

Abbreviations: Au NP, gold nanoparticle; C+, positive control (0.1 μ g/mL MMC); C-, negative (untreated) control; CBMN Cyt, cytokinesis-block micronucleus cytome; CBPI, cytokinesis block proliferation index; MMC, mitomycin C; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

Chromosomal damage

The results of MN, NPB, and NBUD frequencies are summarized in Table 3. The frequency of MN in PBL increased dose-dependently up to 3.8-fold and 6.9-fold after exposure to 5 nm and 15 nm Au NPs, respectively, while the number of MN increased up to 3.2-fold and 11.9-fold when Raw264.7 were treated with 5 nm and 15 nm Au NPs; 15 nm Au NPs showed a dose-dependent relationship in Raw264.7. In both cell types, a more severe MN induction was observed after treatment with 15 nm Au NPs. Chromosome rearrangements, detected as NPB formation, were more pronounced in PBL than in Raw264.7. Furthermore, in PBL following 5 nm Au NPs exposure and for both sets of Au NPs delivered to Raw264.7, the ratio NPB/MN showed that the lowest dose (0.1 µg/mL) exhibited higher chromosome breakage than 100 µg/mL. The frequencies of NBUD, biomarker of amplified DNA elimination, increased more in lymphocytes than in macrophages exposed to 5 nm Au NPs, while after exposure to 15 nm Au NPs the macrophages were more sensitive than lymphocytes.

Taken together, these data indicate that 15 nm Au NPs induced more significant chromosomal damage than 5 nm Au NPs.

FISH analysis

The exposure to Au NPs induced significant increase in centromere-positive MN formation in both cell types, indicating the induction of chromosome segregation anomalies rather than clastogenic effects (Figure 2). Both the Au NPs

induced higher chromosome loss in PBL than in Raw264.7; however, in PBL the lowest dose of 5 nm Au NPs showed minor, but still significant, aneuploidogenic effects. Raw264.7 cells showed no significant aneuploidogenic events at the lowest 5 nm Au NPs dose, while 15 nm Au NPs induced linear dose-dependent effects.

DNA strand breaks

Figure 3 shows the data obtained evaluating the Au NPs-induced DNA damage by the comet assay. Significant increase in DNA migration was detected after 2 hours and 24 hours of exposure in both cell types, and the increase was dose dependent with the exception of PBL exposed for 24 hours to 5 nm Au NPs.

PBL were more sensitive to 5 nm and 15 nm Au NPs than Raw264.7: at 100 µg/mL, in fact, a 3.6-fold and 2.7-fold increase in DNA damage was observed following 24 hours of exposure to 5 nm Au NPs in PBL and Raw264.7, respectively. Accordingly, when exposed to 15 nm Au NPs for 24 hours, PBL were more sensitive than Raw264.7, presenting respectively up to 4.7-fold and 2.7-fold increases in DNA strand breaks. Interestingly, at 2 hours of exposure, PBL were more sensitive to 5 nm Au NPs than 15 nm Au NPs, while after 24 hours of exposure to the highest doses, the opposite was observed.

Oxidative DNA damage

By using the repair enzymes EndoIII and Fpg, the oxidative DNA damage was investigated at 2 hours and

Table 3 Genotoxicity evaluated by CBMN Cyt assay in PBL and Raw264.7 cells exposed to increasing mass concentrations of Au NPs

	MN		NPB		NBUD	
	PBL	Raw264.7	PBL	Raw264.7	PBL	Raw264.7
C-	3.25±0.75	2.3±0.5	0.5±0.5	1.2±0.4	0±0	0.5±0.5
C+	43.3±7.2***	32.4±5.6***	14.4±0.8***	13.4±1.0**	2.8±0.5***	7.0±1.5***
Au NP 5 nm						
0.1 µg/mL	5.0±1.4	5.1±1.1	2.2±0.5**	1.8±0.5	1.5±0.5**	1.3±0.6
1 µg/mL	7.0±2.0	4.5±1.2	2.2±0.2**	1.4±0.8	4.3±0.3***	1.0±0.8
10 µg/mL	8.5±0.35*	5.4±1.4	2.0±0.4**	1.4±0.8	2.5±0.4***	2.0±0.8
100 µg/mL	12.5±0.35**	7.4±1.1**	3.2±0.6**	1.8±0.4	6.7±0.9***	1.3±0.4
Au NP 15 nm						
0.1 µg/mL	9.2±1.2*	17.5±1.5**	2.0±0.5**	4.2±1.2*	0.5±0.5	5.5±0.5***
1 µg/mL	12.5±0.5**	25.0±2.0***	2.8±0.7**	4.8±0.5*	2.5±0.5***	7.0±1.0***
10 µg/mL	14.3±1.2**	27.5±2.5***	2.8±1.0**	4.8±0.8*	5.5±0.7***	7.5±1.5***
100 µg/mL	22.4±2.3***	Toxic	3.2±1.5**	Toxic	12±1***	Toxic

Notes: MN, NPB, and NBUD frequencies scored by CBMN Cyt assay after exposure of PBL and Raw264.7 to increasing mass concentrations of Au NPs (0.1–100 µg/mL). In PBL, 5 nm and 15 nm Au NPs induced dose-dependent effect in MN ($P=0.0014$ and $P=0.03$, respectively), NPB ($P=0.012$ and $P=0.033$; respectively), and NBUD frequency ($P=0.05$ and $P=0.02$, respectively). MN frequency of Raw264.7 exposed to 15 nm Au NPs showed a dose-dependent relationship ($P=0.05$). Statistically significant differences from the control were determined by the Student's *t*-test (* $P<0.05$; ** $P<0.01$; *** $P<0.001$). Data represent the mean ± SEM (PBL: $n=4$; Raw264.7: $n=6$).

Abbreviations: Au NP, gold nanoparticle; C+, positive control (0.1 µg/mL mitomycin C); C-, negative (untreated) control; CBMN Cyt, cytokinesis-block micronucleus cytochrome; MN, micronuclei; NBUD, nuclear buds; NPB, nucleoplasmic bridges; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

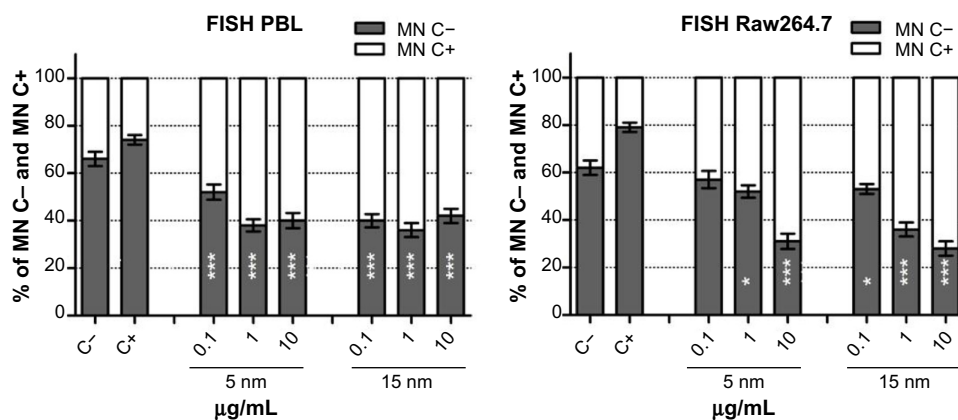


Figure 2 FISH analysis in PBL and Raw264.7 after exposure to 5 nm and 15 nm Au NPs.

Notes: Using pancentromeric probes applied to the CBMN Cyt slides, the centromeric spots were analyzed and the exposure of Raw264.7 to 15 nm Au NPs showed dose-dependent ($P=0.01$) aneuploidogenic events. Data are shown as mean \pm SEM (PBL: $n=4$; Raw264.7: $n=6$). Student's *t*-test: * $P<0.05$; ** $P<0.01$; *** $P<0.001$.

Abbreviations: Au NP, gold nanoparticle; C+, positive control (MMC 0.1 $\mu\text{g/mL}$); C-, negative (untreated) control; CBMN Cyt, cytokinesis-block micronucleus cytome; FISH, fluorescence in situ hybridization; h, hours; MN C+, micronuclei centromere +ve; MN C-, micronuclei centromere -ve; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

24 hours exposure, and oxidized pyrimidines were detected in both cell systems by the presence of EndoIII sensitive DNA sites (Figure 4). Compared to PBL, at 2 hours of exposure Raw264.7 were particularly susceptible, showing a dose-dependent pyrimidines oxidation in the presence of 15 nm Au NPs; interestingly, the 5 nm Au NPs treatments were the less damaging to Raw264.7 and most harmful to PBL. After 24 hours of treatment, in both cell types the two Au NPs induced oxidized pyrimidines in a dose-dependent way; Raw264.7 were more sensitive than PBL (4.2-fold and 7.8-fold increases in Raw264.7 and 2.9-fold and 3.3-fold increases in PBL

at the highest dose following 5 nm and 15 nm Au NPs, respectively).

Raw264.7 were more sensitive in detecting oxidized purines by Fpg restriction enzyme: 5 nm Au NPs at 2 hours of exposure induced a 4.8-fold increase in Raw264.7 and 3.9 in PBL. Similarly, after 24 hours of exposure, in Raw264.7 the purines oxidation increased 4.1-fold, while in PBL the increase was up to 1.5-fold. At 2 hours of exposure to 15 nm Au NPs, purines oxidation was up to eight times higher in Raw264.7 and up to four times higher in PBL compared to negative control, while at 24 hours of exposure the oxidative damage was up to seven times higher in Raw264.7 and 2.3 times in PBL.

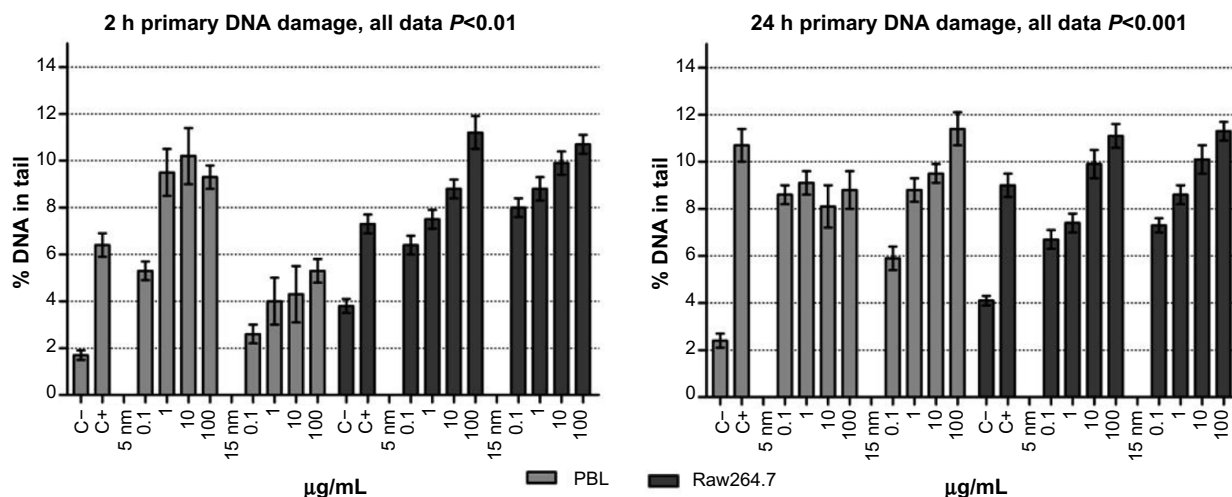


Figure 3 Comet assay to evaluate primary DNA damage/breakage effects in PBL and Raw264.7.

Notes: The 2-hour treatments with 5 nm and 15 nm Au NPs showed a dose-effect relationship in PBL ($P=0.05$ and $P=0.002$) and in Raw264.7 ($P=0.0014$ and $P=0.03$, respectively). The 24-hour exposure to 5 nm and 15 nm Au NPs showed a dose-effect relationship in Raw264.7 ($P=0.0016$ and $P=0.004$, respectively), but only 15 nm Au NPs in PBL ($P=0.006$). Data are shown as mean \pm SEM (PBL: $n=4$; Raw264.7: $n=6$) of percent DNA in tail. Student's *t*-test: 2-hour data, $P<0.01$; 24-hour data, $P<0.001$.

Abbreviations: Au NP, gold nanoparticle; C+, positive control (10 μM hydrogen peroxide); C-, negative (untreated) control; DNA, deoxyribonucleic acid; h, hours; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

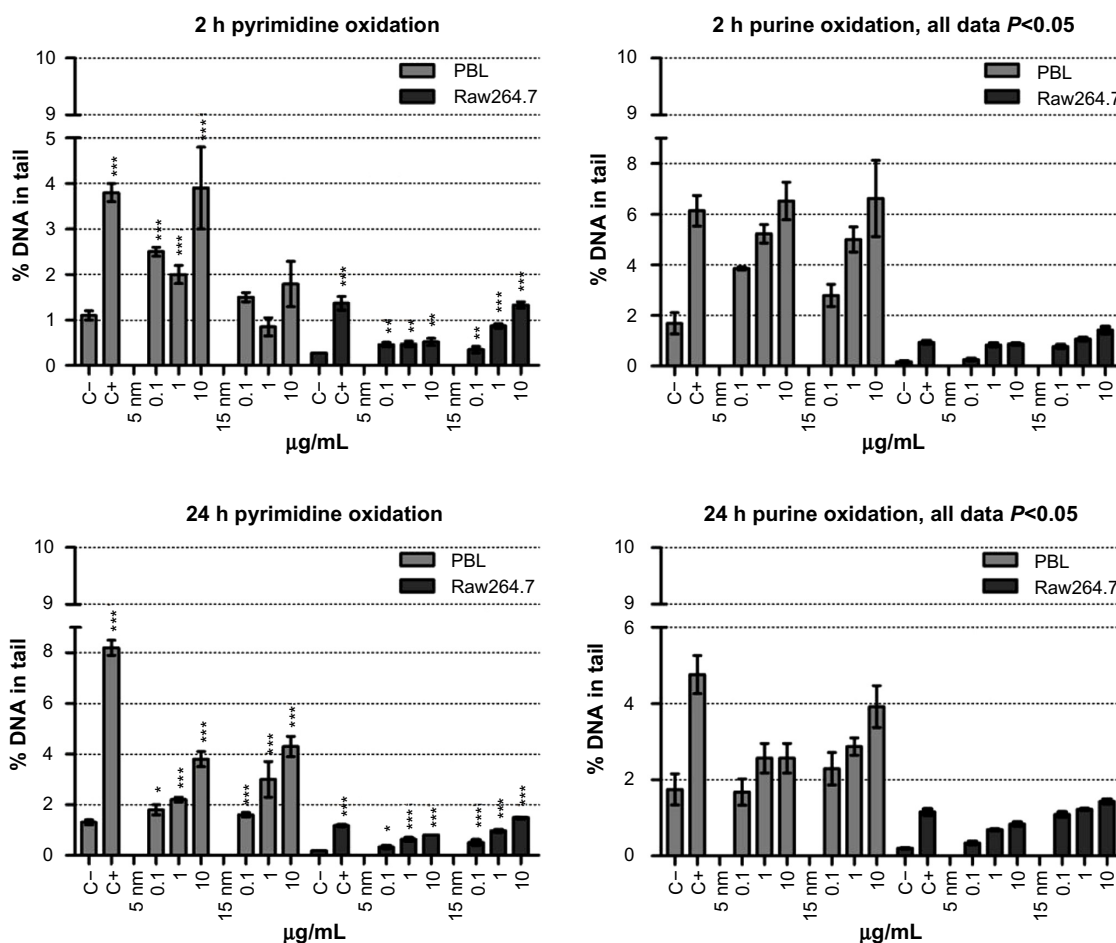


Figure 4 Oxidative DNA damage evaluated using the enzyme-modified comet assay in PBL and Raw264.7.

Notes: Dose-dependent pyrimidines oxidation was detected in PBL after 24 hours of exposure to 5 nm and 15 nm Au NPs ($P=0.05$ and $P=0.028$, respectively). In Raw264.7 the pyrimidines oxidation, detected by the EndoIII enzyme, showed a dose-dependent effect after 2 hours of exposure to 15 nm Au NPs ($P=0.035$) and at 24 hours to 5 nm and 15 nm Au NPs ($P=0.009$ and $P=0.006$, respectively). By Fpg, dose-dependent purines oxidation was observed in PBL after 2 hours of exposure to 5 nm and 15 nm Au NPs ($P=0.009$ and $P=0.007$, respectively), as well as after 24 hours of exposure to 5 nm Au NPs ($P=0.014$). In Raw264.7, a dose-effect relationship was detected following 2 hours of exposure to 15 nm Au NPs ($P=0.01$) and 24 hours of incubation in the presence of 5 nm Au NPs ($P=0.015$). Data are shown as mean \pm SEM (PBL: $n=4$; Raw264.7: $n=6$) of percent DNA in tail. Student's t -test: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

Abbreviations: Au NP, gold nanoparticle; C+, positive control (10 μ M hydrogen peroxide); C-, negative (untreated) control; DNA, deoxyribonucleic acid; EndoIII, endonuclease-III; Fpg, formamidopyrimidine-DNA glycosylase; h, hours; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

Comparison between mass concentrations and numbers of Au NPs

Data obtained by mass concentration exposure indicate that the particle size plays a role in determining cytotoxicity and genotoxicity. However, the same mass concentration of 5 nm and 15 nm Au NPs corresponds to different absolute particle numbers. Consequently, if mass concentrations are expressed as theoretically calculated numbers of Au NPs, MTT assay excludes a size-dependent effect on the cell viability (Figure S1). Following this approach, by MTT the same theoretical number of Au NPs showed cytotoxic effects on PBL and Raw264.7 independently of the dimension. To further evaluate the importance of size in determining cytotoxicity, CBPI was analyzed taking into account the number of Au NPs theoretically calculated from the given

mass concentrations and showed size-independent antiproliferative effects (Figure S2A and B). In contrast, in both cell types the more severe MN induction was observed after the treatments with 15 nm Au NPs either considering the given mass concentrations (Table 3) or the theoretically calculated particle numbers (Figure S2C and D).

To verify these indications, exposures to Raw264.7 cells were carried out by controlling the effective number of Au NPs, ranging from 1×10^9 to 1×10^{11} , that corresponds to different mass concentrations (Au NP 5 nm: from 0.00126 to 0.126 μ g/mL; Au NP 15 nm: from 0.034 to 3.4 μ g/mL). By MTT the size-independent effects exerted by absolute numbers of 5 nm and 15 nm Au NPs on the viability of Raw264.7 cells, both after 2 hours and 24 hours of exposure, were confirmed (Figure 5). CBMN Cyt showed that Au NPs

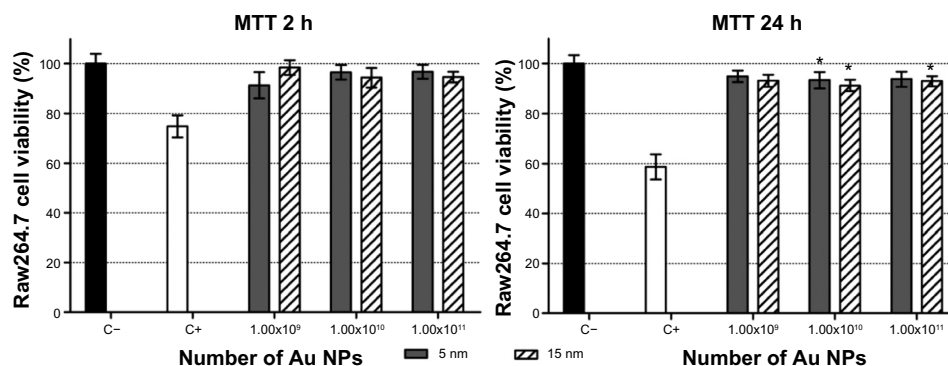


Figure 5 Cell viability in Raw264.7 after 2 hours and 24 hours of exposure to absolute numbers of 5 nm and 15 nm Au NPs.

Notes: Cells were exposed to 1.00×10^9 , 1.00×10^{10} , and 1.00×10^{11} Au NPs and evaluated by MTT assay. The data show the percent of living cells normalized to untreated control (\pm SEM, $n=9$). Statistically significant differences from the C- were determined by the Student's *t*-test ($*P < 0.05$). There were no statistically significant differences between the same number of 5 nm and 15 nm Au NPs.

Abbreviations: Au NP, gold nanoparticle; C+, positive control (10 μ M hydrogen peroxide); C-, negative (untreated) control; h, hours; MTT, methylthiazol tetrazolium; SEM, standard error of the mean.

decreased the proliferation of Raw264.7 cells without statistically significant size effect, while significant differences in genotoxicity and especially in MN induction were found (Table 4). In fact, MN frequencies after exposure to 15 nm Au NPs showed a dose-dependent relationship and differed significantly from 5 nm Au NPs. NPB and NBUD increased in a particle number-dependent way, but only the response obtained with the highest 15 nm Au NPs dose differed significantly from the negative control, and the effects of 15 nm Au NPs were more severe than the damage induced by 5 nm Au NPs.

Table 4 Proliferation and genotoxicity evaluated by CBMN Cyt assay in Raw264.7 cells exposed to an absolute number of Au NPs

	CBPI	MN	NPB	NBUD
C-	1.810 \pm 0.017	2.3 \pm 0.3	1.0 \pm 0.4	0.8 \pm 0.4
C+	1.295 \pm 0.046***	28.7 \pm 2.4***	5.8 \pm 0.8**	6.2 \pm 0.6***
Au NP 5 nm				
1 \times 10 ⁹	1.681 \pm 0.005**	4.3 \pm 0.7	1.0 \pm 0.4	0.8 \pm 0.4
1 \times 10 ¹⁰	1.673 \pm 0.013**	3.7 \pm 0.3*a	1.3 \pm 0.4	1.2 \pm 0.2
1 \times 10 ¹¹	1.661 \pm 0.014**	4.7 \pm 0.3**a	1.5 \pm 0.3 ^a	1.0 \pm 0.4 ^a
Au NP 15 nm				
1 \times 10 ⁹	1.678 \pm 0.005**	6.3 \pm 0.6**	1.3 \pm 0.2	1.3 \pm 0.2
1 \times 10 ¹⁰	1.672 \pm 0.011**	6.0 \pm 0.5**b	2.0 \pm 0.4	1.8 \pm 0.4
1 \times 10 ¹¹	1.665 \pm 0.021**	9.3 \pm 1.2**b	2.3 \pm 0.2* ^b	2.3 \pm 0.2* ^b

Notes: CBPI, MN, NPB, and NBUD frequencies scored by CBMN Cyt assay after Raw264.7 cells exposed to absolute Au NPs numbers. MN frequency in Raw264.7 exposed to 15 nm Au NPs showed a dose-dependent relationship ($P=0.008$), as well as the NPB after exposure to 5 nm and 15 nm Au NPs ($P=0.05$ and $P=0.02$). Dose-dependent NBUD formation was scored in the presence of 15 nm Au NPs ($P=0.008$). Statistically significant differences from C- were determined by the Student's *t*-test ($*P < 0.05$; $**P < 0.01$; $***P < 0.001$), as well as the differences between respective tested conditions ($*P < 0.05$ versus 15 nm Au NPs; $^bP < 0.05$ versus 5 nm Au NPs). Data represent the mean \pm SEM ($n=6$).

Abbreviations: Au NP, gold nanoparticle; C+, positive control (0.1 μ g/mL mitomycin C); C-, negative (untreated) control; CBMN Cyt, cytokinesis-block micronucleus cytome; MN, micronuclei; NBUD, nuclear buds; NPB, nucleoplasmic bridges; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

Discussion

Our results suggest that Au NPs of different size (5 nm and 15 nm) can induce inhibition of cell proliferation mediated by apoptosis, as well as chromosomal damage, aneuploidogenic events, DNA strand breaks, pyrimidines and purines oxidative lesions in human PBL and in a murine macrophage cell line. The chromosomal damage was more severe in Raw264.7 macrophages exposed to 15 nm Au NPs, and overall both Au NPs induced dose-dependent genotoxic effects. Aneuploidy was induced irrespective of the Au NPs size, and the lymphocytes showed a relatively higher percentage of chromosome mis-segregation than macrophages. PBL showed higher DNA strand breaks than Raw264.7, while the macrophages were more sensitive to pyrimidines and purines oxidation. Moreover, our data allowed a correlation between the cytotoxic mechanisms assessed with the tested mass concentration and the absolute number of Au NPs, showing a size-independent relationship. The size of Au NPs, on the other hand, played an important role in determining genotoxic effects in PBL and Raw264.7 cells.

Recent reports on the reactivity of Au NPs focused on their effects on biocompatibility, uptake, toxicity, and mutagenic effects.²⁶⁻³⁰ However, in general, there is a lack of conclusive information on the size as the key factor for Au NPs cytotoxicity and genotoxicity, and on the relationship between size, mass concentration and the absolute particle number.

The cytotoxicity of Au NPs on A549 and NCIH441 cell lines was described, suggesting that the presence of sodium citrate can affect the cell viability and proliferation rather than the size of the particles, which appears to have no effect on active Au NPs endocytosis.⁶ Accordingly, it was shown that not the size of the Au NPs, but the amount of citrate on the

particle surface impaired the viability of the endothelial cells HDMEC and hCMEC, which, in addition, exhibited different uptake behavior for citrate-stabilized gold nanoparticles.⁷ In contrast, the uptake of Au NPs coated with nucleic acid and polyethyleneimine into mammalian cells demonstrated that changes in the nanoparticle size can lead to significant differences in the number of molecules delivered per cell.³¹ Furthermore, the uptake efficiency and the intracellular localization of Au NP-peptide conjugates was shown being directly dependent on the peptide itself, and the intracellular destination was shown to be further determined by Au NP size.³² Au NPs size-dependent bioaccumulation and (eco) toxicity has been demonstrated by metallothionein induction, increased activities of catalase, superoxide dismutase and glutathione S-transferase by three sizes of citrated-coated Au NPs (5, 15, 40 nm) in *Scrobicularia plana*, indicating the activation of response to oxidative stress.¹⁹

Results obtained exposing Balb/3T3 cells to the same citrate-stabilized 5 nm and 15 nm Au NPs here investigated revealed cytotoxic effects, measured by colony forming efficiency, for 5 nm Au NPs but not for 15 nm Au NPs.²⁰ Coradeghini et al showed also that the changes in cytoskeleton structure of Balb/3T3 cells resulted in pronounced modification of the cell shape and reduction in the expression and degradation of the clathrin heavy chain, but without significant effects in the expression of caveolin.²⁰ Citrate-stabilized Au NPs (20 nm) influenced cell cycle pathways in human embryonic lung fibroblasts, reducing the expression of critical checkpoint proteins and causing oxidative stress and alteration of genes associated with genomic stability and DNA repair.³³ Citrate-stabilized Au NPs downregulated, in a size-dependent manner, cellular responses induced by interleukin 1-beta both in vitro in THP-1 human leukemia monocytic macrophages and in vivo in C57BL/6 male mice.³⁴

Nevertheless, the size dependency of adverse effects in human blood is not linear.³⁵ Studies with human blood cells showed that 30 nm Au NPs induced concentration-dependent effects on hemolysis, reactive oxygen species generation, and platelet aggregation.³⁶ In addition, studies in mice showed that the bioaccumulation and toxicity of differently sized (5 nm, 10 nm, 30 nm, and 60 nm) PEG-coated Au NPs was not size dependent.³⁷

In conclusion, both the literature and the results presented in this study demonstrate no clear correlation between the cytotoxicity of differently sized Au NPs and the equivalent spherical diameter. We found that at early time points (2 hours), the highest mass concentration of 5 nm Au

NPs showed a more severe cytotoxicity both in PBL and Raw264.7 compared to the corresponding doses of 15 nm Au NPs. However, taking into account the theoretically calculated numbers of Au NPs corresponding to the different tested mass concentrations, it was not possible to conclude that 5 nm Au NPs induced higher cytotoxicity than 15 nm Au NPs. These indications were further confirmed by experiments carried out in Raw264.7 cells with a fixed absolute number of differently sized Au NPs.

In addition to size, other physicochemical properties of NPs, such as surface charge, are proposed to be critical determinants of their toxic potential.^{38,39} For instance, the evaluation of differently charged Au NPs (neutral, positively, and negatively charged) in a human keratinocyte cell line (HaCaT) showed that the cell morphology was disrupted in a dose-dependent way by all three NPs: the surface charge was the major determinant of the cellular processes, with the charged NPs inducing cell death through apoptosis and neutral Au NPs through necrosis.⁴⁰ However, as highlighted in this paper, the cell type also plays an important role in activating apoptosis: PBL, in fact, showed higher apoptotic responses than Raw264.7 macrophages.

Interestingly, considering both the mass concentrations and the absolute number of Au NPs, from our study it is clear that the size of Au NPs plays an important role in determining genotoxic mechanisms, although in a nonlinear manner. Both cell systems were more sensitive to 15 nm Au NPs than to 5 nm Au NPs, as evaluated through the different biomarkers of chromosomal damage (MN, NPB, and NBUD). The evaluation of the genotoxic effects in Raw264.7 cells, based on the absolute numbers of Au NPs, confirmed nonlinear size-dependent chromosomal damage. However, this result cannot explain the time-dependent primary DNA damage, as the 5 nm Au NPs were the most harmful both in PBL and in Raw264.7 macrophages. In addition, the lowest dose of 5 nm Au NPs showed in both cell types minor aneuploidogenic effects compared to 15 nm Au NPs, and the more clastogenic behavior at low doses was consistent with the described NPB/MN ratio, which provides an important fingerprint for distinguishing the genotoxic mechanisms of different clastogenic and aneuploidogenic agents.⁴¹

To date, few genotoxicity studies have been performed with Au NPs. Although minor focal inflammatory changes occurred after a single instillation of 2 nm, 20 nm, and 200 nm Au NPs into the lung of male adult Wistar rats, genotoxicity was not observed as assessed by the erythrocyte micronucleus test and the comet assay.⁴² In addition, 2 nm, 20 nm, and 200 nm Au NPs failed to induce MN formation in

human blood and did not damage DNA in the liver and lung of exposed adult male Wistar rats.⁴³ The dosing, however, was limited by the stability of the gold suspensions, and Downs et al stated that the results would have looked different upon increasing the gold doses.⁴³

Au NPs capped with sodium citrate or polyamidoamine dendrimers interacted with hepatocytes HepG2 and peripheral blood mononuclear cells, and exhibited in vitro cyto- and genotoxicity even at low concentrations.⁴⁴ In zebrafish *Danio rerio* exposed to low doses of 12 nm and 50 nm Au NPs, size and exposure length modulated the genotoxicity and the expression of genes involved in DNA repair, apoptosis, and oxidative stress.⁴⁵ Flow cytometry and real-time PCR analysis of apoptotic genes and ATP depletion measurements suggested that 17 nm Au NPs induced cell damage through extrinsic and intrinsic apoptotic pathways in human lung adenocarcinoma A549 cells.⁴⁶

Au NPs (12 nm) were internalized by the endolysosomal pathway in Balb/3T3 fibroblasts after 10 days of exposure, and although they were not a severe cytotoxicant, Au NPs induced DNA damage.⁴⁷ Moreover, the comet assay showed DNA damage in MRC-5 human lung fibroblasts treated with 20 nm Au NPs, and FISH analysis revealed that the majority of the chromosomal breaks were undetectable telomeres.⁴⁸ Li et al explained this genomic instability with the differences in the oxidative stress-related proteins as well as with proteins associated to the cell cycle regulation and cytoskeleton and DNA repair.³³ Similarly, our studies directly demonstrated that apoptosis, DNA oxidation, and aneuploidy could explain the genotoxicity observed in PBL and Raw264.7 cells. In addition to the alterations of the mitotic spindle, to the weakness of mitotic checkpoints and to cytoskeletal defects, a growing body of evidence suggests that aneuploidy can be also induced by epigenetic mechanisms.^{49,50} Chromatin condensation and reorganization, as well as global gene expression alterations, were observed in fibroblasts exposed to Au NPs, providing further insights into the molecular mechanisms underlying toxicity of Au NPs and their impact on epigenetic processes.^{51–53}

Conclusion

We can conclude that differently sized Au NPs induced cytotoxicity and genotoxicity through different mechanisms of action. While Au NPs size did not play a pivotal role in determining cytotoxicity, the size was a fundamental factor in inducing genotoxicity. In fact, the most severe genotoxicity in PBL and Raw264.7 was observed following exposure to 15 nm Au NPs. These observations were confirmed, also

taking into account the particle numbers as dose-metric, and these results might represent groundwork for further studies aimed at investigating the mechanistic interactions between Au NPs and DNA. Nevertheless, cytotoxic and genotoxic mechanisms, as well as the exposure time, could be ascribed to the different cell type-related reactivity. Au NPs induced cytotoxicity through apoptotic pathways, and the premutagenic DNA oxidation appears to be an important factor in inducing genotoxicity. Finally, to our knowledge, this study reports for the first time increased aneuploidy after Au NPs exposure in vitro in a human and in a murine cell model.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

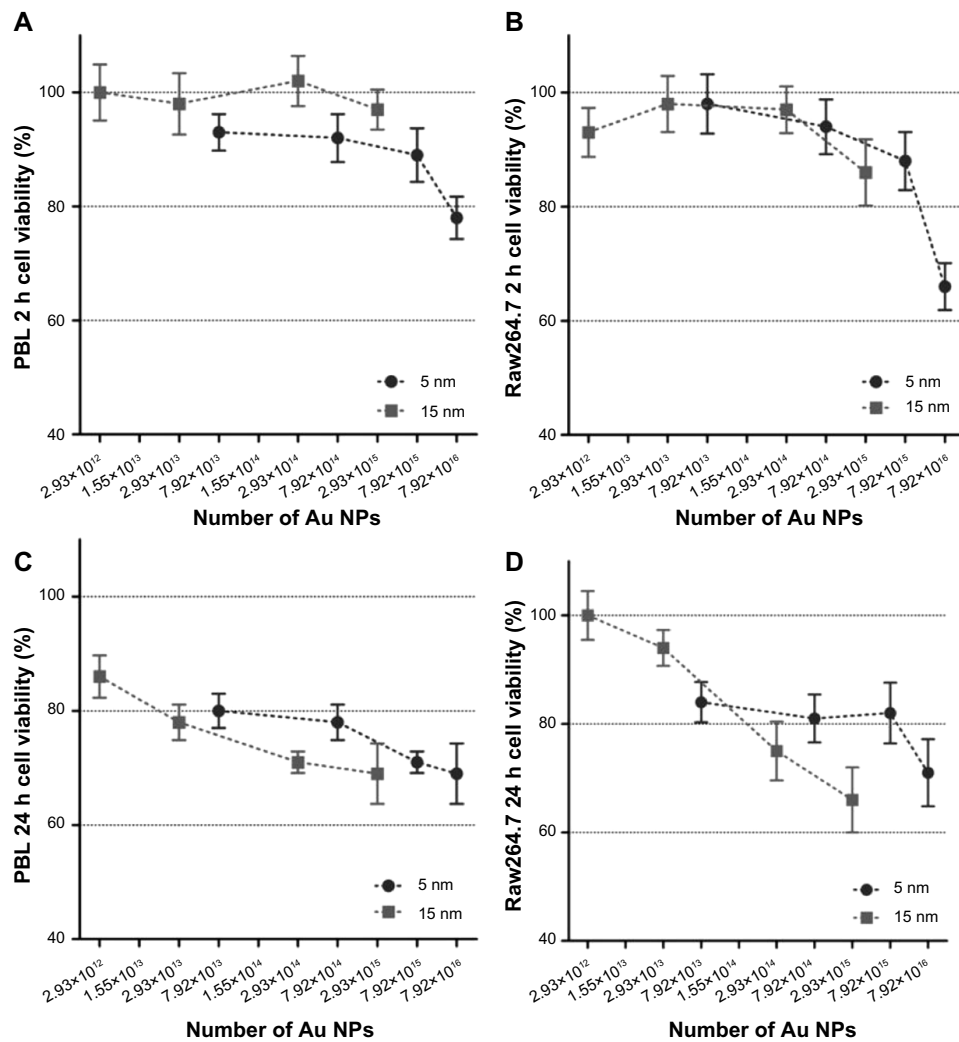


Figure S1 Cell viability evaluated by MTT assay in function of the theoretically calculated number of 5 nm and 15 nm Au NPs.

Notes: Mass concentrations (0.1–100 $\mu\text{g/mL}$) were expressed as theoretically calculated number of Au NPs ranging from 2.93×10^{12} to 7.92×10^{16} . In PBL (A and C) and Raw264.7 (B and D), after 2 hours (A and B) and 24 hours (C and D) treatment, no statistically significant differences among the cytotoxicity of 5 nm and 15 nm Au NPs were observed. Data show the percent of viable cells normalized to untreated control (\pm SEM, $n=9$). Statistical analysis was performed by Student's *t*-test.

Abbreviations: Au NP, gold nanoparticles; MTT, methylthiazol tetrazolium; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.

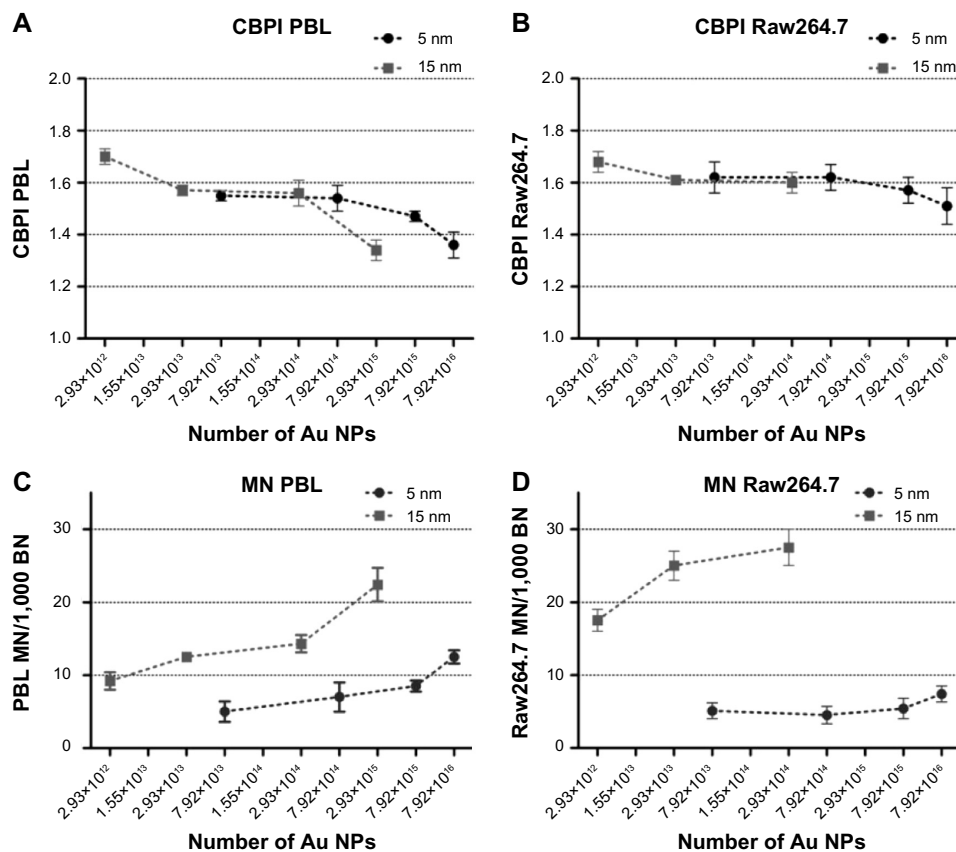


Figure S2 CBPI and MN frequency evaluated by CBMN Cyt assay in function of the theoretically calculated number of 5 nm and 15 nm Au NPs. **Notes:** While CBPI shows no differences in antiproliferative activity of 5 nm and 15 Au NPs in PBL (A) and Raw264.7 (B), MN frequency was higher in PBL (C) and Raw264.7 (D) exposed to 15 nm Au NPs than to 5 nm ($P < 0.05$). Data are shown as mean \pm SEM (PBL: $n=4$; Raw264.7: $n=6$). Statistical analysis was performed by Student's *t*-test. **Abbreviations:** Au NP, gold nanoparticle; BN, binucleated cells; CBMN Cyt, cytokinesis-block micronucleus cytome; CBPI, cytokinesis block proliferation index; MN, micronuclei; PBL, peripheral blood lymphocytes; SEM, standard error of the mean.