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Ultrastructural hepatocytic alterations induced by silver nanoparticle toxicity

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ABSTRACT

Silver nanoparticles (SNPs) are widely used in nanomedicine and consuming products with potential risk on human health. While considerable work was carried out on the molecular, biochemical, and physiological alterations induced by these particles, little is known of the ultrastructural pathological alterations that might be induced by nanosilver materials. The aim of the present work is to investigate the hepatocyte ultrastructural alterations that might be induced by SNP exposure. Male rats were subjected to a daily single dose (2 mg/kg) of SNPs (15–35 nm diameter) for 21 days. Liver biopsies from all rats under study were processed for transmission electron microscopy examination. The following hepatic ultrastructural alterations were demonstrated: mitochondria swelling and cristolysis, endoplasmic reticulum disruption, cytoplasmic vacuolization, lipid droplets accumulation, glycogen depletion, karyopyknosis, apoptosis, sinusoidal dilatation, Kupffer cells activation, and myelin figures formation. The current findings may indicate that SNPs can induce hepatocyte organelles alteration, leading to cellular damage that may affect the function of the liver. These findings might indicate that SNPs potentially trigger hepatocyte ultrastructural alterations that may affect the function of the liver with potential risk on human health in relation to numerous applications of these particles. More work is needed to elucidate probable ultrastructural alterations in the vital organs that might result from nanosilver toxicity.

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Introduction

Silver nanoparticles (SNPs) have been rapidly involved in nanomedicine, consuming products, and environmental issues due to their unique antimicrobial, antifungal, and antiviral properties [1,2]. While the number of commercial and industrial products employing SNPs was 30 in 2006, it has grown to 1300 by the beginning of 2014 in various applications in one way or another [3]. SNPs have been widely used in alginate fibers and wound dressing in treating wounds, burns, skin ulcers, and catheter-related infections [4,5]. Furthermore, nanosilver is also used in coating medical instruments and tools together with commercial brands of sunscreen, medical masks, gels, cosmetics, bone cement, tooth paste, deodorants, dental resin composite, and shampoo [5]. Moreover, these fine particles are invested in medical cloth, footwear, athletic shirts, and textile manufacturing to limit odor due to sweating [6,7]. In addition, SNPs have unique optical, electrical, and thermal properties and are being used in biological and chemical sensors [8].

The smaller nature of SNPs together with their high surface area to mass ratio enables them to penetrate the cell easily by diffusion or endocytosis as biological molecules do [9,10]. Their toxicity might be related to the charge and functional groups on the surface of these particles and due to the release of silver ions. The released ions are neutralized by binding or interacting with the cell component leading to depletion of dissolved oxygen and generating reactive oxygen

species (ROS) together with reactive nitrogen species (RNS) leading to molecular and biochemical alterations [11–17]. *In vitro* toxicological studies showed that SNPs could initiate ROS production that could damage plasma membrane and cell organelles stimulating organelle oxidative damage and cytoskeleton disruption [18]. Moreover, SNPs can also produce cytotoxic RNS with negative impact on the mitochondrial function, membrane integrity, glutathione depletion, and inflammatory cytokine release [19–22]. In addition, SNPs can interact with the cellular proteins with possible adsorption of these molecules at the surface of the particles forming what is known as the nanoparticle-protein corona [23]. The toxicity of SNPs is related with corona composition and associated with the particle size, shape, and surface area [24]. In addition, probable heat alteration due to the plasmonic effect of SNPs might result in conformational changes in the macromolecules of the cell. On the other hand, Asharani et al. [25] concluded that SNPs could induce DNA damage, negatively affect DNA integrity, increase chromosomal aberrations, disturb G2/M cell cycle, and reduce metabolic activity and ATP production in treated cells. Moreover, other toxicological studies have demonstrated size-dependent genotoxic and cytotoxic consequences associated with SNP exposure [20–22].

SNPs can have the same dimensions of the biological molecules, with the possibility of being adsorbed on the surface of these molecules in the tissues and cells [7]. Nanoparticles less than 100 nm in diameter can enter cells, while those less than

40 nm can enter the nucleus [26, 27]. Recent studies indicated that small SNPs were more toxic than the larger ones [28–36]. Particle size-dependent cellular toxicity was reported by Kim et al. [37], where 10 and 20 nm SNPs were more toxic than the larger ones (50 and 100 nm). Other reports indicated that SNPs could stimulate inflammation and oxidative damage affecting cellular metabolism and membrane integrity via lipid peroxidation and protein denaturation [21,38,39].

The liver regulates body metabolism and fight body toxification by many exogenous challenges that make it highly susceptible to their adverse and toxic effects [40]. The hepatocytes form the main parenchymal tissue of the liver are involved in protein, cholesterol, bile salts, and phospholipids synthesis together with the detoxification, transformation, and modification of exogenous and endogenous materials. Some reports explained that SNPs could accumulate mainly in the liver and other vital organs, with the mitochondria as primary organelle target [8,13,28–33]. Moreover, toxicological reports showed that SNPs were capable to induce injuries in the liver, kidney, testes, heart, brain, and lungs [10,28,29]. In addition, some histopathological reports explained that SNPs can induce cytoplasmic accumulation, focal necrosis, bile-duct hyperplasia, fibrosis, edema, and inflammatory cell infiltration in the hepatic tissues [30,32]. Furthermore, SNPs were reported to induce hematological and biochemical alterations, while case studies revealed that some silver-coated wound dressing elevated liver enzymes [33–36]. In addition, exposure to SNPs increased the serum levels of alanine aminotransferase (ALT), aspartate aminotransferase (AST), and alkaline phosphatase together with DNA damage and apoptosis [32,39]. Morphometric and histological hepatic alterations were also induced by 20-nm SNP subjection [10,28,31].

Exposure to SNPs is becoming part of our lives with wide use of these particles in different sectors, making these particles persisted in our environment with possible potential risk to our health. These concerns need to be considered before determination if SNP benefits outweigh their potential risks with special attention needed toward size, composition, and behavior of these particles. Limited information, if any, is available on the ultrastructural alterations induced in the hepatic tissues by SNPs, a concern to be clarified by the present study.

Materials and methods

Animals and conditions

Twenty healthy male Wistar albino rats of King Saud University colony of the same age (10–12 weeks old) weighing 210–230 gm were used. The animals were housed at 24°C ± 1°C, on 12 h light–12 h dark/light cycle, randomly assigned and separately caged to one test group and a control one (10 rats each).

Silver nanoparticles

Spherical SNPs (15–35 nm diameter) dissolved in deionized water containing 0.1 mM sodium citrate were obtained from the Department of Nanotechnology of Laser Zentrum-Hannover, Germany. SNPs were fabricated by picosecond-pulsed

laser ablation in liquids (PLAL). The ablation process was carried as follows: a silver foil (8 × 8 × 0.1 mm; purity: 99.99%, Goodfellow GmbH, Germany) was fixed within a self-constructed ablation chamber made of Teflon which was filled with 35 ml of deionized water containing 0.1 mM sodium citrate, resulting in a liquid column of 10 mm. The laser beam was coupled into a galvanometric scanner laser (HurrySCAN II-14, Scanlab AG Germany), which allowed deposition of laser pulses with controlled interpulse distance on the silver target after focusing with a 56-mm telecentric lens. Liquid stirring was applied using a Teflon rotor for continuous liquid agitation and quick removal of ablated nanoparticles from the process zone. Determination of laser spot size on the target for fluence calculation was performed according to Farkas et al. [41]. Ligand-free SNPs were generated by applying 250 µJ pulse energy with a repetition rate of 33 kHz and laser fluency of 12.7 Jcm⁻². Target position was selected with respect to the focus position in air and to yield the highest ablation rate in deionized distilled water. The optical extinction spectra of the fabricated particles were measured by using UV-Vis spectroscopy (Shimadzu 1650), while the nanoparticle micrographs were obtained by the use of a scanning electron microscope (Quanta 400 FEG).

Experimental protocol

Following a period of stabilization (7 days), the control animals received a daily i.p. injection of the vehicle (400 µl deionized water containing 0.1 mM sodium citrate) only, while test group members received a daily i.p. injection of 400 µl of SNPs (2 mg/kg) for 21 days.

All animals were handled and all experiments were conducted in accordance with the protocols approved by King Saud University animal care ethical committee. In addition, the experimental procedures were carried out in accordance with international guidelines for care and use of laboratory animals.

Electron microscopic investigation

Small pieces of liver from each rat minced into small cubes of 1 mm in length were fixed in 2.5% glutaraldehyde fixative in 0.1 M phosphate buffer (pH 7.4) for 24 h at 4°C. Specimens were then post-fixed in 2% osmium tetroxide (OsO₄) in cacodylate buffer for 90 min at room temperature. Tissues were washed in the buffer and dehydrated at 4°C through a gradual series of acetone and embedded in Epon-araldite resin mixture [42–44]. Semithin sections (500–1000 nm) were obtained in a Leica EM UC6 ultramicrotome mounted on glass slides, stained with toluidine blue (1%), examined, and photographed using light microscopy. Ultrathin sections (60 nm thick) were obtained with a diamond knife on Leica EM UC6 ultramicrotome and mounted on 200-mesh hexagonal copper grids, with 0.5% uranyl acetate and 3% lead citrate using the Leica automated EM stainer [45]. Ultrasection examination and structural abnormality evaluation was carried out by using Jeol transmission electron microscope at 80 kv (JEM-1011, Japan).

Results

Silver nanoparticle characterization

As represented in Figure 1(a–c), the used SNPs demonstrated spherical morphology with size variation range of 15–35 nm in diameter where the maximum particle count was at 20 ± 5 nm size. In addition, the UV-visible spectrum of the particles presented single symmetric absorption band with maximum peak of absorbance around ~400 nm that correspond to the normalized absorption spectrum of SNPs.

Control rats

Semithin and ultrathin sections of the control rats demonstrated normal hepatocytes as well as normal sinusoids lined with Kupffer cell. The hepatocytes revealed normal nucleoplasm with nuclei surrounded by even distinct nuclear envelop demonstrating heterochromatin adjacent to the border. The cytoplasm of these hepatocytes appeared crowded with organelles mainly the mitochondria, endoplasmic reticulum (ER), free and bounded ribosomes, lysosomes, and glycogen particles (Figures 2a,b). The control hepatocytes

demonstrated mitochondria in orthodox conformation together with regular parallel organized ER.

Rats exposed to 20 nm SNPs

In comparison with the control group, the hepatic tissues of SNP-treated rats exhibited the following remarkable alterations:

- Sinusoidal dilatation: Examination of semithin sections stained with toluidine blue reveals sinusoidal dilatation in the hepatic tissues of rats treated with SNPs (Figures 3a, b). Sinusoidal dilatation was also demonstrated in the electron micrograph sections where red blood cells and debris were observed in the dilated sinusoids (Figure 4).
- Alterations in the rough endoplasmic reticulum: Partial destruction and reduction profile together with a loss of arrays, stacks, and shortening of rough ER were demonstrated (Figure 5a). Occasional vesiculated and dilated ER were also seen (Figure 5b). Moreover, degranulation of the rough ER and ribosomal dropping were also observed.
- Mitochondrial alterations: The pleomorphic mitochondria of SNP-treated rats demonstrated swelling, cristolysis,

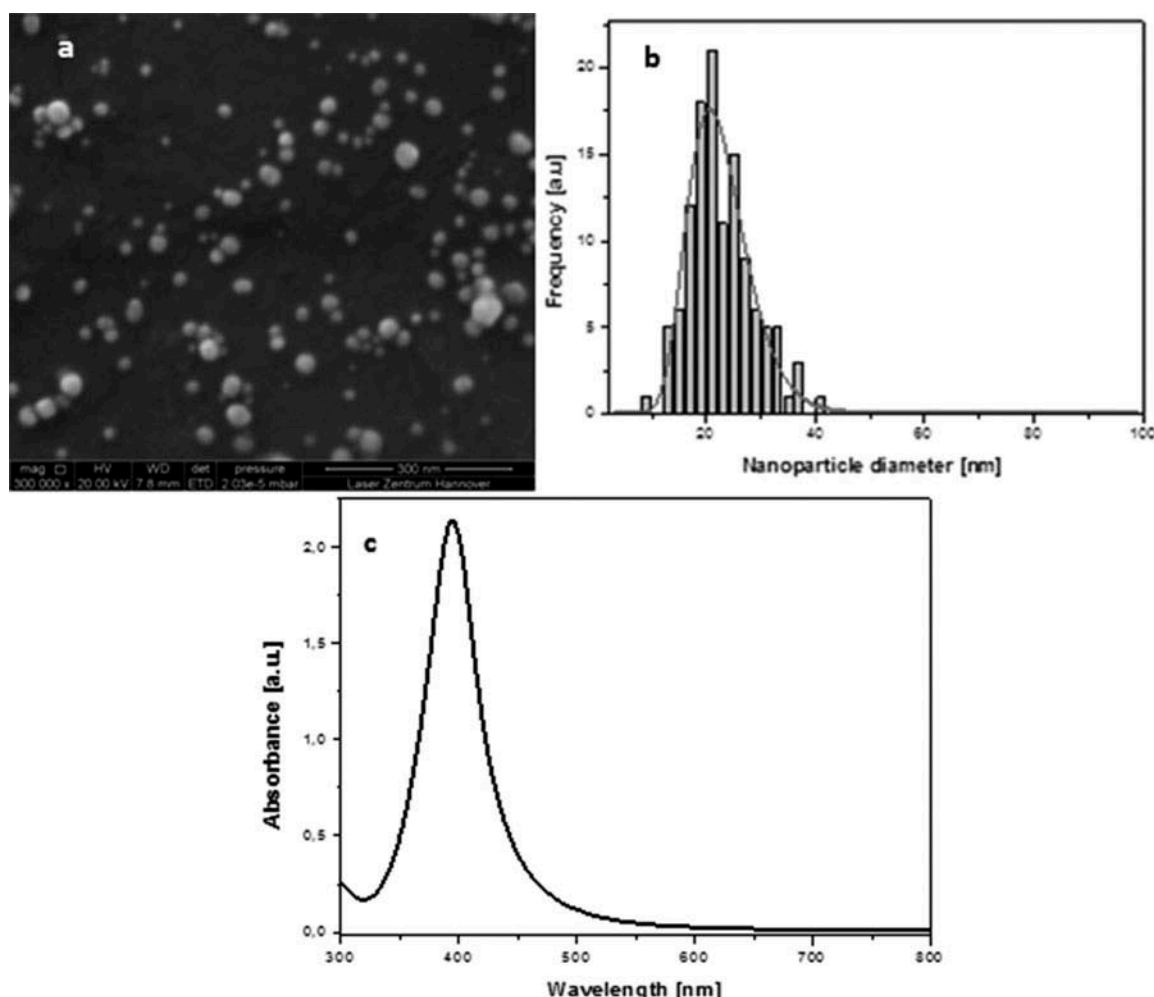


Figure 1. (a–c) Physical characterization of the used SNPs demonstrating: (a) Scanning electron micrograph demonstrating spherical morphology of the used SNPs. (b) Histogram showing that the particle size distribution of the used SNPs is within the range of 15–35 nm in diameter, with maximum particle count at 20 ± 5 nm size. (c) A plot UV-visible spectrum of the used particles showing single symmetric absorption band with maximum peak of absorbance around ~400 nm which corresponds to the normal absorption spectrum of SNPs.

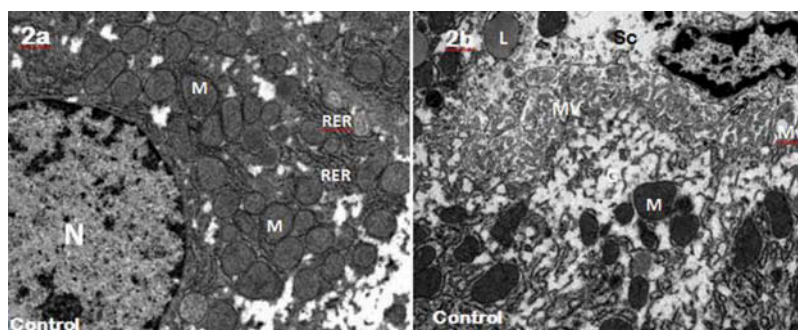


Figure 2. (a–b). Transmission electron micrographs of control rats demonstrating: (a) Control hepatocyte showing normal organelle ultrastructure with rounded nucleus (N), numerous of mitochondria (M), rough endoplasmic reticulum (RER), and bounded ribosomes. $\times 12,000$. (b) Control hepatocyte showing lipid droplets (L), glycogen particles (G), sinusoidal cell (Sc), and microvilli (Mv). $\times 15,000$.

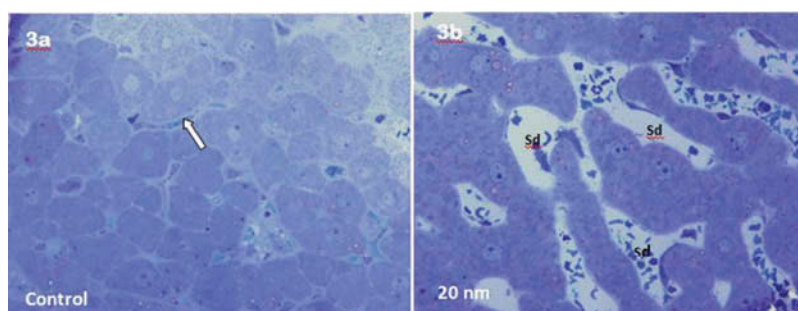


Figure 3. (a–b). Toluidine blue stained semithin sections of: (a) Control rat demonstrating normal sinusoids (arrow). $\times 1000$. (b) SNP-treated rats demonstrating dilated sinusoids (Sd) with different architecture from that of the control rat indicating vascular lesion. $\times 1000$.

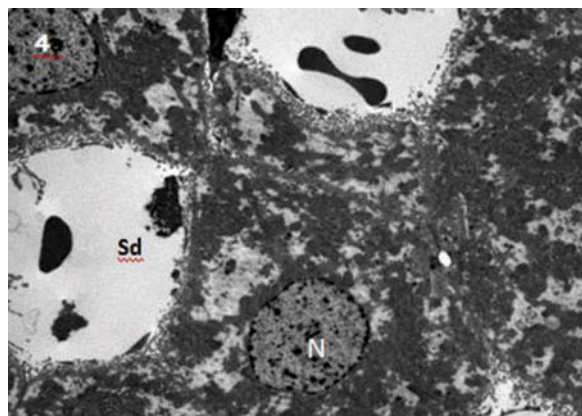


Figure 4. Transmission electron micrograph of SNP-treated rats demonstrating red blood cells and debris in the dilated sinusoids (Sd). $\times 4000$.

angulations, and elongation along their longitudinal axis with occasional ruptured membranes (Figures 6a–c). Some swollen mitochondria were short and emarginated in comparison to well-preserved mitochondria in the control hepatocytes. In addition, some mitochondria demonstrate SNPs within their matrices.

- **Cytoplasmic vacuolization:** The cytoplasm of the affected hepatocytes appeared disorganized with variable sizes of cytoplasmic vacuoles (Figure 7).
- **Lipid droplet accumulation:** In comparison with the control hepatocytes, marked elevation number of variable sizes of lipid droplets was demonstrated in the cytoplasm of some affected hepatocytes (Figure 8).

- **Glycogen depletion:** In comparison with the control hepatocytes, evident glycogen depletion was clear in the hepatocytes of rat subjected to SNPs (Figure 9).
- **Multilamellar body formation:** Some affected cells demonstrated membranous phagolysosomes multi-layered myelin figures. Some of these bodies were embedded in cytoplasmic vacuoles (Figure 10).
- **Nuclear alterations:** Some hepatocytes of SNP-treated rats demonstrated heterochromatic micronuclei, karyopyknosis, nuclear membrane irregularity, and occasional apoptosis (Figures 11a–d).
- **Multiple lysosomes:** Hepatocytes of SNP-treated rats demonstrated multiple lysosomes some of which were membranous phagolysosomes (Figure 12).
- **Kupffer cell alterations:** Semithin and ultrathin sections of the hepatic tissue of SNP-treated rats demonstrated hypertrophoid Kupffer cells with irregular pyknotic nuclei (Figure 13). Some Kupffer cells showed vacuoles, multiple lysosomes, occasional phagocytic deposits, and minute electron-dense bodies.

Discussion

SNPs are important materials in industry and nanomedicine because of their unique biological and physiochemical characteristics. The results of the present work indicated that SNPs could induce variable ultrastructural alterations in most organelles of the hepatocytes. This finding is in consistence with some previous studies that indicated that the liver is a target organ for SNPs [28,29,31].

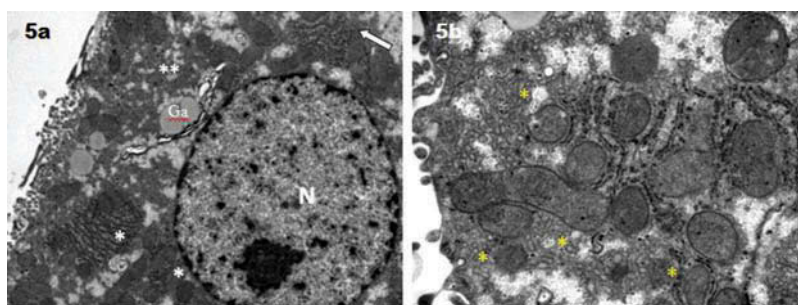


Figure 5. (a–b). Transmission electron micrographs of SNP-treated rats demonstrating: (a) Endoplasmic reticulum disruption (arrow), loss of array (*), ER reduction profile (**) and Golgi apparatus trans face (Ga). Note that the ER cisternae are broken up and fragmented. $\times 10,000$. (b) Vesiculated endoplasmic reticulum (stars) that might indicate fluid ingress or storage products. $\times 30,000$.

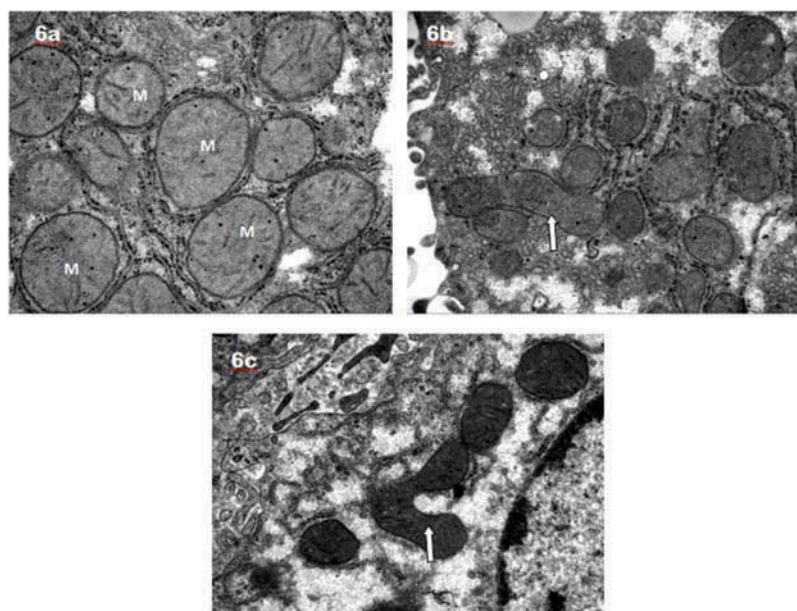


Figure 6. (a–c). Transmission electron micrographs of SNP-treated rats demonstrating: (a) Swollen mitochondria (M) with partial cristolysis. Note the lucent mitochondrial matrix and disarrangement of cristae. In addition, some mitochondria demonstrate SNPs (dark dots) within their matrices. $\times 40,000$. (b) Elongated mitochondrion (arrow) along its longitudinal axis with other mitochondria exhibiting variable size and shape. $\times 30,000$. (c) Angular mitochondrion (arrow). Note the rupture of the mitochondria cristae together with glycogen depletion in the cytoplasm of the demonstrated hepatocyte. $\times 30,000$.

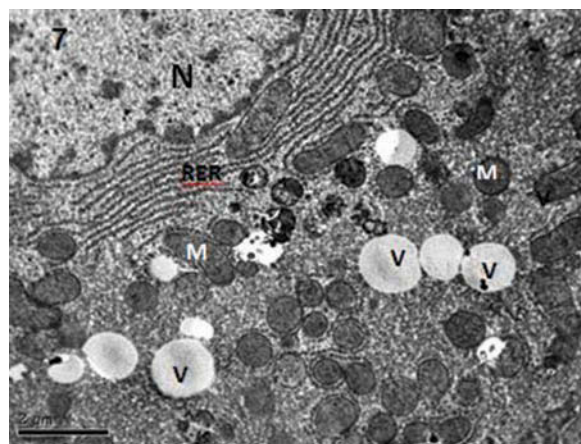


Figure 7. Transmission electron micrograph of SNP-treated rat demonstrating multiple cytoplasmic vacuoles (V) of different sizes. $\times 15,000$.

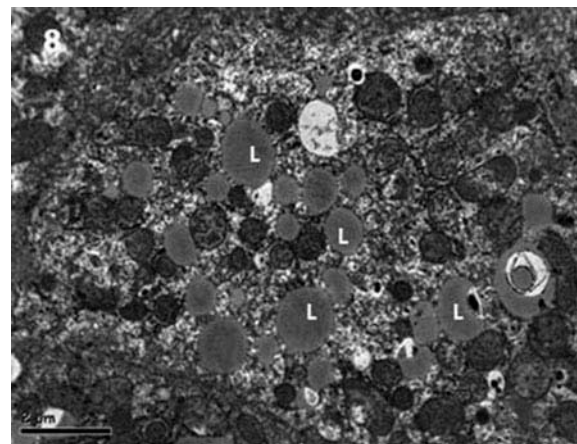


Figure 8. Transmission electron micrographs of SNP-treated rat demonstrating numerous lipid droplets (L) occupying most of the cytoplasm. $\times 15,000$.

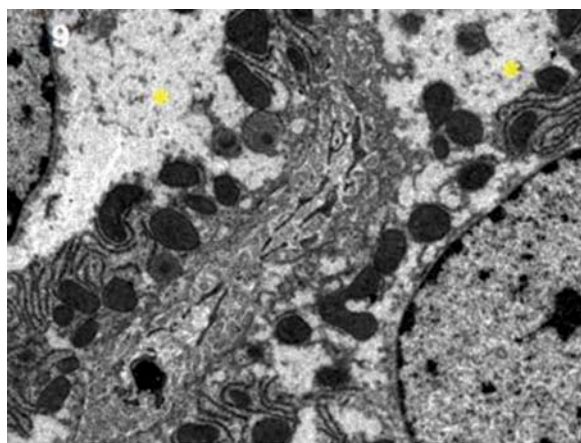


Figure 9. Transmission electron micrograph of SNP-treated rats demonstrating glycogen depletion (*). $\times 15,000$.

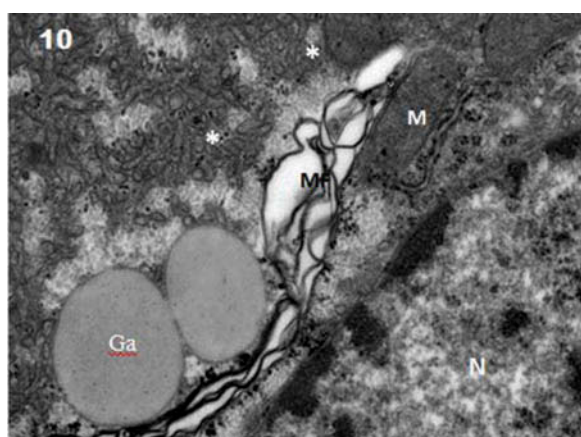


Figure 10. Transmission electron micrograph of SNP-treated rat demonstrating myelin figures (MF) in whorled arrays membranous structure. Note Golgi apparatus trans face (Ga) and dilated degenerative rough ER (*) in the cytoplasm. $\times 40,000$.

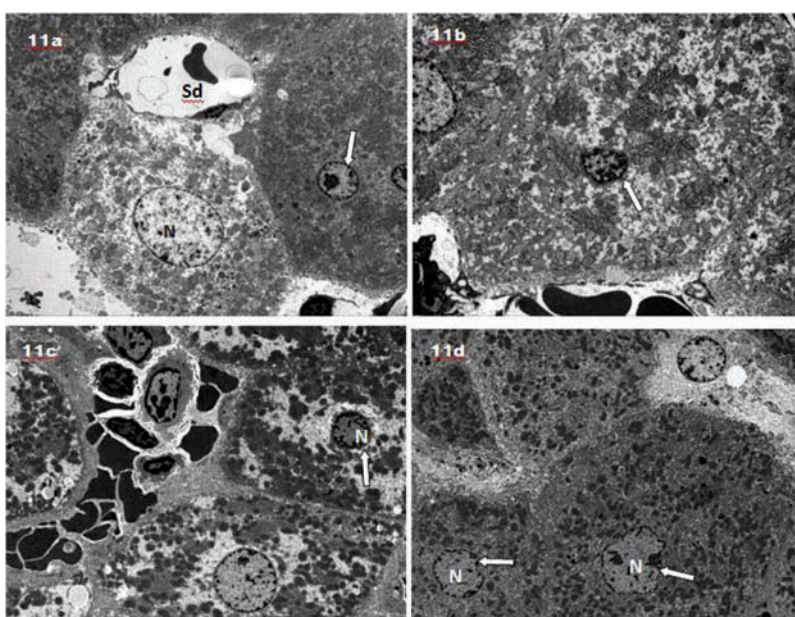


Figure 11. (a–d). Transmission electron micrographs of SNP-treated rats demonstrating hepatocytes with: (a) Micronucleus (arrow). Note the nucleus size ratio of the indicated hepatocyte with that of the hepatocyte on the left showing normal nucleus. $\times 3000$. (b) Apoptotic hepatocyte (arrow). Note the formation of granular emarginated chromatin masses against the nuclear membrane. $\times 5000$. (c) Karyopyknosis (arrow). Note chromatin condensation and nucleus shrinkage. $\times 3000$. (d) Nuclear membrane irregularity (arrows). Note the difference from the smooth nuclear membrane of the control hepatocyte (Figure 2a). $\times 3000$.

SNPs enter the cells by endocytosis targeting the organelles and deposited in the nucleus with probable consequence of oxidative damage to DNA. The results of the present study illustrated that the mitochondria were the most affected organelles by the toxicity of these particles. This is in agreement with other findings where Asharani et al. [25] reported that SNPs could penetrate into cellular organelles specially the mitochondria. The present study showed that mitochondrial damage was demonstrated in the form of swelling and cristolysis due to SNP exposure. These together might indicate alteration in the osmolarity of the hepatocyte causing salt influx to the mitochondrial matrix and affecting the integrity of the inner membrane. Mitochondrial cristae play an important role in the function of this organelle [46–48]. Cristolysis induced by SNPs might affect oxidative phosphorylation and ATP production due to the effect of these particles in the efficiency of the mitochondrial electron transport system [49–52]. The access of SNPs into the mitochondria may stimulate cytotoxicity by ROS generation. Several reports indicated that high level of ROS in the mitochondria could cause damage and depolarization in the mitochondrial membranes causing ATP synthesis interruption [53–54]. In addition, ROS generation by SNPs may affect the DNA repair, induce DNA damage, and activate the apoptotic gene P_{53} . Furthermore, some previous studies indicated that SNPs might induce metallothionein upregulation and heme oxygenase-1 gene upgrading as a defend mechanism of detoxification and protection from free radical stress induced by these particles [52–53]. Some investigators reported DNA damage in human mesenchymal stem cell after 1, 3, and 24 h exposure to SNPs at 0.1 $\mu\text{g/ml}$ by both comet assay and aberration tests [49].

The results of the present work indicated that SNPs could affect hepatocyte rough ER in the form of destruction, loss of array, vesiculation, and stack shortening. This stress might be resulted due to hepatocyte redox disturbance in the ER

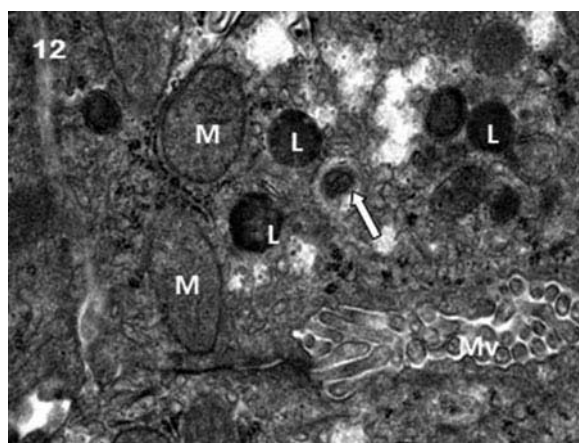


Figure 12. Transmission electron micrograph of SNP-treated rat demonstrating multiple lysosomes (L), phagolysosomes (arrow), mitochondrial cristolysis (M), and microvilli (Mv) in Disse's space. $\times 40,000$.

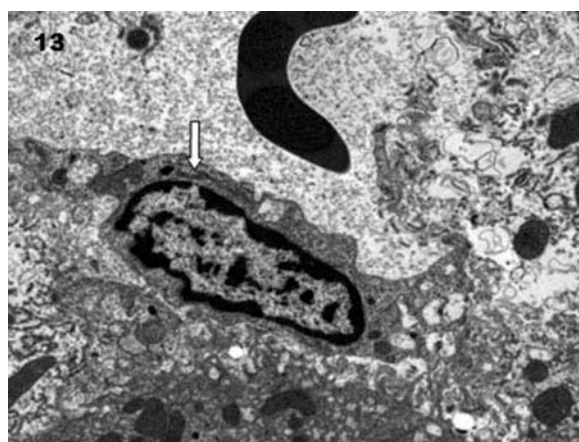


Figure 13. Transmission electron micrograph of SNP-treated rat demonstrating hypertrophied Kupffer cell with irregular pyknotic nucleus (arrow). Note the phagocytized deposits in the cytoplasm. $\times 10,000$.

membranes induced by SNPs. These together would indicate that SNPs would decrease the surface density of the endoplasmic reticulum that may negatively impact protein and lipid synthesis by the affected hepatocytes. Shannhan et al. [55] reported affinity of SNPs toward hydrophobic proteins for protein corona formation and electrostatic interaction. Recent studies indicated that SNPs always interact with cellular proteins forming corona prior to their biological action [24].

The hepatocytes of SNP-treated rats demonstrated partial glycogen depletion. This may indicate that the oxidative stress induced by SNPs may affect the activity of some enzymes necessary for glycogen synthesis. Moreover, lipid droplet accumulation in the cytoplasm of the affected hepatocytes might indicate impairment of fatty acid removal due to SNP toxicity or inability of the affected cells to catabolize fats. In addition, SNP exposure induced lysosomal autophagy which might indicate the need of the affected hepatocytes to degrade the foreign materials and damaged organelles. This finding is in consistence with some studies indicating lysosomal autophagy induced by nanotoxicity [56].

SNP exposure induced hepatocyte nuclear alterations including apoptosis. Some previous studies described

apoptosis as major mechanism of cell death by oxidative stress of nanoparticles [20,57]. This programmed cell death might be resulted from the reduction of ATP production due to the mitochondrial damage induced by SNPs. One study showed that exposure of rats with single intraperitoneal dose of 2 mg/kg SNPs (100 nm diameter) induced apoptosis after 48 h of administration [58]. According to Kim et al. [59], SNPs could induce apoptosis by ROS generation via mitochondrial caspase-dependent pathways. Furthermore, some toxicological studies explained that SNP exposure could induce necrosis and gene alterations [60,61]. In addition, the findings of the current work showed that some hepatocytes of SNP-treated rats demonstrated micronuclei which is in consistence with the report of Song et al. [62]. Micronuclei formation is an indication of DNA damage, ultimately leading to cell death and usually seen in chemical induced genotoxicity.

The results of the present investigation revealed that SNPs could induce multilamellar body formation. Lamellar myelin figures were found to have lysosomal character consisting mainly of phospholipids, glucosylceramide, and lytic enzymes with autophagic activities. Formation of these figures might indicate pathologic hallmark of phospholipidosis and cytotoxicity associated with SNP exposure [63–65].

The present work demonstrated Kupffer cell hyperplasia due to SNP exposure. These cells are hepatic macrophages activated by subtoxic concentration of hepatotoxins and always demonstrated with liver injuries [66]. The activation and enlargement of these cells are always related with stress induced in the hepatic tissues by some drugs and environmental pollutants.

One would conclude from the results of the present study that exposure to SNPs would induce hepatocytes cytoplasmic and nuclear ultrastructural alterations might be associated to disturbance in the pro-oxidant/antioxidant of the hepatocytes system leading to cellular damage and affecting the function of the liver. In addition, the results of present work may raise the concerns about the potential risk on human health that might be related with numerous applications of SNPs. More work is needed to elucidate probable pathophysiological alterations and hepatotoxicity that might be resulted from SNP exposure.

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