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Toxic effect of iron oxide nanoparticles, silver nanoparticles and their mixture on heart, brain and lung of male rats

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Humans are exposed to nanoparticles (NPs) from ambient air and certain workplaces. The data on the potential health hazards of NPs exposure is limited. From the literature, there is not enough data on cardiotoxicity, neurotoxicity, and lung toxicity induced by the co-exposure to iron oxide nanoparticles (Fe2O3NPs) with silver nanoparticles (AgNPs). Therefore, the present study aimed to investigate the toxic effect of Fe2O3NPs, AgNPs alone or in combination on the brain, heart, and lungs of male rats. Animals were divided into 4 equal groups. Group 1 served as control, group 2 was administered orally with Fe2O3NPs (5 mg/kg BW; >50 nm), group 3 was treated intraperitoneally with AgNPs (50 mg/kg BW; >100 nm) and group 4 was administered with the mixture of Fe2O3NPs plus AgNPs. Animals were treated every day for 79 days. The present results showed that at the molecular level Fe2O3NPs, AgNPs, and their mixture showed marked DNA fragmentation as a hallmark of cell death. At the gene expression level Fe2O3NPs, AgNPs, and their mixture showed significant suppression of the mitochondrial transcription factor A (mtTFA) gene, while showing significant induction of peroxisome proliferator activator receptor gamma-coactivator 1α (PGC- 1α) gene. Both genes; mtTFA and PGC- 1α are involved in the regulation of mitochondrial biogenesis and function. Fe2O3NPs, AgNPs, and their mixture caused a significant decrease in final body weight, body weight gain, serotonin, dopamine, acetylcholine esterase, paraoxonase 1, antioxidant enzymes (GST, SOD, CAT, and GPX), total antioxidant capacity, and reduced glutathione in brain, heart, lung, and plasma. Whereas, Fe2O3NPs, AgNPs, and their mixture resulted in a significant increase in norepinphrineiron, acetylcholine, creatine kinase, thiobarbituric acid-reactive substances, nitric oxide, tumor suppressor gene p53, tumor necrosis factor-α, interliukin-6, and lipid profiles. Fe2O3NPs, Ag-NPs, and their mixture showed histology changes alteration in the brain, heart, and lung. In conclusion, the obtained data showed that Fe2O3NPs, and AgNPs alone and in combination induced neurotoxicity, cardiotoxicity, and lung toxicity.

The toxic effects of the combination of Fe2O3NPs with Ag-NPs were more pronounced than each one.

Keywords: Iron oxide nanoparticles; Silver nanoparticles: Male rats; Oxidative stress; Antioxidants; cardiotoxicity, neurotoxicity, and lung toxicity, Cytokines; Mitochondrial transcription factor A peroxisome proliferator activator receptor gamma-coactivator 1α , Biochemical and histology changes.

Recent Publications

- Mosa IF, Abd HH, Abuzreda A, Yousif AB, Assaf N. (2021) Chitosan and curcumin nanoformulations against potential cardiac risks associated with hydroxyapatite nanoparticles in Wistar male rats. International Journal of Biomaterials. Jul 29;2021
- Yousif AB, Mosa IF, Abd HH, Abuzreda A, Assaf N, (2020) Bioevaluation of the role of chitosan and curcumin nanoparticles in ameliorating genotoxicity and inflammatory responses in rats' gastric tissue followed hydroxyapatite nanoparticles' oral uptake. Toxicology Research. Jul;9(4):493-508.
- Yousef MI, Abuzreda AA, Kamel MA, (2019). Cardiotoxicity and lung toxicity in male rats induced by long-term exposure to iron oxide and silver nanoparticles. Experimental and therapeutic medicine. Dec 1;18(6):4329-39.
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Biography

Abdelsalam Abuzreda is a Ph.D. graduate in Nanotoxicity on Molecular and Physiological Characteristics and he is the Assistant Professor and Postdoctoral Research fellow at Benghazi University. Currently, he is a researcher in the Department of Health Safety and Environmental (HSE), Arabian Gulf Oil Company (AGOCO). He has contributed to various conferences and many publications in his research interests such as Material Characterization, Nanoparticle Synthesis, Nanoparticle Preparation, Nanomaterials Synthesis, Nanomaterials, Nanoscience, Nanostructured Materials, Nanobiotechnology, Polymers, and Biomaterials.

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