



MoS₂ nanoparticles induce behavioral alteration and oxidative stress mediated cellular toxicity in the social insect *Oecophylla smaragdina* (Asian weaver ant)

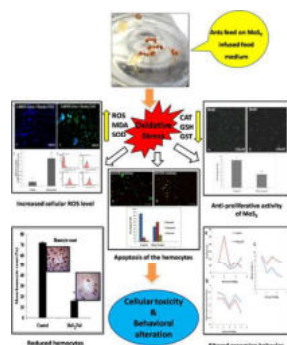


CC Sheeja^a, Anusri Ambali^a, Levna Chacko^b, PM Aneesh^{b,*}, Lekha Divya^{a,*}

^a Department of Zoology, Central University of Kerala, India

^b Department of Physics, Central University of Kerala, India

GRAPHICAL ABSTRACT



ARTICLE INFO

Editor: R. Debora

Keywords:

Insect

MoS₂

Nanoparticle

Behavioral alteration

Oxidative stress

Toxicity

ABSTRACT

The study evaluates molybdenum disulfide (MoS₂) nanoparticles (NPs) induced oxidative stress during cellular toxicity in an invertebrate *in vivo* system, the weaver ant. The lethal concentration was checked and LC₅₀ was obtained as 50 µg/mL. Feeding assay and the photoluminescence activity confirmed the ingestion of MoS₂ NPs by the organism. Behavioral assays showed altered grooming behavior in the MoS₂ NP fed ants. A drastic decrease in the hemocyte count in the MoS₂ NP fed ants revealed the anti-proliferative role of MoS₂. This was further confirmed by 5-bromo-2'-deoxyuridine (BrdU) labeling assay. MoS₂ NPs induced apoptotic activity was also observed in the hemocytes by acridine orange/ethidium bromide (AO/EB) staining. The level of oxidative stress during cellular toxicity was observed. An increased reactive oxygen species (ROS) level was observed in the MoS₂ NP fed ants when compared to the control group. The increased activity of superoxide dismutase (SOD) and the lipid peroxidation (LPO) product were observed. While, the activities of catalase (CAT) and glutathione-S-transferase (GST) and the glutathione content (GSH) were decreased by MoS₂ NPs. The transcript levels of SODs, CAT and GST were up regulated in the treated group. Our results suggest that MoS₂ NPs induced oxidative stress mediates the cellular toxicity in the foragers of the weaver ant.

* Corresponding authors.

E-mail addresses: aneeshpm@gmail.com (A. PM), divyal@cukerala.ac.in (D. Lekha).

<https://doi.org/10.1016/j.jhazmat.2019.121624>

Received 23 August 2019; Received in revised form 19 October 2019; Accepted 5 November 2019

Available online 21 November 2019

0304-3894/ © 2019 Elsevier B.V. All rights reserved.