Metal contents of cement dust contribute to lung disease prevalence in cement factory workers

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Introduction: Exposure to cement dust is one of the most common occupational dust and poses health threats to the public. This has been associated with the development of laryngeal cancer and other metal related diseases. The mechanism of injury to lung cells (Alveolar macrophages and type II epithelial cell) and disease development by this particulate is still unclear.

Objectives: We evaluated the immunotoxicity of two cement dust samples (CDN and CDU) and clinker (CN) using alveolar macrophages and type II epithelial cell relative to their metal contents.

Methods: Metals (chromium, copper, lead, manganese, nickel, cadmium and mercury) were quantified using graphite furnace atomic absorption spectrophotometry, while hexavalent chromium (Cr (VI)) was determined by colorimetric method. Endocytosis of particles was assessed using transmission electron microscope. Additionally, apoptosis (annexin-V-PI), intracellular reactive oxygen species (iROS) generation and reduced glutathione (GSH) were determined using flow cytometry. Tumor necrosis factor-α (TNF-α), interleukin-1β (IL-1β) and macrophage inflammatory protein-2 (MIP-2) secretion from NR8383 were evaluated by ELISA technique.

Results: Our results indicated that Cu, Ni and Mg were significantly higher in CDU relative to CDN. Both total Cr and Cr (VI) were also higher in CDN than in CN. Cadmium was higher in both CDN and CN. Mercury was more in both CDN and CN, while lead (Pb) was only significantly higher in CN. Alveolar epithelial cells internalized clinker predominantly at the membrane bound vacuoles. The CDU induced more apoptosis, intracellular ROS generation (22% higher) and reduced GSH compared with control, which may be related to the significant Cr (VI) level in CDU. Increase in IL-1β and TNF-α secretion were consistent in both CDN and CDU, while MIP-2 was not significantly increased in cells exposed to both CDU and CDN but significant in cells exposed to clinker.

Conclusion: These data suggest that the high metals and Cr (VI) concentrations; known carcinogens may be contributory to the pathologic basis of cement dust toxicity, which may be factory dependent. Endocytosis of cement dust particulates, oxidative stress induced-apoptosis and induction of pro-inflammatory cytokines may be the key mechanisms of cement dust immunotoxicity in lung cells. This study revealed that cement dust exposure is a public health threat to both cement factory workers and people located near such factories.

Biography
Anthony O Okorodudu completed his Doctorate degree in Pathology from University of Medicine and Dentistry of New Jersey and Post-doctoral studies in Clinical Chemistry from Hartford Hospital, CT, USA. He is the Medical Director of Clinical Chemistry Division at the University of Texas Medical Branch and Medical Director of Laboratory Services for the UTMB/Correctional Managed Care System. He has trained 25 Doctoral students and Post-doctoral fellows and several pathology residents. His publication record includes 100+ peer reviewed publication in reputed journals and 11 book chapters. He currently serves on Editorial Board of two scientific journals and he has served on the board of directors for several scientific organizations.

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