

# Investigating the regulatory effect of omega-3 polyunsaturated fatty acid supplementation on the immune imbalance induced by crystalline silica (cSiO<sub>2</sub>) in mice

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## Outline

- I. Introduction
- II. Omega-3 polyunsaturated fatty acid intervention against established autoimmunity in a murine model of toxicant-triggered lupus
- III. Dietary docosahexaenoic acid supplementation inhibits acute pulmonary transcriptional and autoantibody responses to a single crystalline silica exposure in lupus-prone mice
- IV. Conclusion

## Abstract

Systemic lupus erythematosus (SLE) is an autoimmune disease mediated by autoantibodies (AAb), characterized by disruption of immune tolerance and overactivation of B cells, leading to damage to organs. Recent studies have pointed out that eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA) in omega-3 polyunsaturated fatty acids (PUFAs), may have a regulatory effect on immune imbalance. In the first study, the authors used cSiO<sub>2</sub>-induced NZBWF1 mice to simulate the onset of autoimmune diseases. The results showed that DHA intervention can inhibit the production of inflammatory cytokines and regulate the differentiation of B cells, and reduce the production of AABs, indicating that omega-3 can delay the course of lupus. The second study focused on the acute pulmonary immune response after cSiO<sub>2</sub> exposure. High-dose DHA can effectively intervention inhibit the elevation of antibodies and downgrade the antigen presentation of inflammation-related pathways, thereby reducing lung inflammation and immune imbalance.

Combining the two studies, the protective effect of omega-3 is mainly by inhibit B cell differentiation and AABs production, as well as control antigen-presenting cells (APCs) activation and inflammatory gene expression. These results show that omega-3 has potential nutritional adjuvant therapeutic value for autoimmune diseases.

## Reference

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