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**Endothelial Dysfunction and Atherogenesis Are Induced by Chronic Carbon Disulfide (CS<sub>2</sub>) Exposure**

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We hypothesized, that the endothelial dysfunction as result of free radical reactions (oxydative and nitrosative stresses) is the main factors predisposing to 'response-to-injury' and atherogenesis.

**Goal:** to analyze the markers of endothelial dysfunction and lymphocytes chromatin topography at chronic CS<sub>2</sub> exposure.

**Objectives and methods:** 61 worker of viscose industry, chronically exposed by CS<sub>2</sub> more 10 years. The markers of endothelial dysfunction (ET-1, sE- and sP-selectin, sICAM-1 and sVCAM-1 levels) and members of proinflammatory cytokines network (IL-1a, IL-1b, IL-6, IL-8 and IL-10) and their functional response were measured by ELISA. The chromatin of lymphocytes nuclei was studied using Computer TV Morphodensitometry System 'DiaMorph' (Russia) in the smears dyed especially for DNA.

**Results:** Cell coordinate communication disturbances (intracell and intercell) are the basic proinflammatory motif of chronic CS<sub>2</sub> exposure, characterized by atherogenic levels of all studied molecules. So, we detected the prevalence of cerebral atherosclerosis with its specific features: increase of ET-1, sICAM-1, sVCAM-1 and IL-6 levels and their functional response, proapoptotic lymphocytes intranuclear rebuilding and complete apoptotic reactions after additional heat stress at chronic CS<sub>2</sub> exposure.

**Conclusion:** The possible elevated expression of stress proteins is considered to be a universal response to adverse conditions. Obtained results indicate on cerebral (mainly) atherosclerosis development in result of free radicals-induced progression of endothelial dysfunction as main coordinator of cytokines network and cell-cell communications. Theremore, the cell crosstalk on the blood-brain barrier level may present the additional factors, triggering the cell coordination imbalance.

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