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Immunophenotypical Aspects of Peritoneal and Liver Macrophages Derived Animals with the Model of Alloxan Diabetes (Type I) and Their Correction by Sodium Aminodigydrophtalazindione *in vitro*

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In this research morphological and functional characteristics of macrophage cell cultures obtained from different localization in intact animal (IA) and animal with the model of type 1 diabetes mellitus (DM1) were investigated. The research was carried out on the macrophage cell cultures isolated from rat liver and peritoneal cavity. The macrophages were stimulated *in vitro* for 24 and 72 hours with a sodium aminodigydrophtalazindione *in vitro*. Cells, nucleus, cytoplasm area were measured and nuclear cytoplasmic ratio (NCR) were calculated. The phenotype was determined by expression of CD163 (M2-macrophages) and CD80 (M1-macrophages) receptors. Cytokine activity of macrophages was determined by IL-1a, IL-10 μ TNF-level. As a result, the ADPH changes morphometric parameters (a decrease in the size of the nucleus and cells, an increase in NCR) and synthetic cell activity (an increase in IL-10 in macrophages of the peritoneal cavity; IL-1 α and TNF- α in macrophages of the liver) in the first 24 hours of cultivation. ADPH stimulation for 72 hours leads to a decrease in the levels of IL-10, TNF- α and an increase in the level of IL-1 α in all cell populations. ADPH does not affect the expression level of markers of M1 and M2 macrophages.

Keywords: liver macrophages, peritoneal cavity macrophages, sodium aminodigydrophtalazindione

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