The Role of IL-6 in the Immune Response of Glial Cells Against Neospora caninum

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Neosporosis, caused by Neospora caninum, is an important cause of abortion in cattle and neurological diseases in dogs. The systemic immune response to this parasite is characterized by a Th1 phenotype, but in central nervous system (CNS), the resistance to the infection is related to an anti-inflammatory pattern. We have already shown that there is a high production of IL-6 and IL-10 by glial cels infected by N. caninum and parasitemia in these cells became less intensive with an anti-IL-10 monoclonal antibody treatment. This study aimed to evaluate the role of IL-6 in glial cells infected by N. caninum. Astrocytes and microglia cultures from rat brain cortex were treated with anti-IL-6 monoclonal antibody and infected with N. caninum tachyzoites. After 72h of infection, parasite proliferation and cell viability, by Lactate Dehydrogenase (LDH) activity, were evaluated. It could be observed that, with IL-6 blocking, the number of tachyzoites reduces 43.7%, showing that this cytokine contributes to maintain the parasite infection in CNS. Regarding LDH activity, there was no difference in glial viability of infected cells treated or not with anti-IL-6 antibodies, suggesting a weak participation of this cytokines in maintenance of cellular integrity during parasitism. These results demonstrate the importance of IL-6 in the immunopathogenesis of Neoporosis in CNS, and more studies are needed to clarify how N. caninum modulates the production of this cytokine in glial cells.