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IMMUNE STATUS ON HAEMODIALYSIS PATIENTS AFFECTED WITH COVID19 INFECTION

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BACKGROUND AND AIMS: End-stage renal disease patients on haemodialysis (HD) seem more likely to develop severe COVID19 disease. Over the course of COVID disease, we observed a poor tolerance to HD sessions with a marked tendency of clinical deterioration over them.

The objective is to evaluate changes on immunological system over HD session on patients affected with COVID19 compared with patients without COVID19.

METHOD: Fourteen HD patients were studied including 9 confirmed COVID19 infection and 5 healthy controls. Predialysis and postdialysis blood samples were compared to study alterations on immune status. We identified cytokines by Luminex (CCL2, CXCL10, IL1Ra, IL10, IL12p70, TNF α , IL17Ra, IL6, IL7) and adaptive lymphocyte subsets (CD4/CD8 naïve, CD4/CD8 MC, CD4/CD8 MP, CD19, CD56). Monocyte subsets (CD14+CD16-, CD14+CD16+, CD14-CD16+) were detected from peripheral blood mononuclear cells (PBMC), as well as immune activation (CD11b, HLA-DR, CD86) and migration factors (CCR2, CCR5). The supernatant of isolated CD14+ cells after 4-hour stimulation with LPS where analysed by Luminex to measure cytokines (CCL2, CXCL10, GM-CSF, IL10, IL12p70, IL17Ra, IL6, IL7, TNF α).

RESULTS: Patients with COVID19 presented predialysis: (1) higher plasmatic levels of IL12p70, TNF α e IL7, (2) lymphopenia and neutrophilia, (3) higher percentage of intermediate monocytes and lower of non-classical, (4) lower membrane expression of CCR2, HLA-DR y CD86 over Cd14+ cells, and (5) higher production of CCL2, GM-CSF, IL10, IL12p70 y IL17Ra by LPS stimulated monocytes compared with patients without COVID19. When analysed the fold-change between pre and postdialysis values, patients with COVID19 infection present a: (a) higher plasmatic levels of IL6, IL1Ra, CCL2 e CXCL10, (b) reductions of total lymphocytes, (c) higher membrane expression of CCR2, CD33 y CD86 on CD14+ cells, and (d) higher production of TNF α , GM-CSF, IL10, IL17, IL6 e IL7 by LPS stimulated monocytes compared with patients without COVID19. No differences on lymphocyte subset were found.

CONCLUSION: The clinical deterioration on COVID19 infected patients over HD session could be related with monocyte activation and pro-inflammatory cytokines secretion.