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**Title:** BALF TNF $\alpha$  level in relation to inflammatory status and phenotype of sarcoidosis

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**Body:** Determining the inflammatory status of the disease may be helpful in predicting the prognosis and responsiveness to treatment. Aim: To evaluate the relation of BALF TNF $\alpha$  levels to the inflammatory status and phenotype of sarcoidosis. Material and methods We have measured: TNF $\alpha$ , TGF $\beta$ , IL2, IL2R, IL12, IL10 in BALF and TNF $\alpha$ , TGF $\beta$ , IL2, IL12, IL10 levels in serum in 184 sarcoid patients. We checked for the correlations between BALF TNF $\alpha$  and cytokines, selected systemic inflammatory markers and clinical factors (age, duration of the disease, PFT, 6MWT, extrapulmonary involvement). Results We have found important positive correlation between concentration of BALF TNF $\alpha$  and: - BALF: IL2R(r=0,4676, p=0,0001), IL12 (r=0,4545, p=0,0001) and serum: IL12 (r=0,26, p=0,001), IL10 (r=0,2832, p=0,001), TNF $\alpha$  (r=0,1871, p=0,12). - serum CRP(r=0,2427, p=0,003),  $\gamma$ globulins(r=0,1648, p=0,033), Ddimers(r=0,1685, p=0,035). - the age of patients (r=0,16, p=0,029), No relation was found between the BALF TNF $\alpha$  and the duration of the disease, PFT (except for FEV1 r=-0,1464, p=0,05), 6MWT, extrapulmonary sarcoidosis (liver and spleen dimensions, hepatic enzymes, protrombine), Ca and P serum nor urine levels. Weak but important negative correlation was observed between BALF TNF $\alpha$  and monocytosis(r=-0,1788, p=0,015). Conclusion In our group of patients: The BALF cytokine inflammatory status reflects the systemic inflammation (measured by the serum cytokine network and non-specific inflammatory biomarkers). The BALF TNF $\alpha$  status has no relation to the phenotype of sarcoidosis.