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Cardiorespiratory fitness is inversely associated with the inflammatory status in the general population

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Background/Introduction: Low-grade systemic chronic inflammation is a major risk factor for cardiometabolic diseases. Higher physical activity has athero- and cardioprotective effects, potentially through its anti-inflammatory properties. However, the relation between resting inflammatory status and cardiorespiratory fitness (CRF) in population-based settings remains to be elucidated. While previous research has shown inverse associations between CRF and high-sensitive C-reactive protein (hs-CRP) levels, relations with other inflammatory markers are not yet well examined. Purpose: We assessed the relation between markers of CRF (i.e. maximal oxygen uptake [VO2peak], oxygen uptake at the anaerobic threshold [VO2@AT], oxygen pulse [O2HRmax] and maximal workload [max. Watt]) and hs-CRP as well as total white blood cell (WBC), monocyte, neutrophil, lymphocyte, eosinophil and basophil counts.

Methods: Data of the population-based cohort Study of Health in Pomerania (SHIP) was used (n=1,481; 51% male; age range 20–81 years). CRF was assessed using standardized cardio-pulmonary exercise testing (CPET) according to a modified Jones protocol. The Dimension Vista 500 analytical system was used to measure hs-CRP concentrations. Fluorescence-activated cell sorting was used to assess total as well as subpopulation WBC count. We excluded subjects with missing data, anti-rheumatic/steroid/anti-inflammatory medication as well as chronic inflammatory and hepatic diseases, gastritis, hepatitis infection, severe renal dis-

ease, chronic lung disease, asthma, chronic bronchitis, previous myocardial infarction, left ventricular ejection fraction <40% and previous cancer. Linear regression models, adjusted for age, sex, current smoking and leanmass, were used to assess the relation between CPET and inflammatory parameters.

Results: A 14 ml/min (95%-confidence interval [CI] -23 to -6, p=0.004) lower VO2peak was associated with a one mg/L higher hs-CRP. A lower VO2peak was also related with a one Gpt/L greater WBC (β : -42. 95% CI: -55 to -27 ml/min, p<0.001) as well as neutrophil (β : -57, 95% CI: -77 to -36 ml/min, p<0.001), lymphocyte (β : -63, 95% CI: -107 to 19 ml/min, p=0.005) and monocyte count (β : -204, 95% CI: -360 to -47 ml/min, p=0.011). No significant associations were found for eosinophil and basophil count. Similar results were found for VO2@AT, O2HRmax and max. Watt.

Conclusion: Results suggest an inverse association between CRF and resting inflammatory status. Our non-significant findings with regards to eosinophils and basophils may be expected since these cell types are generally involved in type 2 immune responses related to allergic skin and airway inflammation as well as immunity against parasites. Overall, our results imply that potential biological mechanisms underlying the atheroand cardioprotective effects of high CRF may be related to lower chronic inflammation in fitter individuals.