Screening of yo T cells in breast cancer patients

Chiara Massa 1,2, Anja Müller 2, Katharina Biehl 2, Barbara Seliger 1,2

Breast cancer (BC) is the most common cancer among women worldwide and a leading cause of death. In order to improve BC patients' outcome, different immunotherapeutic strategies are currently tested, for which predictive and prognostic biomarkes are urgently needed. T cells carrying the gammadelta ($y\delta$) T cell receptor (TCR) represent a minor population of T cells, which are increasingly being evaluated for tumor immunotherapy. Indeed, they can directly recognise a variety of biomolecules, like several lipids and protein antigens, without the requirement of antigen processing and presentation by classical MHC class I molecules and display cytotoxic activity against malignant cells. Analogous to the conventional αβ T cells, yδ T cells exist in different polarisations, thus their evaluation as a "pan" population results in both positive and negative correlation with patients' outcome depending on the clinical setting. Therefore, the aim of this study is to perform a deep characterisation of the frequency and phenotype of yδ T cells within the peripheral blood of patients with different forms of BC and to correlate these data with the patients' clinical outcome and therapy response. A CyTOF panel including the major V_γ and Vδ chains, together with innate receptors, functional molecules, activation markers and different immune checkpoint inhibitor receptors have been established to perfom such characterisation. The acquired knowledge will allow a more rational use of $y\delta$ T cells for the therapy and / or stratification of BC patients.

¹ Institute for Translational Immunology, Brandenburg Medical School Theodor Fontane, Brandenburg an der Havel, Germany

² Institute for Medical Immunology, Martin Luther University Halle-Wittenberg, Halle (Saale), Germany