Heart Failure & Arrhythmias









Focus of research group (I)

Name PI: Esther Lutgens (e.lutgens@amc.uva.nl) Department, UMC, Medical Biochemistry, AMC Size of research group: 8 PhDs, 4 PD, 1 staff, 3 techs (2 fte)

Current mission, vision and aims

To better understand the role of the immune system in atherosclerosis and the metabolic syndrome.

Discover novel immunotherapeutic targets for CVD, design and test potential therapeutics.

Focus on immune checkpoints (**CD40, GITR, CD27, CBL-B**) and their signaling intermediates (**TRAFs, NFkB**)











Focus of research group (II)

Current expertise

Mouse models of atherosclerosis, diet induced obesity, EAE Immune system: FACS, sorting, cell culture, luminex Histology and morphometry, pathology Drug design: in collaboration with G. Nicolaes (UM)

Collaboration in ACS

Menno de Winther Mat Daemen Stephan Huveneers Erik Stroes, Jeffrey Kroon Ex ACS: Willem Mulder, Niels van Royen

Current funding ERCcon, VICI, GENIUS II (CVON)











Future plans

Short term (1-2 year) plan

Cell type specific functions of CD40 in atherosclerosis and obesity CD40-TRAF6 compound: first in human Validation of CD40-TRAF2 compounds Cell type specific role for CBL-B in CVD, design of CBL-B agonist Side effects of cancer immunotherapy

Long term (>2 year) plan

Identify the immune (checkpoint) landscape in hyperlipidemia in murine models and humans: regulators, novel checkpoints, therapy.

- CyTOF
- (Single cell) RNAseq
- Metabolomics

Checkpoint ATHERO (leDucq) TURNOVER (Zwaartekracht) Cardio-oncology (Tom Seijkens)