



**RESEARCH TOPIC-MEM8
IMMUNE AND GENETIC MECHANISMS OF HEART FAILURE**

Curriculum MEM Clinical

Clinical Unit name and address: Department of Cardiovascular Medicine, Humanitas Clinical and Research Center-IRCCS

Laboratory name: Molecular Cardiology, Humanitas University

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Abstract

Our group has been studying the molecular mechanisms of cardiovascular diseases, in particular heart failure and atherosclerosis. In our laboratory we try to approach the complexity of these diseases through an integrative approach aimed at understanding the genetics and epigenetics beneath the pathological processes. More recently, we started defining the relevance of immune mechanisms underlying cardiovascular diseases through single cell sequencing and FACS analysis approaches, focusing in particular on the interaction between the immune system cells and other parenchymal cells of the heart and in the arteries in disease. We plan to create a clinical database of human cardiovascular diseases integrated with human specimen correlates in order to translate information gathered from basic science studies. The role of the immune system and of genetics will be studied in human cardiovascular diseases

Main technical approaches

FACS analysis of immune cells; single cells genetic analysis through RNA sequencing and bioinformatics; human genetic technologies (HT-DNA sequencing and SNP analysis); deep clinical and imaging phenotyping; human biobanking and clinical data-base integration

Scientific references

- 1.Papait R, Serio S and Condorelli G: The role of the epigenome in heart failure, *Physiol Rev*,10.1152/physrev.00037.2019
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3.Salvarani N, Crasto S, Miragoli M, Paulis M, Kunderfranco P, Forni A, Dal Ferro M, Sinagra G, Vezzoni P, Faggian G, Condorelli G*, Di Pasquale E: Lamin A/C mutations induce cardiac conduction defects through an epigenetically-mediated reduction of sodium currents: a study in a iPSC-derived model of cardiac laminopathy, *Nature Comm*, May 22;10(1):2267. doi: 10.1038/s41467-019-09929-w

Type of contract

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