Direct reprogramming of human cardiac fibroblasts to cardiomyocytes using microRNA mimics

C. Paoletti¹, C. Divieto², F. Di Meglio³, D. Nurzynska³, V. Chiono¹ Presenting Author: Camilla Paoletti, camilla.paoletti@polito.it

¹Department of Mechanical and Aerospace Engineering, Politecnico di Torino, Turin, Italy ²INRIM, Metrology for Life Quality, Turin, Italy ³Department of Public Health, University of Naples "Federico II", Naples, Italy

INTRODUCTION: The combination of four different microRNAs (miR-1, 133, 208 and 499), named "miRcombo", has been used for the direct reprogramming of murine fibroblasts into cardiomyocytes (CMs) for myocardial infarction (MI) treatments.[1,2] Here, we evaluated miRcombo mediated reprogramming of human adult cardiac fibroblasts (AHCFs) into CMs in 2D and 3D culture.

METHODS: For digital droplet PCR (ddPCR) analysis, 3x10⁵ AHCFs were plated in 6-well plates, for Immunocytochemistry (ICC) 5x10⁴ cells were plated in 24-well plates. AHCFs were transfected with miRcombo (mirVana) using DharmaFECT1 (Dharmacon). Untrasfected and NegmiR (mirVana) transfected AHCFs were used as controls. After 24 hours, medium was changed to medium with 1 μM Jak1 Inhibitor for 4 days for 2D experiments; for 3D experiments, cells were cultured in fibrin-based hydrogels.

RESULTS & DISCUSSION: ddPCR analysis showed significant increase expression of early cardiac transcription factors (TFs) Hand2 and Mef2c (p < 0.005) slight increase expression of Tbx5 and Nkx2.5, although non-significant (p > 0.05), and reduced Vimentin expression (p < 0.05) in miRcombo-transfected AHCFs compared to controls after 4 days in 2D culture. ICC analysis showed increased expression of late cardiac markers α -sarcomeric actinin and cTnT in miRcombo-transfected AHCFs after 10 and 20 days of culture in 2D. However, ddPCR showed no significant differences of late cardiac markers Myh6 and cTnI expression between the groups after 15 days in 2D culture. On the other hand, cells cultured in 3D fibrin-based hydrogels showed enhanced cardiac TFs expression compared to 2D. However, miRcombo transfection did not significantly enhance cardiac gene expression in AHCFs cultured in 3D hydrogels respect to controls after 4 days. After 15 days, AHCFs cultured in 3D hydrogels showed a strongly enhanced expression of cardiac genes such as cTnI and Myh6 compared to 2D.

CONCLUSIONS: Together, these results showed that a 3D environment was found to play a key role in enhancing direct reprogramming of AHCFs into CMs.

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