

Supplemental figures

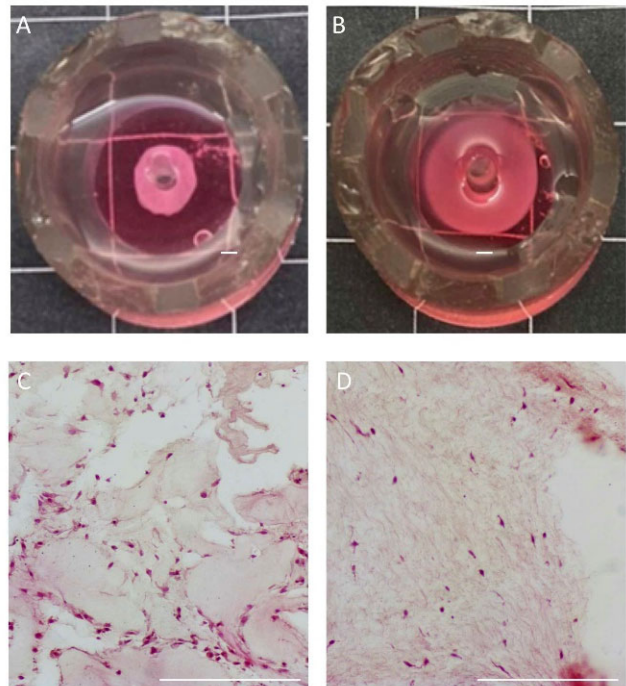
Culturing of cardiac fibroblasts in Engineered Heart Matrix reduces myofibroblast differentiation but maintains their response to cyclic stretch and transforming growth factor $\beta 1$

Meike C. Ploeg¹, Chantal Munts¹, Tayeba Seddiqi¹, Tim J.L. ten Brink², Jonathan Broomhaar³, Lorenzo Moroni², Frits. W. Prinzen¹ and Frans. A. van Nieuwenhoven^{1*}

¹Department of Physiology, Cardiovascular Research Institute Maastricht (CARIM), Maastricht University, the Netherlands, ²MERLN, Institute for Technology-Inspired Regenerative Medicine, Maastricht University, the Netherlands, ³MosaMeat, Maastricht, the Netherlands.

*Correspondence: f.vannieuwenhoven@maastrichtuniversity.nl

Supplemental Figure S1. Effect of initial collagen concentration on EHM compaction. EHM-rings using 1 mg/ml (A) or 3 mg/ml (B) initial collagen concentrations (2000 cells/ μ L) in 250 μ L gel after 24 h. Images are taken on the black and white grid of 1 x 1 cm. Shown is the silicone mold, the central pole containing the ring in light pink tones, surrounded by CFMM. Bars represent 2 mm. HE staining of the EHM-ring after 24 h using 1 mg/ml (C) and 3 mg/ml (D) initial collagen concentrations (2000 cells/ μ L in 250 μ L gel) showing the cell density difference between the two collagen concentrations. Magnification 20x. Bars represent 200 μ m.



Supplemental Figure S2. Effect of initial collagen concentration (1 or 3 mg/ml) on RNA yield per cell (pg/cell) and gene expression of Acta2, Ctgf and Tgf $\beta 1$ of the EHM-rings after 24 h of culture (2000 cells/ μ L in 250 μ L gel) (n=2). Bar represents mean.

