Reduction of Cardiac Fibrosis by Interference With YAP-Dependent Transactivation

Strain-sensitive fibrotic programming of human cardiac stromal cells can be reverted by interfering with YAP-dependent transcriptional activation

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Abstract

Background - Conversion of cardiac stromal cells into myofibroblasts is typically associated with hypoxia conditions, metabolic insults, and/or inflammation, all of which are predisposing factors to cardiac fibrosis and heart failure. We hypothesized that this conversion could be also mediated by response of these cells to mechanical cues through activation of the Hippo transcriptional pathway. The objective of the present study was to assess the role of cellular/nuclear straining forces acting in myofibroblast differentiation of cardiac stromal cells under the control of YAP transcription factor and to validate this finding using a pharmacological agent that interferes with the interactions of the YAP/TAZ complex with their cognate transcription factors TEADs, under high-strain and pro-fibrotic stimulation.

Methods - We employed high content imaging, 2D/3D culture, atomic force microscopy mapping and molecular methods to prove the role of cell/nuclear straining in YAP-dependent fibrotic programming in a mouse model of ischemia-dependent cardiac fibrosis and in human-derived primitive cardiac stromal cells. We also tested treatment of cells with Verteporfin, a drug known to prevent the association of the YAP/TAZ complex with their cognate transcription factors TEADs.

Results – our experiments suggested that pharmacologically targeting the YAP-dependent pathway overrides the pro-fibrotic activation of cardiac stromal cells by mechanical cues *in vitro*, and that this occurs even in the presence of pro-fibrotic signaling mediated by TGF-β1. *In vivo* administration of Verteporfin in mice with permanent cardiac ischemia reduced significantly fibrosis and morphometric remodeling but did not improve cardiac performance.

Conclusions - Our study indicates that preventing molecular translation of mechanical cues in cardiac stromal cells reduces the impact of cardiac maladaptive remodeling with a positive effect on fibrosis.

Non-standard abbreviations and acronyms

YAP: Yes-associated protein

TAZ: Transcriptional co-activator with PDZ-binding motif

TEAD: TEA domain transcription factor

TGF-β1: Transforming growth factor beta-1

ECM: Extracellular matrix

MI: Myocardial infarction

LV: Left ventricle

VTP: Verteporfin

CF: cardiac fibroblast

CM: Cardiomyocyte

Introduction

Maladaptive remodeling of the myocardium is one of the earliest hallmarks of heart failure (HF). This is characterized by inflammation and a progressive fibrosis that leads to replacement of the parenchyma with a stiff, fibrotic, tissue[1]. In the adult healthy myocardium, the turnover of the extracellular matrix (ECM) is controlled by stromal interstitial cells, which can be mapped into several different phenotypes based on transcriptional and functional features[2-4]. Under conditions causing ventricular mechanical overload (e.g. hypertension), in the presence of metabolic alterations such as hyperglycemia, injury such as hypoxia, or just depending on the aging process, resident un-activated stromal cells evolve into pro-fibrotic cells, the so called myofibroblasts (Myo-Fbs) that contribute to inflammation, altered ECM accumulation, and myocardial stiffening[5-10].

The term mechanosensation refers to the ability of the cells to sense the physical characteristics of the surrounding environment through the activation of intracellular signaling cascades elicited by mechanical cues[11]. The Hippo signaling pathway, an essential component of the machinery translating cell mechanical responses into discrete transcriptional activation is well contextualized in cancer biology as a *primus movens* in cancer stem cells determination, metastatic activity, drug resistance and cell plasticity[12]. A direct connection between static/dynamic mechanical cues and transcriptional activation of downstream targets has been established with the finding that cytoskeletal tensioning resulting from cellular adhesion to extracellular components with specific geometric arrangement and viscoelastic properties[13] translates into reversible nuclear shuttling of the main Hippo transcriptional component, the YAP/TAZ complex[14-16]. In addition to promoting shuttling from the cytoplasm to the nucleus of the YAP/TAZ complex by inhibiting the Hippo kinase pathway, the acto-myosin cytoskeleton also directly forces the YAP/TAZ into the nucleus by physically deforming the nuclear lamina and opening the nuclear pores[17]. The relevance of the Hippo signaling for cardiac biology has emerged from studies highlighting the role of YAP in neonatal myocytes proliferation and cardiac regeneration[18-22]. Moreover, specific deletion of upstream components of the Hippo pathway Lats1/2 in cardiac fibroblasts determines a permanent activation into Myo-Fbs[23, 24], strongly suggesting implication of the YAP/TAZ complex in homeostatic control of cardiac matrix, and a specific function in myocardial remodeling after injury. In keeping with these evidences, elevated levels of nuclear YAP have been found in infarcted hearts in mice[25], and blockade of YAP/TAZ complex activity with Verteporfin (VTP), a drug that interferes with the binding of the complex to TEADs DNA binding proteins, attenuates injury-dependent cardiac fibrosis[26, 27]. The link between the abnormal distribution of the straining forces in the infarcted heart and activation of the YAP-dependent transcriptional activation is, however, not clear. It is also not clear whether human myocardial stromal cells respond directly to strain forces with activation of YAP-dependent pro-fibrotic signaling. In the present study, we provide evidences that the straining forces acting upon remodeling of the myocardial matrix after infarction activate the Hippo transcriptional pathways through a mechanism determining nuclear straining and preferential cellular polarization in the infarct scar. We also show that cardiac stromal cells (cSt-Cs) activate a pro-fibrotic pathway in response to sensing of tissue compliance and geometry, thus linking YAP nuclear translocation to the force-dependent matrix remodeling process occurring

during the cardiac healing after infarction. This process can be reverted by treatment with VTP even under high-strain conditions and in the presence of TGF-β1, a key paracrine stimulator of cardiac fibrosis[28]. These findings were obtained in mice carrying permanent ligature of the left anterior descending coronary, in which treatment with VTP reduced fibrosis at short- and long-term after ischemia.

Methods

Ethics

Experimental investigations involving human-derived cells and tissues were approved by the local ethical committee at Centro Cardiologico Monzino, IRCCS (approval date: 19 May 2012 and subsequent renewal on 16 May 2016) and Policlinico San Donato, IRCCS (protocol 2438, 27/01/2009 and CE 85/int/2016 9/6/2016). Collection of material and experiments were performed in compliance with the Declaration of Helsinki. Patients were required to sign an informed consent to donate small fragments of cardiac tissue (right atrial appendage), during coronary bypass grafting or aortic valve substitution interventions. Experiments in mice were conducted in keeping with the guidelines from Directive 2010/63/EU of the European Parliament on the protection of animals used for scientific purposes, and in accordance with experimental protocols approved by the University Committee on Animal Resources at the University of Milan (668-2015). The protocol of VTP *in vivo* administration in mice with permanent MI was approved by the Italian Ministry of Health (Authorization number 861-2021) and were executed in an authorized facility.

Mouse model of myocardial infarction (MI)

A mouse model of permanent ligature of the left anterior descending coronary artery (LAD) was employed to create a myocardial infarction. Mice were sacrificed at short (3 days), intermediate (7 days) and long (4weeks) follow-up times, after which hearts were prepared for histological analysis. The protocol of *in vivo* treatment of mice with VTP was performed by administrating VTP dissolved in corn oil at a concentration of 50 mg/kg every third day. Monitoring of the treatment effect was performed by echocardiography at specific steps during the observation period. At the end of the treatment, the hearts were harvested and processed for histology. For further information on animal procedures see the extended methods available in the supplementary material.

Microfabrication methods

For the manufacturing of the controlled stiffness gels with polyacrylamide (PAA), we used a protocol that was previously employed by us[29]. For investigating the traction forces generated by single cells, the micropillar method was adopted[30, 31]. The manufacturing of the micropatterned substrate of poly(dimethylsiloxane) (PDMS, Sylgard 184, Dow Corning, United States), a dedicated silicon mold with a negative pattern of the micropillar arrays was designed using the software Solidworks (Dassault Systemes, Vélizy-Villacoublay, France). Further details about design and fabrication of the microfabricated structures are provided in the extended methods available in the supplementary material.

Primary cell culture, human tissue harvesting and main immunofluorescence methods.

Primary human cardiac stromal cells were derived from "cardiospheres" as previously described[32, 33]. Briefly, cellular outgrowths of small myocardial explants were sub-cultured to obtain cardiospheres and cardiospheres-derived primitive cSt-Cs. Cardiospheres and cSt-Cs were used for immunofluorescence staining and high content image analysis using CARE, an algorithm originally developed by us to analyze microscopic images[34]. Controls of immunofluorescence/histochemistry included samples stained with pre-immune antibodies/sera and also secondary antibody-only stained samples to determine the background level. For exhaustive information on cell culture methods and cell analyses see the extended methods. For the observations on human myocardium, left ventricle cardiac biopsies were harvested from patients affected by dilated hypokinetic ischemic cardiomyopathy undergoing surgical ventricular reconstruction procedure. Formalin-fixed paraffin-embedded consecutive serial sections (1 µm thickness) of the border zone samples (peri-infarct area) were dewaxed and hydrated through graded decrease alcohol series and stained for histological analysis. Hematoxylin/Eosin staining (Mayer's Hematoxylin/Eosin; Bio Optica, Milan, Italy) and Masson Trichrome staining (Masson trichrome with aniline blue Bio-Optica) were carried out using standard protocols. The microscopic images contained in the figures are those that best represent the observed phenotypes.

Atomic force microscopy

Force spectroscopy measurements were performed with a commercial AFM mounted on an inverted microscope using a borosilicate glass microsphere of about 18 µm in diameter manually glued at the end of the cantilever. Cell rigidity was evaluated in 30 randomly chosen cells for each condition, acquiring three force spectroscopy curves in the center of each cell nuclei. Further details are provided in the online supplementary material.

Molecular methods

For Western blotting analyses, cells were lysed in standard sample buffer and run (30µg) into 4-12% gradient SDS polyacrylamide gels. Proteins were transferred onto nitrocellulose membranes followed by incubation with the indicated primary and secondary antibodies (see Major Resource Table). For RNA-sequencing analysis, total RNA was extracted from 6 independent cSt-Cs cultured ±TGF-β ±VTP using TRIzol. After quality checking and quantification, the Poly(A)$^+$ RNA was enriched and then processed for RNAseq. Differential gene expression analysis was performed using the R software. Validation was performed in independent RNA samples of cSt-Cs by RT-qPCR. Indications about the primers sequences and reagents are are provided in the Major Resource Table. The effects of TGF-β1±VTP on cSt-Cs contraction were determined using a cell contraction assay kit (Cell Biolabs), as per the Manufacturer's instructions. Collagen plug areas were measured using ImageJ software. Soluble collagen release was quantified using Sircol colorimetric assay (Biocolor) on conditioned medium of cSt-Cs. A full description of the methods employed in molecular analyses is provided in the online supplementary material.

Statistical analyses

Data were graphed and analyzed using the Prism Graph Pad statistical software. Normality tests on normal/log-normal distributions were run using the Shapiro-Wilk methods fixing $\alpha = 0.01$. The number of replicates, the

167 *P* values, and the type of tests that were used for statistical comparisons are specified by the dots/numbers over
168 the histograms in the figures and the figure legends, respectively.

Results

Orientation- and cell strain-dependent activation of YAP in the infarct scar

To assess the activation of the YAP-dependent transcriptional pathway in the context of the late myocardial remodeling occurring after ischemia, we performed immunohistochemistry on hearts of mice with permanent ligature of the left coronary artery at early (3 days) and late (4 weeks) follow up[35]. **Figure 1A** shows the results of YAP immunolocalization in transversal sections of the infarcted heart at three days post-MI. Cells with the morphology of monocytes/macrophages invading the infarct area exhibited, in line with the literature[36], high levels of the protein in the nucleus. Cells showing nuclear-localized YAP included also putative myofibroblasts and cardiomyocytes (CMs). In the remote zone, a reduced number of cells were positive for YAP (independently of nuclear/cytoplasmic localization), suggesting an ischemia- and or inflammatory-dependent activation of the pathway in the infarcted region. At 4 weeks after infarction, infarct scars with abundant collagen deposition were clearly visible (**Figures 1B**). In this fibrotic tissue, collagen bundles and cells with Myo-Fbs morphological characteristics were clearly recognizable with a preferential equatorial/circumferential orientation (see insets in the figure). Analysis of YAP expression in the fibrotic tissue by immunohistochemistry revealed cells with Fibroblasts (Fbs) morphology bearing high levels of YAP nuclear localization (YAP^+) or negative for expression of the transcription factor (YAP^-) (**Figure 1C**). Examining the IHC images at high magnification, it came to our attention that YAP^+ cells were not evenly distributed in the collagen-rich matrix, but exhibited a preferential nuclear orientation along the equatorial cutting plane of the left ventricle (LV), suggesting a relationship between nuclear translocation of the transcription factor and strain forces transmitted within the circumferentially arranged collagen fibers (**Figure 1B**)[37]. To quantify this finding we determined the nuclei orientation and deformation in these cells using CARE, an algorithm that we previously tailored to perform automatic segmentation of microscope images[34] (**Figures 1C, D, S2**).We then correlated these values to the presence of nuclear YAP. **Figure 1E** contains the results of this analysis, showing a significantly higher nuclear deformation and a more frequent equatorial alignment of YAP^+ compared to YAP^- cells. This suggests that strain forces, known to direct anisotropic collagen deposition in the forming scar[37], are also involved in the activity of the YAP/TAZ transactivation pathway connected to nuclear deformation and equatorial alignment of cells with CFs morphological characteristics in the fibrotic areas[17]. These results were validated in myocardial tissue of patients with severe heart failure undergoing surgical ventriculoplasty. As shown in **Figure 1F**, the interstitial fibrosis, well evident in the tissue, was associated with presence of cells with YAP staining in the nucleus, suggesting activity of the pathway in cells of the fibrotic scar.

YAP nuclear translocation is mechanically regulated by cytoskeleton tensioning in human cardiac stromal cells and is correlated to expression of fibrotic markers

To prove that YAP nuclear translocation is mechanically connected to cytoskeleton tensioning in cells with the ability to differentiate into Myo-Fbs, we derived cardiac stromal cells from human primary explant cultures from human right atrial fragments[32], selected through spontaneous spheroid growth (the so-called

'cardiosphere')[38] for a primitive mesenchymal phenotype. In line with the existing literature[39], cardiosphere-derived cells exhibited a variety of mesenchymal markers including CD29, CD44, CD90 and CD105, but not endothelial markers CD31 and CD144 (**Figure S2**). We first employed 2D poly-acrylamide gels (PAGs) with a discrete stiffness in the range of 17 - 58 kPa (Young's elastic modulus, detected by atomic force microscope - AFM)[29] and, as a reference, glass surfaces to promote cell attachment onto Fibronectin. Results of cytoskeleton staining by phalloidin-TRITC, revealed a decrease in stress fibers polymerization and consequent changes in cellular forms factors (area and circularity) onto softer gels (**Figure 2A**). To unravel the role of environmental mechanics on YAP-dependent nuclear signaling[15, 16], we performed YAP staining of the cells plated onto PAGs followed by quantitative evaluation of the cells with nuclear-localized YAP ($nYAP^+$ cells) and of the cytoplasmic/nuclear YAP signal ratio (**Figure 2B, S3**). As expected[16, 40], results showed a stiffness-dependent trend of YAP to be confined in the nucleus, and this was also directly connected to cell proliferation, as detected by labeling cells with Ki-67 antibody (**Figure 2C**).

Since the changes in proliferation and YAP nuclear/cytoplasm ratio observed in cells plated onto PAGs with controlled stiffness could be due to variations in cytoskeleton polymerization/tensioning[17] we explored the possibility that cSt-Cs in a 3D structure could sense geometric or positional information translating into discrete YAP nuclear translocation and activation of pro-fibrotic signaling. To validate this hypothesis, we analyzed the distribution of YAP localization inside the cardiospheres. These structures have been in fact described as 3D models of cardiac 'niches' with undifferentiated cells located in the 'core' of the sphere and more differentiated cells with mesenchymal characteristics located in the external shell[32, 41, 42]. **Figures 3A-B** show, respectively, the structure of the cardiospheres and the localization of YAP protein and Ki67 proliferation marker in the 3D structures. By employing CARE[34] to analyze these structures in high resolution images serially acquired along the vertical axis of the spheroid cell clusters (*z*-stack), we were able to clearly distinguish in the 3D volume an external 'shell' of the cardiospheres, defined as the outer volume encompassing one-third of the radius of the spheroids, where cells with intra-nuclear YAP and Ki67 localization were preferentially localized, and an inner 'core' where YAP was mainly cytoplasmic and Ki67 signal was absent (**Figures 3A, S4**). As a first control in this experiment, we used immunostaining with antibodies specific for GATA-4, a cardiac-specific transcription factor that is not known to be mechanically regulated and, indeed, showed no preferential nuclear localization neither in the shell nor in the core of the cardiospheres (**Figure 3C**, **S4**). Second, given that the asymmetric distribution may simply reflect from possible metabolic stress of the cells in the core, due to limited distribution of nutrients and oxygen, we cultured cSt-Cs in hypoxia, low glucose and low serum conditions, and various combinations thereof. As shown in **Figure S5**, none of these conditions affected the YAP nuclear distribution in the cells, suggesting that asymmetric YAP nuclear localization results from a differential response of the cells to compression forces resulting from topological cues. Finally, since YAP nuclear translocation is directly connected to cell mechanics-controlled proliferation and pro-fibrotic activation[43], we assessed the distribution of Ki67 and markers typically expressed in pro-fibrotic cells in the heart (Collagen I and αSMA) in the cardiospheres. Results showed an overlapping between the regions of the cardiospheres where nuclear YAP, Ki67 and fibrotic

markers were more frequently detected (**Figures 3B, D, E**), suggesting a cause relationship between topological activation of YAP signaling and differentiation of cSt-Cs in pro-fibrotic cells.

To establish a direct relationship between YAP localization and transmission of forces from the cytoskeleton to the nucleus, we investigated the nuclear ellipticity of cSt-Cs plated on 2D gels with differential stiffness and, in parallel, in the 3D cardiospheres volume. As shown in **Figures 4A, B**, there were significantly more round nuclear shapes in cells plated onto gels with low stiffness and in the core of the cardiospheres compared to gels with higher stiffness and the outer cardiospheres layer. We then directly interfered with the polymerization of stress fibers by treating cSt-Cs plated onto glass (stiffness in the MPa range) with Blebbistatin (BB) and Y27632[44, 45]. Although these two compounds inhibit cytoskeleton tensioning with different mechanisms, both determined a completely reversible effects on stress fibers generation and nuclear YAP localization (**Figure 4 C, D**). To assess whether nuclear straining by cytoskeleton tensioning is directly associated with modifications in nuclear geometry and mechanics we used confocal imaging and AFM. Results of these analyses showed that both inhibitors reversibly increased the dimension along the *z*-axis of the nuclei of cSt-Cs, and determined a relaxation of their chromatin as detected by a decrease in nuclear stiffness (**Figure 4E**). Finally, to contextualize the results in the framework of the YAP-dependent transactivation, we analyzed the expression of three YAP canonical targets (*CTGF*, *CYR61*, *ANKRD1*) in cells treated with BB or Y27632 using RT-qPCR (**Figure 4F**). This experiment showed a completely reversible reduction in the expression level of the three genes *vs.* controls. This establishes a connection between the expression of *pro*-fibrotic genes and strain-dependent YAP transcriptional signaling in human cSt-Cs.

Override of the pro-fibrotic mechanical programming of human cSt-Cs by Verteporfin

Results showing the reversible changes in nuclear shape and relaxation in cells treated with Actin cytoskeleton inhibitors indicated a possible way to interfere with YAP signaling and, thus, reverse the cSt-Cs pro-fibrotic activation controlled by cell and nuclear mechanics. On the other hand, these inhibitors are quite unspecific and display an elevated cytotoxicity *in vivo*, thereby calling for more specific pharmacological inhibitors of the YAP nuclear functions. A more specific inhibitor was found in the Verteporfin (VTP) molecule, a drug with anti-fibrotic properties[46, 47], recently found to inhibit cardiac remodeling in mice[26, 48].

In the first series of experiments, cSt-Cs were treated with VTP under the maximal strain condition (culture onto glass) and then assayed for the expression of YAP target genes and genes involved in cardiac fibrosis[26]. To exclude that VTP has toxic effects on cells, we performed live/dead staining of cells treated with increasing amounts (1 – 10 μM) of the drug for 5 hours, followed by a 48 hours observation period (**Figure S6**). Having excluded toxic effect of VTP, we then analyzed its biological effects. **Figure 5A** shows that a 5hrs treatment with the drug did not modify the cellular shape, and did not affect YAP nuclear localization. By contrast, RT-qPCR experiments clearly showed a reduction in the expression of *CTGF*, *CYR61* and *ANKRD1*, as well as of *Coll1A1*, *CollA3* and *Thy-1*, which are typical markers of cells with a fibrotic phenotype[49].

We then analyzed the effects of a more chronical treatment with VTP (3 days) in the context of the signaling induced by TGF-β1, one of the most potent inducers of cardiac fibrosis[50]. In these experiments, we co-treated cSt-Cs with TGF-β1 (±VTP) and assessed the expression of fibrotic genes after 72 hours using an RNA

sequencing approach. Following data processing and raw count filtering criteria, we identified 17781 expressed genes, which included 13112 protein coding genes (74%), 2823 pseudogenes (16%), 1770 long non-coding genes (10%), and a small fraction of small non-coding genes (<<1%) (Gene ID details in **supplementary bioinformatics data – a**). Paired-sample analysis adjusted for confounding "latent" variables allowed reducing the effects of heterogeneity among subjects, thus unveiling specific changes between each treatment *vs.* CTRL. We report the overall results and statistics for each statistical model in the supplementary material, and summarize the main findings in **Figure S7A, B** with a representation of the differentially expressed genes in the comparisons of the treated cells *vs.* control cells, or in the VTP+TGF-β1 *vs.* TGF-β1 condition. In brief, we found a substantial number of differentially expressed (DE) genes by comparing the mRNAs of VTP-treated cells (±TGF-β1) *vs.* CTRL and to a lesser extend also for the treatment with TGFβ alone. On the other hand, 103 out of the 196 genes up-regulated by TGFβ were down-regulated by co-treatment with VTP, indicating that VTP has a down-regulatory effect on genes induced by the pro-fibrotic cytokine (**Figure S8A**). This evidence was supported by the results of unsupervised hierarchical clustering, which allowed us to distinguish between the diverse treatment conditions and, in particular, indicated a clear distinction in the transcriptional signature of cells treated with VTP (±TGF-β1) and those that were cultured without the drug (**Figure 5B**). We validated the transcriptional signature by performing single RT-qPCR assays with specific primers for *CTGF*, a pro-fibrotic factor expressed under the direct control of YAP/TAZ/TEAD complex[51], for the two collagen-encoding genes *ColA1*, *ColA3*, and for *Thy-1* and *ACTA2*, the genes encoding for CD90 and *αSMA*, two markers of cardiac Myo-Fbs[49, 52]. In all cases, treatment of the cells with VTP reverted the upregulation of these genes even below the level exhibited in control cells **(Figure 5B**), again confirming a potent transcriptional inhibitory effect of VTP in human cSt-Cs. Since interference with YAP signaling may also result in an imbalance in the expression of the Hippo pathway, we investigated the relative expression of genes with a functional annotation in the pathway. As shown in F**igure 5C**, a number of DE genes were up- or downregulated in VTP-treated (±TGF-β1) *vs.* untreated cells (±TGF-β1). Among the upregulated genes, we noticed the expression of *DVL2*, that in analogy to DVL, is involved in balancing the YAP cytosol/nuclear ratio[53, 54] and that of *AMOTL1*, a member of the Angiomotin proteins family, which has a specific inhibitory function of YAP activity by promoting phosphorylation via LATS proteins [55]. It was also remarkable that VTP treatment determined downregulation of the transcripts of *LATS1/2*, the transcripts encoding for the kinases of the Hippo pathway directly promoting YAP phosphorylation and functional activity[56], of *TJP1*, encoding for a tight junction protein positively regulating the TAZ/TEAD function[57] and of *YAP* mRNAs itself. Together, these results suggest that treatment with VTP determines an imbalance in numerous checkpoints of the finely regulated mechanism tuning the activity of the YAP/TAZ transcriptional function. In order to substantiate the action of VTP on cSt-Cs with the attribution of specific biological functions, we performed a Gene Ontology (GO) enrichment analysis of the DE genes in the various treatments. As expected, this showed a majority of pathways that were downregulated with high significance in VTP-treated cells (±TGF-β1) connected with extracellular matrix organization, cell migration, inflammatory responses and cytokines production (**Figure S9** and complete description of the pathways in **supplementary**

bioinformatics data – b), consistent with a reduction of the inflammatory and matrix remodeling activity of cells treated with VTP. Finally, in order to contextualize the inhibitory effect of VTP in the framework of the TGF-β-dependent transcriptional signaling, we performed *cis*-regulatory sequence analysis on down-regulated genes by VTP, and identified binding motifs and tracks associated with TEAD 1/3/4 and SMAD4 transcription factors among the most significant. As shown in **Figure S9** (see also **supplementary bioinformatics data – c**), the analysis identified groups of genes that are likely co-downregulated by VTP, due to connection of the YAP/TAZ complex with transcriptional mediators downstream of TGF-β1 (SMADs), through cooperation of TEADs.

VTP inhibits proliferation and reduces matrix remodeling activity of human cSt-Cs.

Since conversion of cardiac stromal cells into Myo-Fb is accompanied by transitioning into a rapid proliferation phase under the control of TGF-β1 signaling[50], or even mechanical cues regulated by YAP [15, 43], we determined the percentage of cells exhibiting PCNA staining in cSt-Cs cultured with or without TGF-β1±VTP onto glass slides. Results evidenced a sharp increase in the percentage of proliferating cells in the presence of TGF-β1 (**Figure 6A**), blunted by VTP treatment. In a previous study, we established a method to assess cell strain-dependent Myo-Fb differentiation onto substrates with discrete stiffness, by determining the loading of αSMA onto the F-actin cytoskeleton as a measure of the ability of the cells to evolve toward a contractile phenotype[29]. Adopting this system, we found a clear effect of TGF-β1 treatment in increasing the number of the fluorescence peaks showing co-localization of F-Actin and αSMA onto stress fibers, using confocal imaging (**Figure 6B**), thus confirming promotion of a Myo-Fb phenotype. On the other hand, cells treated with VTP reduced, at least in part, the loading of αSMA onto the actin cytoskeleton, suggesting reversion of the Myo-Fb phenotype due to TGF-β1 treatment. To further validate the effect of VTP in reversing the Myo-Fb activation of cSt-Cs, we assessed the expression of αSMA using fluorescence quantification on confocal images and Western blotting, and by measuring the collagen secretion in the culture medium (**Figure 6C**). In line with the previous results, VTP reduced the expression of the Myo-Fb marker and the secretion of collagen even below the level of control cells. Finally, since the ability to remodel the extracellular matrix in the context of myocardial infarction is a specific activity of Myo-Fbs, we measured the matrix compacting ability of the cSt-Cs treated with TGF-β1, and the reversal of this activity by VTP, using a collagen compaction *in vitro* assay. Results (**Figure 6D**) showed a clear contraction of the 3D gel containing cSt-Cs treated with TGF-β1, as opposed to control cells, and cells treated with TGF-β1+VTP. Interestingly, cell force analysis by measuring the bending of PDMS micropillars did not show differences between the average forces developed by the cells treated under the different conditions (**Figure 6E**). Together, these results show that interfering with YAP-dependent transcriptional activity blocks differentiation of cSt-Cs into myofibroblasts induced by TGF-β1 and limits their matrix remodeling and compacting ability, without affecting the physiological intracellular transmission of mechanical forces.

Anti-fibrotic activity of VTP in a mouse model of ischemic cardiac fibrosis

To verify the possible anti-fibrotic activity of VTP *in vivo*, we set a mouse model of ischemic cardiac remodeling in mice. The treatment protocol of mice was in line with existing literature on the anti-fibrotic

effects of VTP in other organs, such as kidney[58]. Two time-points were chosen to assess the effects of the continuous drug administration in relationships with the known biphasic myocardial remodeling process characterized by a precocious inflammatory phase (7 days), followed by an anti-inflammatory phase characterized by scar formation and collagen deposition (day 28). The experiment was controlled by evaluating the cardiac function by echocardiography and, finally by histological examination of the explanted heart at 7 and 28 days. The results of this experiment are summarized **in Figure 7**, where it is clearly shown that administration of VTP reduced the extent of the fibrotic areas at either 7 or 28 days post MI (Figure 7A-D). It was interesting to note that the fibrotic areas in VTP-treated mice were less compact (compare insets in panels **7C** *vs.* those in **7D**) and, in some cases, exhibited a higher number of CMs resembling the interstitial fibrosis detected in patients with ischemic cardiomyopathy (**Figure 1**). The anti-fibrotic effect of VTP also determined a significant increase in infarct thickness and a reduction of the infarct size at 7 and to a lesser extent at day 28 (**Figure 7D**). The positive effect on fibrosis of VTP was not, however, counterbalanced by an improvement in cardiac function, as shown by the failure of the treatment to increase ejection fraction and left ventricle fractional shortening as well as to decrease the end-diastolic/systolic volumes.

Discussion

Relevance of extracellular matrix remodeling for activation of YAP-pathway *in vivo*

Previous investigations highlighted the importance of the YAP signaling in cardiac fibrosis. For example, it was found that cells in the infarct zone in mice exhibit YAP nuclear staining[25] and that in mice lacking Lats1/2 kinases, two components of the Hippo kinase pathway, cardiac fibroblasts transition into Myo-Fbs spontaneously, giving rise to a hyper-secretory phenotype worsening the outcome of myocardial infarction[24]. Since Lats proteins phosphorylate directly YAP preventing its nuclear translocation[56], the Authors investigated the promoter occupancy of the chromatin in wt and Lats$^{-/-}$ CFs and found significant enrichment of YAP-bound enhancer sites in *pro*-fibrotic genes in knockout cells. This evidence led them to conclude that YAP/TAZ complex promotes cardiac fibroblasts transitioning into myo-Fbs by stable modifications of chromatin architecture and activity. In keeping with these conclusions, selective genetic ablation of YAP in cardiac fibroblasts, reduced cardiac fibrosis and improved cardiac function after MI[26, 59].

To assess whether the active matrix remodeling creates permissive conditions for YAP nuclear translocation in cells with morphological characteristics of Myo-Fbs, we employed a mouse model of myocardial infarction and characterized the features of the cells exhibiting nuclear YAP localization as a criterion to determine its transcriptional activity *in vivo*[25, 56]. In this analysis, we took as a reference evidences emerging from computational models describing the regional deposition of collagen fibers following myocardial injury, where non-random patterns have been observed, likely depending on the alignment of the cells depositing the matrix caused by the anisotropic distribution of the strain forces[37]. Strikingly, as revealed by the nuclear ellipticity and orientation analyses performed in cells with the morphology of CFs in the scars at 4 weeks post-MI (**Figure 1**), we noticed a more frequent occurrence of nuclei with a more elongated shape and a more consistent alignment along the equatorial cutting plane of the left ventricle in cells with nuclear-localized YAP. Since the orientation of the nuclei in the infarct and, more in general, inside 3D matrices subjected to strain with a principally uniaxial component aligns the cells and the ECM deposition along the principal force vector[60], the prevalent nuclear translocation of YAP in cells with a higher nuclear ellipticity and a more frequent cellular circumferential orientation might be part of a strain-dependent anisotropic activation of the *pro*-healing response to infarct reinforcing the cardiac wall and preventing its immediate rupture after infarction, but promoting chronic ventricular remodeling (hypothetical model described in **Figure 8**)[61]. This hypothesis is corroborated by results showing that matrix remodeling in the infarcted heart involves a complex series of structural modifications, changes in mechanical properties and hyper-activation of *pro*-fibrotic signaling[62], as well as by evidences in other tissues, e.g. the cardiac valves, where the deformation of the nuclei in interstitial cells has been connected to cyclic deformations due to straightening and compaction of the fibrous ECM components (i.e. collagen)[63]. This validates YAP as a transcriptional 'sensor' of the dynamic remodeling of the cardiac ECM consequent to ischemic damage. Its expression in nuclei of CFs in the fibrotic areas in myocardial samples of patients with severe heart failure (**Figure 1**) suggests the possible activity of the pathway in the human pathology.

Cell/nuclear strain/compression-dependence of a pro-fibrotic pathway in human cardiac stromal cells

Human primitive cardiac stromal cells have been shown to display pro-fibrotic features when exposed to the native extracellular matrix from remodeled hearts, or even to the altered substrates synthesized *ex vivo* by pathological cardiac fibroblasts from failing hearts[62, 64, 65]. To get insights in the mechanically-activated pro-fibrotic evolution of the primitive human stromal cells, we employed a 2D cell culture systems to test the effects of discrete cytoskeleton tensioning on the activation of YAP target genes, and to assess reversion of the *pro*-fibrotic phenotype by treating cells with inhibitors of the F-Actin cytoskeleton. In line with evidences provided in other studies, performed either on cardiac or valve stromal cells[29, 43, 52, 66], human cSt-Cs exhibited a stiffness-dependent YAP nuclear translocation resulting into cell proliferation (**Figure 2**). When treated with inhibitors of the cytoskeleton tensioning, cSt-Cs reversibly extruded YAP from the nucleus and downregulated canonical transcriptional targets and *pro*-fibrotic genes (**Figure 4**). It was finally interesting to note that nucleus geometry and compliance were also reversibly affected by the release of cytoskeleton tensioning by the two inhibitors (**Figure 3**). The change in nuclear geometry due to relaxation along the *z*-axis of the microscopic view likely reflects the release of the nuclear tensioning - acting mainly in the 2D (*x*, *y*) dimensions - due to adherence of the cells to a rigid flat surface[17], similarly to the effect of plating the cells onto soft substrates (**Figure 2**). The decrease in nuclear compliance suggests a possible effect of releasing cytoskeleton tensioning not only on the reduction of the strain at the level of the nuclear lamina, but also generalized changes in the chromatin compaction possibly leading to modifications in the epigenetic setup. In line with these evidences, it was demonstrated that biophysical cues can alter the methylation and acetylation status of histones, favoring epithelial to mesenchymal transition[67], and that cells embedded into matrices simulating stiff pathological tissues exhibit a mechanical 'memory' establishing permanent activation of pathological programs[68]. Whether, and to what extent, the nuclear straining imposed by cytoskeleton tensioning in human cSt-Cs has an epigenetic readout is the subject of ongoing investigations.

The connection between YAP nuclear translocation, nuclear straining and activation of *pro*-fibrotic signaling was optimally validated in cardiospheres. Apart from the description of this model as an *in vitro* 'niche' of cardiac mesenchymal progenitors[39], the cardiosphere appeared to fulfill the characteristics of a suitable 3D system to assess the impact of cell/nuclear straining on YAP-dependent pro-fibrotic activation in cSt-Cs, in analogy to what we observed in the fibrotic scar in mice (**Figure 1**). We adopted this model by extending concepts validated in the earliest stages of mammalian embryogenesis, where mechanical-dependent inhibition of the Hippo pathway and a robust YAP nuclear translocation have been described as a topological determinant of the primitive trophectoderm cells differentiation in the cellular shell of the forming blastocyst[69]. To validate this model, we computer-analyzed the deformation of the nuclei in the cellular spheroids in relationship with the position of the cells in the core *vs.* the shell of the cardiospheres, the presence of nuclear-localized YAP, the expression of cell proliferation and *pro*-fibrotic cellular markers, and the deposition of ECM components (**Figure 3**). The suitability of this method, based on the CARE algorithm[34], allowed to establish rapidly, and with a high level of significance, a cause-effect relationship between the extent of nuclear deformation, the YAP nuclear translocation and the presence of Myo-Fb features (such as proliferation, expression of αSMA

and collagen) in the cells depending on their 3D positioning, with cells in the shell characterized by higher strain *vs.* those positioned in the core subjected to lower compression forces and strain. These evidences support, finally, the *in vivo* findings showing the effects of topological cues on the activity of the YAP signaling in a 3D multicellular cardiac fibrotic context (**Figure 1**).

Targeting downstream effectors of mechanical cues reverts TGF-β-dependent and independent CF fibrotic programming

The data of our RNAseq analysis support the notion that treating cells with VTP disassembles the bridging of YAP to TEADs regardless of cell strain and has unbalancing effects for the activity of the Hippo pathway with possible readouts on the level of YAP phosphorylation, transcriptional cooperation with the Wnt pathway, *YAP1* gene silencing (**Figures 5B, C**) and downregulation of several pathways connected with extracellular matrix remodeling, inflammation and cell polarity/migration (**Figure S9A**). Interestingly, cells treated with VTP also downregulated more than a half of the genes that were induced by TGF-β1, one of the most potent cardiac fibrosis-inducing factors[50, 62] (**Figure S8**). These genes included canonical YAP targets such as *CTGF (CCN2)* and *ACTA2*, genes connected to the CFs pro-fibrotic phenotype (*Thy-1*), and genes encoding for major scar components such as *COLA1* and *COLA3*. The centrality of the YAP/TAZ transcriptional signaling in the context of fibrotic activation of human cSt-Cs emerged, finally, from the bioinformatics search of the *cis*-regulatory elements potentially involved in the generalized gene downregulation observed in VTP-treated cells (±TGF-β1) *vs.* controls (**Figure S9B**). This search identified TEAD 1/3/4 and the common transcriptional transducer of TGF-β signaling SMAD4 as candidates for the co-regulation of genes involving YAP/TAZ complex as a common factor. While this is in line with the current literature identifying cooperation of these transcriptional modulators[70], it also highlights the centrality of the YAP/TAZ complex in the context of the TGF-β1 transcriptional signaling[71], suggesting effectiveness of 'mechano-therapeutic' approaches to reduce the extent of fibrosis in the cardiovascular system and other organs controlled by the pro-fibrotic factor.

In agreement with the variation in gene expression signatures, the blockade of YAP/TAZ/TEAD interaction by VTP also appeared to completely override the TGF-β1-dependent *pro*-fibrotic activation of cSt-Cs on a phenotypic and functional points of view. This was substantiated by the reduction in cell proliferation (**Figure 6C**), the unloading of SMA from the F-Actin cytoskeleton (**Figure 6B**), and the inhibition of collagen secretion/matrix compaction ability of the cells treated with TGF-β1 (**Figure 6C, D)**. Opposite to our hypothesis that cells treated with TGF-β1 might increase pulling forces, the experiment on micropillar arrays did not show any variation in cell contractility (**Figure 6E)**. This finding is only in apparent contradiction with the reduction in matrix compaction due to VTP treatment, considering that other factors such as secretion of matrix remodeling enzymes may concur to the increased collagen compaction of the cells treated with TGF-β1.

Anti-fibrotic effect of VTP treatment *in vivo* does not coincide with improvement of cardiac function

The interest for manipulating the Hippo pathway as a strategy to repair/regenerate the heart has produced a growing course of studies with remarkable, but sometimes antagonistic results (reviewed in[72]). In fact the existing literature clearly distinguishes between the function of the YAP/TAZ complex in the heart, depending

on the localization of the loss/gain of function effects. For example, while upregulation/stabilization of YAP in CMs primes the contractile cells to divide, with potential cardiac regenerating effects[19, 20, 22], the hyperactivation of the YAP/TAZ complex in cardiac fibroblasts primes these cells toward a chronic scarring phenotype that result in accelerated and injury-independent myocardial remodeling[24] (also discussed in [23]). On the other hand, more recent studies showed that selective ablation of YAP in CFs reduces ischemia or pressure overload-dependent fibrosis with an improvement of myocardial function[26, 59], thus opening the way to possible mechano-therapeutic strategies to limit cardiac fibrosis.

Prompted by the positive effects of VTP administration in models of ischemia/reperfusion-dependent kidney fibrosis[58], in the present study we used a classical VTP continuous administration protocol in mice with a chronic cardiac ischemia setting. Although this clearly reduced the accumulation of collagen and, at least in part, preserved the infarct wall from the extreme thinning observed in control mice, the administration of the drug did not relief the detrimental effects of ischemia on cardiac function (**Figure 7**). Different possibilities may account for this effect, which makes the results of our findings clearly different from those achieved in genetic models of selective YAP inhibition in CFs. A first element that may play a role in this difference is the possibility that VTP alters/retards the timing of the pro/anti-inflammatory phases after cardiac damage. For example Freeman and colleagues showed that selective ablation of YAP in early stage macrophages after MI promoted cardiac repair by shifting the phenotype of toward an anti-inflammatory phenotype[36]. In line with this, our treatment with VTP could have an initially positive effect on selective ablation of pro-inflammatory macrophages but, thereafter, a pleiotropic inhibiting function of anti-inflammatory macrophages with detrimental effects for cardiac functional recovery. A second possibility may derive from the ambivalent function of the YAP/TAZ complex in cardiac myocytes *vs.* fibroblasts. For example, YAP loss of function in CMs leads to impaired survival after cardiac injury with worsening effects on maladaptive ventricular remodeling after myocardial injury[73]. In such a case, the beneficial effect of VTP administration on reduction of fibrosis might be counteracted by a reduced survival of myocytes in the infarcted heart, thus limiting the therapeutic efficacy of bulk administration of the drug. In order to substantiate our results in translational and functional perspectives, we are currently evaluating, ***i)*** other treatment modalities (e.g. to start or interrupt administration of the drug at defined times after the initial pro-inflammatory period) to better distinguish between possible antagonistic effects of the continuous VTP treatment on myocardial functional recovery, ***ii)*** the adoption of imaging-derived parameters of global cardiac function such as the myocardial strain[74], ***iii)*** the use of 3D imaging tools such as the second harmonic generation microscopy[62], and ***iv)*** of force-based mapping by nanoindentation[29] of the cardiac tissue.

Conclusions

The emerging role of YAP in fibrotic progression in several diseases[75], including myocardial remodeling after infarction [25], provides a strong rationale for a potential anti-fibrotic therapy of the failing heart, based on local mechanical desensitization of the *pro*-fibrotic cells. Translation of genetic approaches into protocols of selective pharmacological inhibition of the complex in CFs but not in CMs is, in fact, not currently amenable using systemic administration protocols.

In the present study, we provide evidences that the pro-fibrotic programming of human cSt-Cs *in vitro* is subjected to the cooperation of mechanical, topological and paracrine cues likely originating from the known effects of strain forces resulting from the anisotropic arrangement of the collagen matrix occurring during scar formation[37, 76] (**Figure 8**), and the results of the nuclear strain/orientation analyses performed on YAP^+ and YAP^- nuclei in the infarct fibrotic scar. We extend this conclusion also to an *in vivo* cardiac remodeling situation, where the administration of VTP caused a significant inhibition of the fibrotic progression in the ischemic hearts, even though this improvement was not accompanied by a net increase in cardiac performance.

A final and more general conclusion of our investigation is that, in analogy to what already described in other cardiovascular pathological settings such as the aortic valve disease[29, 77] or vascular pathological conditions[78, 79], understanding the cooperation between mechanical cues and paracrine factors in cardiovascular diseases will be a possible key to achieve innovative and targeted anti-fibrotic therapies. Indeed, if administered with systems specifically designed to perform drug delivery in the fibrotic tissues (e.g. nanotechnology), these therapies will be optimal candidates to mechanically desensitize the pro-fibrotic cells, with hopes for robust reverse remodeling effects.

Novelty and significance

Emerging evidences suggest that mechanical signaling is crucial for acquisition of pathological phenotypes in cardiovascular tissues remodeling. While the identity and the function of mechanically regulated transcriptional activators in cardiac pathology has been clarified mainly with genetic studies, a direct connection between cell mechanics and progression of ischemia-dependent fibrosis was missing. In the present contribution, we show that nuclear translocation of the Yes-Associated Protein (YAP) transcription factor occurs by exposing cardiac fibroblasts to incremental strain/compression forces either in vivo or in vivo. We also provide evidences that pharmacological interfering with nuclear function of YAP is sufficient to override the TGF-β1-dependent pro-fibrotic programming in vitro, and to reduce the extent of cardiac fibrosis in vivo. Our results open the way to ‘mechano-therapeutics’ of the fibrotic heart.

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Disclosures

The Authors declare no conflicts of interest.

Supplemental materials

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Figure Legends

Figure 1. Activity of YAP-dependent signaling is subjected to oriented nuclear straining and cellular alignment in the infarct scar. (**A**) Equatorial section of the infarcted heart showing the left and right ventricles (LV and RV, respectively) at a short follow-up time (3 days post-MI) stained with anti-YAP antibodies. The higher magnification on the left side of the panel shows the presence of cardiomyocytes (cm) and fibroblasts (Fbs) characterized by low levels of nuclear YAP. A preferential localization of YAP^+ cells was observed in the infarct zone (IZ) magnified in the right side of the panel, which at this time point is not yet subjected to extensive remodeling, but is affected by a high degree of inflammation. At this stage, YAP nuclear signal was detected in inflammatory infiltrate cells (infl), in some myofibroblasts (Myo-Fbs) and cardiomyocytes (cm). (**B**) Masson trichrome staining of a terminal stage (four weeks post-MI) cardiac remodeling event, characterized by the presence of a collagen-rich scar extending from the wall of the right ventricle (RV) to a wide portion of the left ventricle (LV). On the left side of the panel it is represented the whole equatorial section of the heart to show the totality of the remodeling process. In the center and the right of the panel are represented two consecutive magnifications of the areas of the infarct zone (IZ) encircled by the blue dotted squares, to show the orientation of the collagen (Coll) bundles along a preferential equatorial plane (red arrows). Note the the presence of numerous fibroblasts exhibiting a similar orientation. (**C**) YAP immunohistochemical labeling of the same heart cut with the same equatorial orientation shown in panel **B**. The magnification of the two areas encircled by the red dotted squares show the presence of collagen bundles (Coll; blue arrows) and of fibroblast-resembling cells characterized by presence of YAP in the nucleus (YAP^+) or absence of the transcription factor (YAP^-), indicated by red arrows. Note in both panels that cells with YAP^+ nuclei appeared in both areas to show a higher nuclear aspect ratio (ellipticity). (**D**) Virtual reconstruction of the nuclei orientation in the infarct scar as detected by CARE. On the left side of the panel it is represented the actual image of an equatorial section of the left and right ventricle (LV and RV, respectively) of a heart with an end-stage remodeled infarct zone (red dotted square), with an indication of the profiles of the external cardiac wall and the internal LV wall (both indicated by blue lines) and the midline of the wall (green line). The panel in the center shows the margins of the scar (green lines) defined by the two concentric lines contouring the collagen-rich region in the scar, and the scar centerline (red). The orientation of each cell nucleus was measured by determining the tilting of the major nuclear axis with respect to the line intersecting perpendicularly the scar midline in manually selected zones with a clearly visible fibrotic appearance in the scars (green boundaries in the right panel; see also **Figure S1**). (**E**) Comparison of nuclear ellipticity and orientation as computed by CARE for YAP^+ and YAP^- nuclei in the terminally remodeled infarct of 3 mice with a 4 weeks post-MI follow-up. The box plot on the left represents the min-max distribution, the median and the mean (+) of the nuclear aspect ratio in YAP^- (blue) and YAP^+ (brown) nuclei. **Data were analyzed** by Mann-Whitney t-test (n = 3329 YAP^- and = 4559 YAP^+ nuclei, respectively). The two distributions on the right side of the panel show the percentage of the cells with nuclei with an orientation (θ) $\pm 10°$ in a $0° < \theta < 180°$ range with respect to the perpendicular direction to the scar center line (See **panel E** and **Figure S1**). Data in the two distributions are indicated as percentage (±SE) of YAP^- (blue) or YAP^+ (brown) nuclei of the

589 total amount of computable nuclei in the scar of 3 mice within the 18 resulting ±10° orientation categories. As
590 shown, the percentage of YAP^+ nuclei with an orientation ±10° within the $70° < \theta < 110°$ range were
591 significantly higher than that of YAP^- nuclei. Data were compared by unpaired multiple t-test analysis. (**F**)
592 Low and high magnification of representative sections of human myocardium from patients with dilated
593 hypokinetic ischemic cardiomyopathy stained with Masson's trichrome and YAP IHC. In the lower insets it is
594 evident the presence of numerous fibroblasts with YAP^+ nuclei populating the collagen scar.

595 **Figure 2. Strain-dependent activation of YAP transcriptional pathway in human cSt-Cs.** (**A**) Geometric
596 features (cell spreading area; circularity) of human cSt-Cs are subjected to stiffness-dependent control. Cells
597 were plated onto glass and onto a series of polyacrylamide gels with controlled stiffness ranging from 58kPa
598 to 17kPa (Young's modulus)[29] and were photographed to derive geometrical information. The pictures show
599 the staining of the nuclei (blue) and the F-Actin cytoskeleton by Phalloidin-TRITC (white color). The two
600 graphs include areas [μm^2] and circularity factor from more than cells from each of the indicated number of
601 experimental replicates performed with cSt-Cs lines from different donors. (**B**) The expression of YAP (green
602 fluorescence) was visualized by IF along with the nuclear and F-Actin staining by DAPI (blue) and phalloidin-
603 TRITC (red), respectively. The percentage of cells with nuclear localized YAP (arrows) and the $YAP_{NUCL/CYTO}$
604 expression ratio were determined using ImageJ and CARE[34], respectively (see also **Figure S3**). Graphs were
605 derived with data obtained in at least 10 cells from each of the indicated number of experimental replicates
606 performed with cell lines from different donors. (**C**) Cells plated onto glass and PAGs with differential stiffness
607 were labelled with Ki-67 proliferation marker (green, arrows)) along with F-Actin (red) and nuclear (blue)
608 staining. The graph shows the quantification of the results and indicates the statistical significance in the
609 comparison between experimental groups. In all graphs in the panels, the P values of the statistical comparisons
610 between the cells plated onto glass *vs.* PAGs with the different stiffness, or between cells plated on the
611 differential stiffness PAGs are calculated by one-way pairwise Anova with Dunnet and Tukey post-hoc tests,
612 respectively.

613 **Figure 3. Topological cues support an asymmetric distribution of cells with high $YAP_{NUCL/CYTO}$ expression**
614 **ratio and proliferation/fibrotic markers in a 3D context.** (**A**) Phase contrast view of primary cardiospheres
615 derived from human myocardial tissue (left) and confocal image of a cardiosphere whole mount-stained with
616 YAP-specific antibody (green, right), F-actin (red), and nuclei. The image on the right shows, on the top, the
617 3D projection of the cardiosphere and the *x*, *y*, *z* dimensions (µm) as detected by high-resolution confocal
618 imaging. The three images on the bottom show three discrete *x*, *y* equatorial images of the internal view of the
619 same cardiosphere at the indicated distances (-35µm, -20µm and -10µm) from the top of the sphere (set at
620 0µm) along the *z* axis. In these images it is evident the asymmetric distribution of cells with YAP nuclear
621 signal in proximity of the surface of the cardiosphere compared to its core. The arrows in different colors show
622 examples of nuclei exhibiting a high content of nuclear YAP in the three equatorial images, also indicated with
623 the same color code in the 3D projection on the top. (**B**) On the right side of the panel it is represented a
624 confocal imaging 3D projection (top) and three *x*, *y* equatorial views (bottom) of a cardiosphere stained with
625 Ki-67 antibody (white fluorescence) and nuclei (blue fluorescence) as described for **panel A**. The two graphs

on the left represent, respectively, the $YAP_{NUCL/CYTO}$ expression ratio (top) and the percentage of the cells expressing Ki67 (bottom) as computed by CARE algorithm. (**C**) The whole mount IF of GATA-4, a cardiac specific transcription factor, which is not known to be mechanically regulated, did not show differences in the nuclear/cytoplasm expression ratio between the shell and the core of the cardiospheres at any of the equatorial *x*, *y* projections. Data computation by CARE confirmed no statistically significant differences in the distribution of the ratios in the two compartments. (**D - E**) Whole mount IF staining of cardiospheres with Collagen-I and αSMA antibodies, respectively, visualized as in **panels B** and **C**. As shown by the images and the *x*, *y* projections, these markers tended to be more expressed at the periphery and not in the center of the cardiospheres. Data computation with CARE performed on the fluorescence distribution in the shell vs. the core of the spheres showed a clear asymmetry, similar to that found for YAP. In all graphs, the *P*-value of statistical comparison by paired student's t-test is shown. The number of cardiospheres (obtained from 4 independent tissue donors) introduced in the analyses are represented by the orange circles overlapped to the bars.

Figure 4. Nuclear geometry and tensioning regulates activity of YAP transcriptional pathway in human cSt-Cs. (**A**) Comparison between the shape of the cSt-Cs when in contact with hard (58kPa) and soft (17kPa) substrates. The low magnifications on the left show the cellular shape along with the YAP labeling (green); the magnifications of the areas encircled with the dotted squares on the right show the major (a) and the minor (b) axes, used to calculate the aspect ratio (Ellipticity) of the nuclei. Note that in cells in contact with the soft substrate, YAP fluorescence was almost localized almost entirely in the cytoplasm, while in cells adhering onto the stiff substrate it was almost entirely into the nucleus (see also **Figure 2** for quantifications). The graph on the right shows the quantification of nuclear ellipticity in cSt-Cs plated onto the whole series of PAGs (plus glass as a control). The *P* values of the statistical comparison of the nuclear aspect ratio in cells the cells plated onto glass *vs.* PAGs with the different stiffness, or between cells plated on the differential stiffness PAGs are indicated. These values were calculated by one-way pairwise Anova with Dunnet and Tukey post-hoc tests, respectively. (**B**) Calculation of the nuclear aspect ratio in the cardiospheres with CARE. The two images show, respectively, the equatorial section of a cardiosphere labeled as in **Figure 3**, with YAP antibody (green), for F-Actin (red) and nuclei. Arrows indicate filamentous Actin labeling, showing putative stress fibers. The image on the right is the virtualization of the nuclei image in the left with an indication of the shell and the core of the sphere (see also **Figure S4**). This image contains a color-coded nuclear structural information according to a nuclear ellipticity scale comprised in this panel between 0.40 and 0.50. It is evident that the majority of the nuclei with higher aspect ratio were present in the shell of the sphere and that more round nuclei were abundant in the core. The graph on the right is a statistical comparison of the average nuclear ellipticity in the shell and the core in the indicated number of cardiospheres. The *P* value of data statistical comparison by paired student's t-test is indicated above the graph. (**C**) Cellular effects of treating cSt-Cs with ROCK inhibitor Y27632 and Myosin II inhibitor Blebbistatin on substrate with maximal stiffness (glass). Cells are labeled with YAP antibody (green), F-Actin label (Phalloidin-TRITC, red) and nuclei (DAPI, blue). Before treatment (CTRL, left), cells exhibited a normal fibroblastoid/mesenchymal phenotype and nuclei with a high

level of nuclear YAP (arrows). Treating them with both inhibitors (center panels in **C**) reduced the number of stress fibers and induced a change in cell shape, with a consistent reallocation of the YAP fluorescence in the cytoplasm. A re-tensioning of the stress fibers was observed with washout of both drugs (right panels), along with a return to normal level of YAP nuclear confinement (arrows). **(D)** Quantification of YAP nuclear confinement. *P* values of data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test are indicated above each graph. (**E**) Nuclear geometry and compliance are affected by release of cytoskeleton tensioning. The pictures on the left show the normal (*x*, *y*) microscopic view of glass-adhering cSt-Cs labeled with YAP antibody (green), F-Actin probe (red) and nuclear label (blue). The pictures on the right of each panel show the projection of the cells circled with a dotted square, along the *x*, *z* and the *y*, *z* axes, as detected by super-resolution confocal imaging. It is evident that treatment with inhibitors determined a relaxation of the nuclei along the *z* axis and that after washout of the cells, the re-tensioning of the stress fibers caused a return to a nuclear flattening condition (in keeping with YAP nuclear segregation). The effect of nuclear geometry on nuclear mechanical characteristics is shown in the panels on the right, where a quantification of the nuclear geometrical changes along the *z*-axis by both treatment (and the treatments washouts) is shown, together with the physical 'softening' of the nucleus, as detected by AFM force imaging. In both graphs, the *P* values of data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test are indicated above each graph. (**F**) Transcriptional readout of cytoskeleton inhibition consisted of downregulation of canonical YAP targets, as shown by RT-qPCR amplification of *CTGF*, *CYR61* and *ANKRD1* gene transcripts. Note that the expression level of these genes returned to baseline after drugs washout. Data are represented as fold changes (FC) in the expression of each gene in the indicated condition with respect to the expression level in untreated cells (indicated with a C corresponding to a level=1) calculated by the $2^{-\Delta\Delta Ct}$ method. Statistics was calculated on the corresponding ΔCt values. Above all graphs, the *P* values of RT-qPCR Delta-CT data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test indicate the significance of differences in the expression of each tested gene in VTP-treated *vs.* control cells.

Figure 5. Global transcriptional effects of VTP. (**A**) cSt-Cs pulse-chased with verteporfin (VTP) for 5 hours exhibited a partially reversible downregulation of YAP target genes (*CTGF*, *CYR61*, *ANKRD1*) and *pro*-fibrotic markers (*Col1A1*, *Col3A*, *Thy-1*). Transcriptional inhibition was not accompanied by cell morphological changes and YAP cytoplasm reallocation, as in experiments performed with cytoskeleton inhibitors Y27632 and blebbistatin (**Figure 4**). Data are represented as fold changes (FC) in the expression of each gene in the indicated conditions with respect to the expression level in untreated cells (indicated with a C corresponding to a level=1) calculated by the $2^{-\Delta\Delta Ct}$ method. Above all graphs, the *P* values of RT-qPCR Delta-CT data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test indicate the significance of differences in the expression of each tested gene in VTP-treated *vs.* control cells. (**B**) Results of an RNAseq analysis of RNA samples extracted from control cSt-Cs, and cSt-Cs cultured ±TGF-β1 (±VTP) for three days. Hierarchical clustering was performed by Euclidean (sample) and 1-pearson correlation (genes) metric and average linkage method; gene expression levels are displayed as gradient colors from higher (dark red) to lower (dark blue). The heat map on the left shows the results of DE gene unsupervised clustering,

exhibiting a nearly perfect clusterization of genes significantly regulated by the treatment with the drug. The table on the bottom indicates the number of the DE genes (adj. P-Value<0.05 and |log2FC|>0.58) for each comparison between treatment vs. CTRL. DE genes are also distinguished between those that are up- or down-regulated in each treatment vs. CTRL. On the top right side of the panel, we report a representation of RT-qPCR analysis of five genes regulated by YAP whose modulation was consistent with the observed changes in the RNAseq profiling, in independent cellular replicates. In all panels, graphs were generated using the fold changes (FC) in the expression of each gene in the indicated conditions with respect to the expression level in untreated cells (indicated with a C corresponding to a level=1) calculated by the $2^{-\Delta\Delta Ct}$ method. Above all graphs, the *P* values of RT-qPCR Delta-CT data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test indicate the significance of differences in the expression of each tested gene in VTP-treated *vs.* control cells or between treatments. The heat map on the bottom right indicates the variation in the expression of the same genes as represented in the general dataset of the DE mRNAs (**bioinformatics data supplement – a**). (**C**) Differential expression of genes with functional annotation in the Hippo pathway. With * are indicated differentially expressed genes (adj. *P*-Value < 0.05 and |log2FC|>0.58) found in at least one of the treatments *vs.* CTRL comparison.

Figure 6. Biological effects of VTP treatment in vitro. (**A**) Treatment with VTP reduces proliferation of human cSt-Cs, as assessed by IF staining with antibodies specific for PCNA (green fluorescence) and F-Actin (red fluorescence)/nuclear staining (blue staining). Note that the drug reduced the percentage of $PCNA^+$ cells (arrows) below the control level even in the presence of TGF-β1, supporting a strong reduction of cellular proliferation, a recurrent feature in conversion of primitive stromal cells into myo-FBs. Above the graph, the *P* values indicate the results of data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test. (**B**) cSt-Cs treated with VTP loose myo-FB characteristics promoted by TGF-β1, as detected by the unloading of αSMA from F-Actin cytoskeleton. The three images on the top show three cells labelled with F-Actin (red fluorescence) and nuclear staining (blue fluorescence) along with αSMA antibody (green fluorescence). As shown by the fluorescence intensity profile along the indicated dotted lines, treatment with VTP reduced the co-localization of αSMA and F-Actin signals (indicated by * in each of the plots corresponding) that was elevated by treatment with TGF-β1. (**C**) Reduction of αSMA protein expression and of collagen-1 secretion by VTP treatment. The top left panel indicates the integration of the αSMA IF signal calculated as the integrated fluorescence density using ImageJ software. The panel on the top right represent an example of a Western blotting analysis performed with whole protein extracts from cSt-Cs treated as indicated. Note the decrease of the αSMA band intensity in VTP-treated cells compared to controls and TGF-β1 treatment, also indicated in the bar graph in the low right, showing quantification of the normalized αSMA/GAPDH in all the conditions. The panel on the low left indicates the reduction in collagen secretion by the cells treated with VTP *vs.* CTRL and TGF-β1 treatment. Also in this case, this reduction occurred also in the combined VTP+ TGF-β1 treatment. Above all graphs, the *P* values indicate the results of data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test. (**D**) The ability of the cells to remodel the matrix was assessed by the collagen plug contraction assay. The panels on the top show the top-view

images of the plugs containing cells treated as indicated, while the bar graph on the bottom shows the quantification of the area occupied by the plug at 24hrs after matrix release. Lower areas are indicative of a higher contraction activity, in response to Myo-Fb differentiation of human cSt-Cs determined by TGF-β1. VTP treatment completely reverted the contractile phenotype of the cSt-Cs. Above the graphs, the *P* values indicate the results of data statistical comparison by one-way pairwise ANOVA with Tukey post-hoc test. (**F**) Untreated and TGF-β1-treated cSt-Cs ±VTP were seeded onto PDMS micropillars to calculate the average traction force. The picture on the top illustrates the phase contrast image of a cell deposited onto the pillar array (left) and its corresponding F-Actin (red)/nuclear (blue) staining. The picture on the bottom is a virtual representation of the traction forces exerted by the cell onto each individual pillar, its direction (arrows) and intensity (color code). The graph on the bottom represents the average force exerted by a total of 12 cells (3 cells per donor) in the four treatment conditions.

Figure 7. Effects of VTP treatment on cardiac fibrosis and function in a mouse model of permanent cardiac ischemia. (**A**) Side by side comparison of equatorial sections of infarcted hearts harvested at 7 days post-MI from control (CTRL) and VTP-treated mice after staining with Masson's trichrome. It is evident the higher extension of the infarct, the higher collagen deposition, and the thinning of the scar in CTRL *vs.* VTP mice. (**B** – **C**) Side by side comparison of the scars at 28 days after MI in CTRL and VTP-treated mice. Also at this time point VTP reduced fibrosis and increased thickness of the infarct. Less evident was the effect of the drug on reduction of the infarct size. (**D**) Quantification of the infarct morphometry as detected in the tissue sections stained with Masson's trichrome at 7 and 28 days after MI. VTP reduced significantly the fibrosis and the infarct size and increased the wall thickness indicative of a lower LV remodeling. This effect was more pronounced at 7 days given that the reduction of the infarct size at 28 days post MI did not reach statistical significance. Above all graphs, the *P* values indicate the results of data statistical comparison by pairwise student's t-test. (**E**) None of the echocardiographic parameters (Ejection Fraction, LV end diastolic/systolic volumes and fractional shortening) were affected by the treatment at neither time.

Figure 8. **Proposed model of strain-dependent YAP transcriptional signaling activation in the infarct scar.** Myocardial remodeling after a transmural infarction involves matrix deposition and this occurs, according to literature, with a preferential equatorial orientation. This deposition pattern is due to a strain-dependent alignment of the collagen depositing cells along the principal strain vector. The findings in this work show that the variation in the nuclear aspect ratio in the collagen-depositing cells is connected to nuclear translocation of YAP. We propose that this creates a permanent activation condition of these cells toward a pro-fibrotic phenotype. Given that existing models of anisotropic collagen deposition in injured hearts describe this phenomenon only for regions of the cardiac wall at a distance from the LV apex[37], we cautiously extend our hypothetical model only to regions of the cardiac wall, where the distribution of strain forces occurs principally with an equatorial direction, and not in the apex. Further studies involving *in vivo* cardiac kinematics and serial sectioning of the scars at different sectioning planes would be necessary to further validate this hypothesis for the whole ventricular wall.

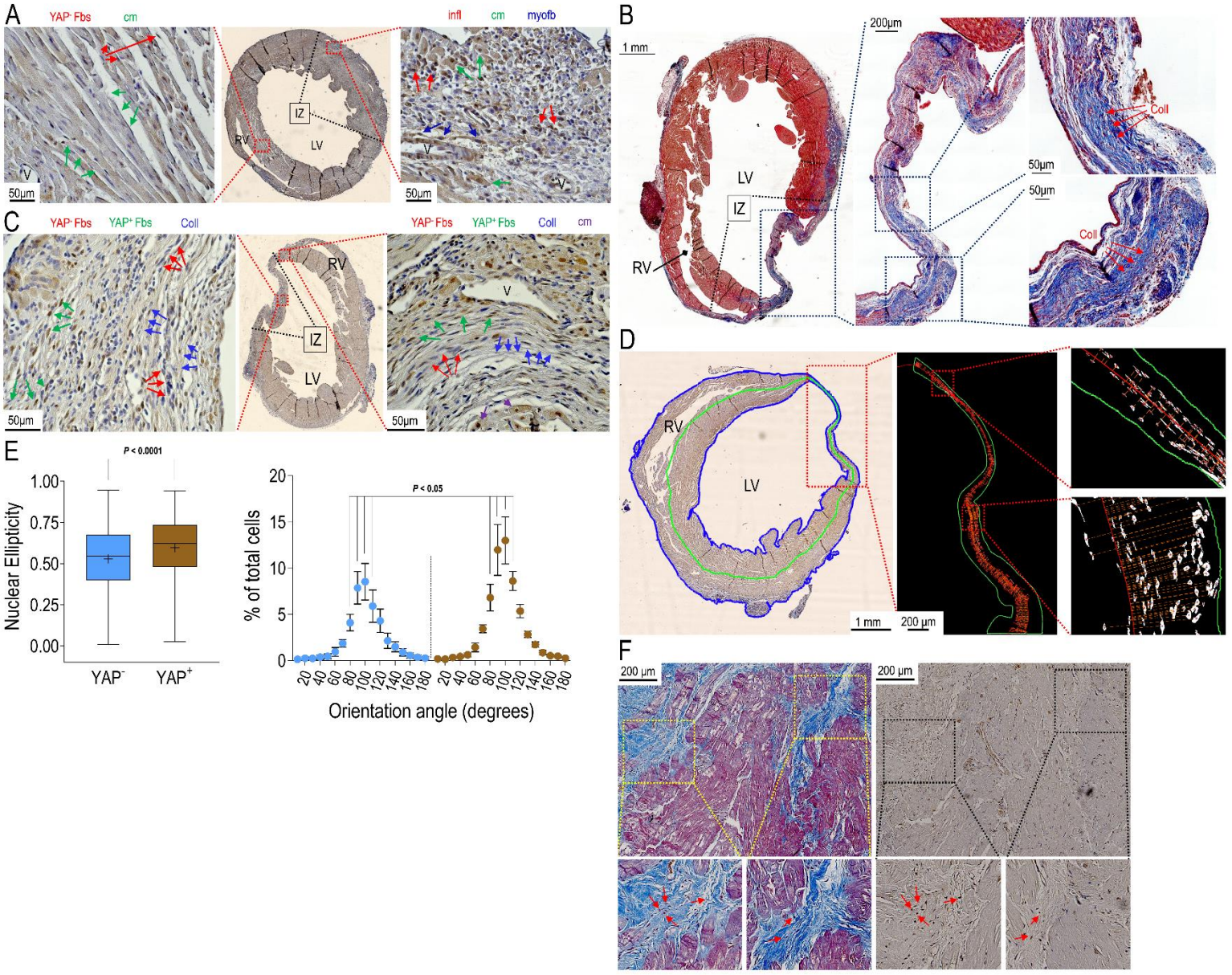

A
YAP- Fbs
cm
infl
cm
myofb
IZ
RV
LV
V
50µm
B
1 mm
200µm
LV
IZ
RV
Coll
50µm
C
YAP- Fbs
YAP+ Fbs
Coll
cm
D
RV
LV
1 mm
200 µm
E
P < 0.0001
Nuclear Ellipticity
1.00
0.75
0.50
0.25
0.00
YAP-
YAP+
P < 0.05
% of total cells
Orientation angle (degrees)
F
200 µm

Figure 1

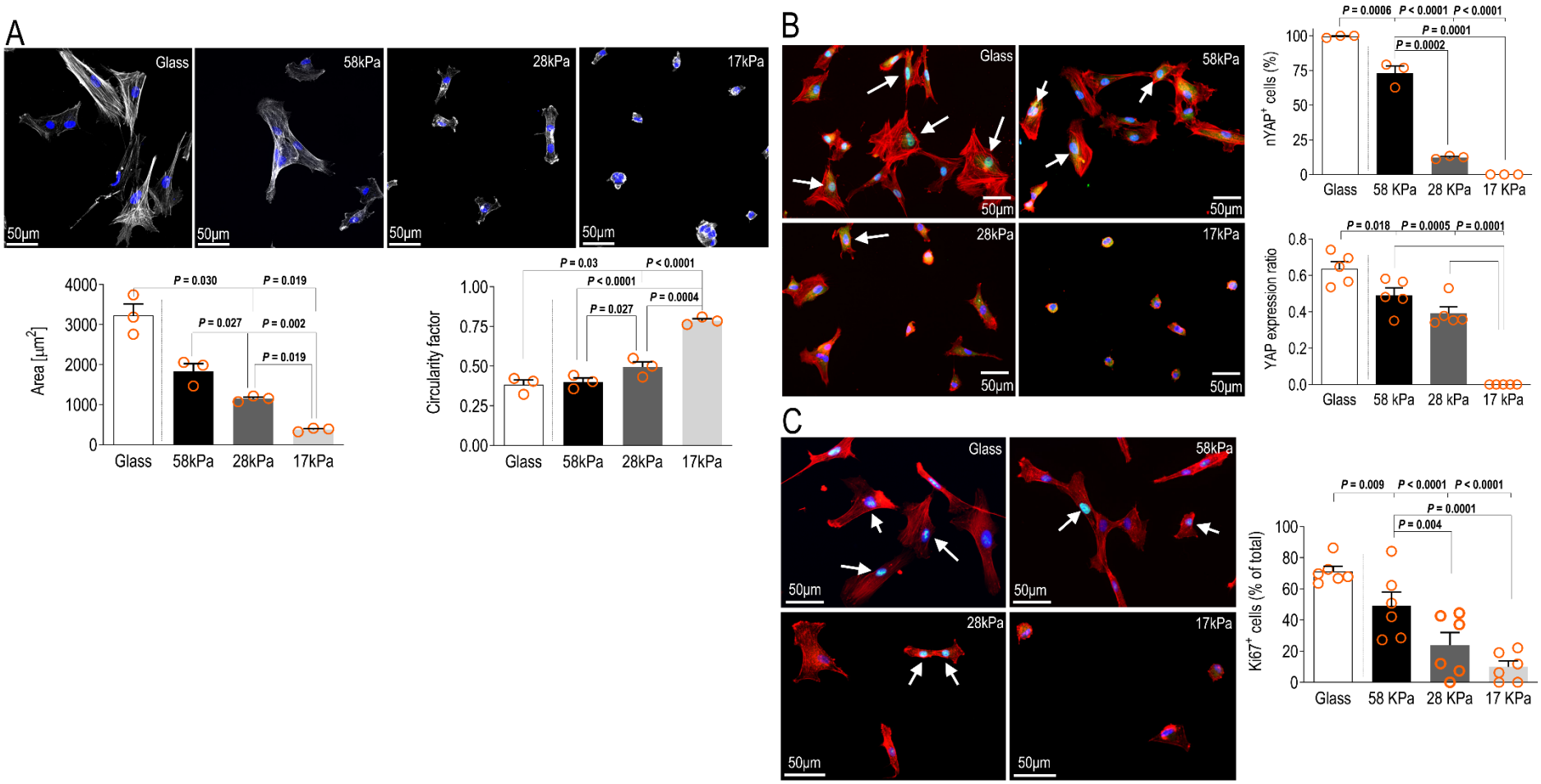

A
Glass
58kPa
28kPa
17kPa
50µm
Area [µm²]
P = 0.030
P = 0.019
P = 0.027
P = 0.002
P = 0.019
Circularity factor
P = 0.03
P < 0.0001
P < 0.0001
P = 0.027
P = 0.0004
B
nYAP⁺ cells (%)
P = 0.0006
P < 0.0001
P < 0.0001
P = 0.0001
P = 0.0002
YAP expression ratio
P = 0.018
P = 0.0005
P = 0.0001
C
Ki67⁺ cells (% of total)
P = 0.009
P < 0.0001
P < 0.0001
P = 0.0001
P = 0.004
58 KPa
28 KPa
17 KPa

Figure 2

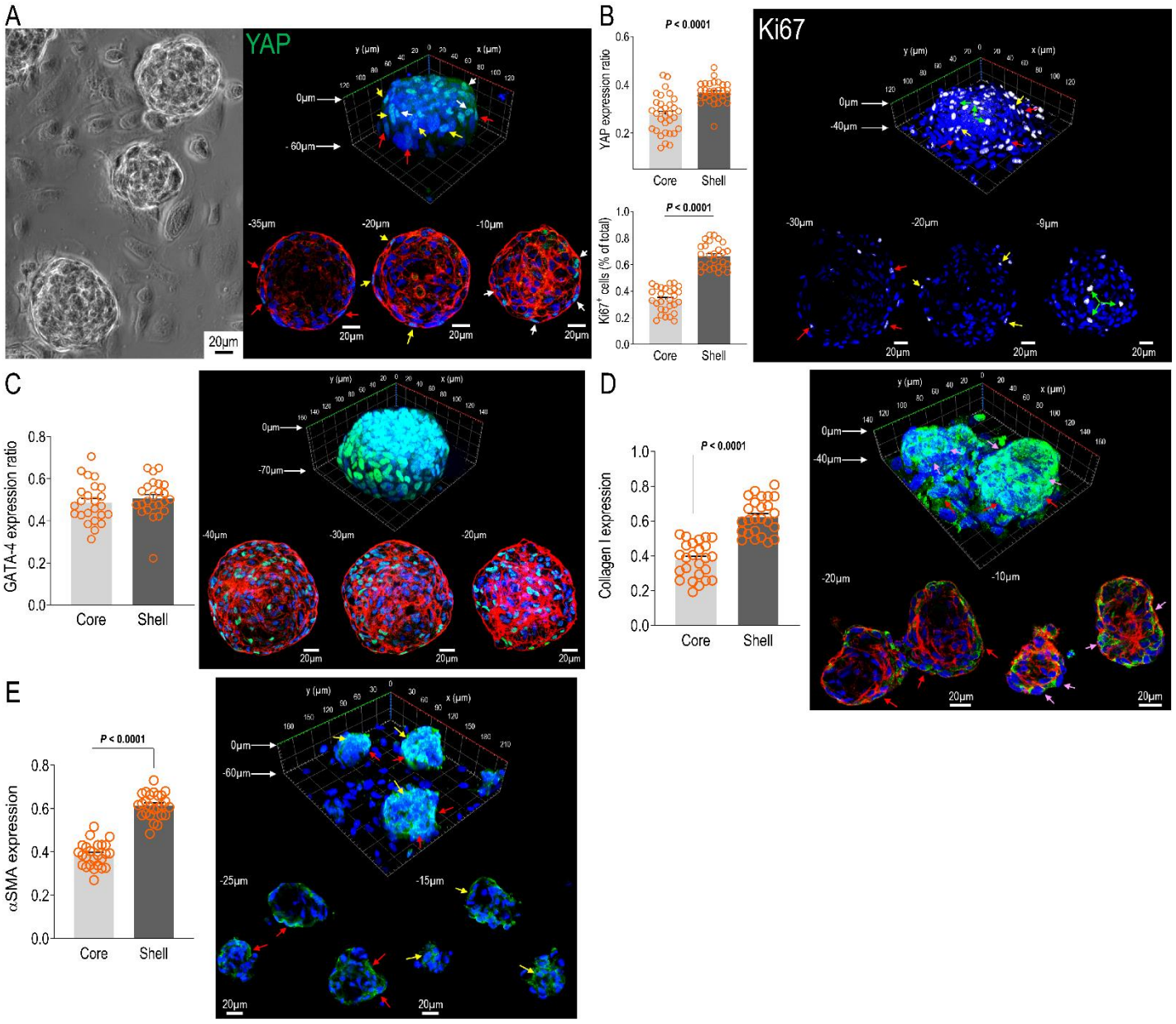

A
YAP
0µm
- 60µm
-35µm
-20µm
-10µm
20µm
B
P < 0.0001
YAP expression ratio
Core
Shell
P < 0.0001
Ki67+ cells (% of total)
Ki67
0µm
-40µm
-30µm
-20µm
-9µm
C
GATA-4 expression ratio
0µm
-70µm
-40µm
-30µm
-20µm
D
P < 0.0001
Collagen I expression
0µm
-40µm
-20µm
-10µm
E
P < 0.0001
αSMA expression
0µm
-60µm
-25µm
-15µm

Figure 3

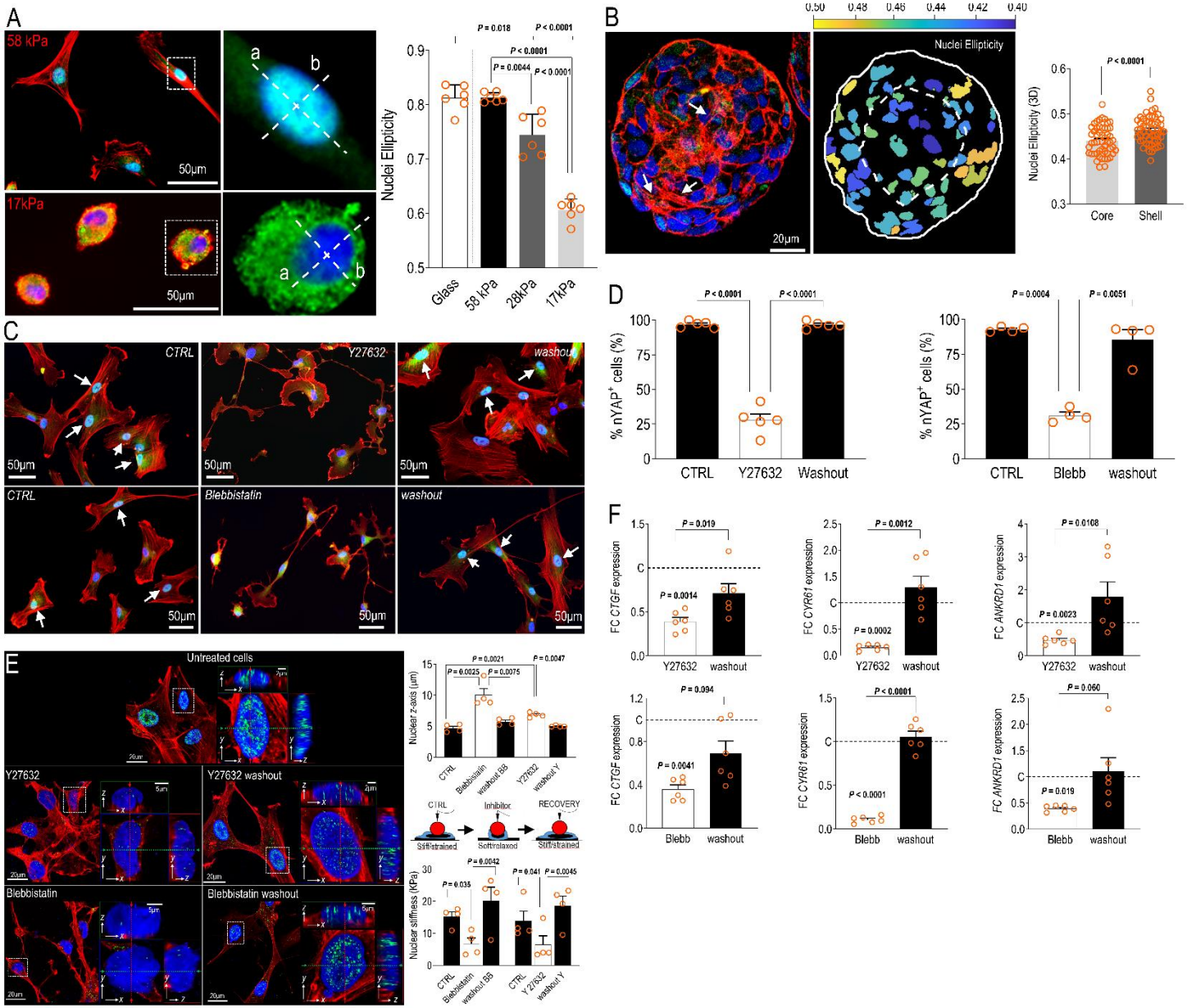
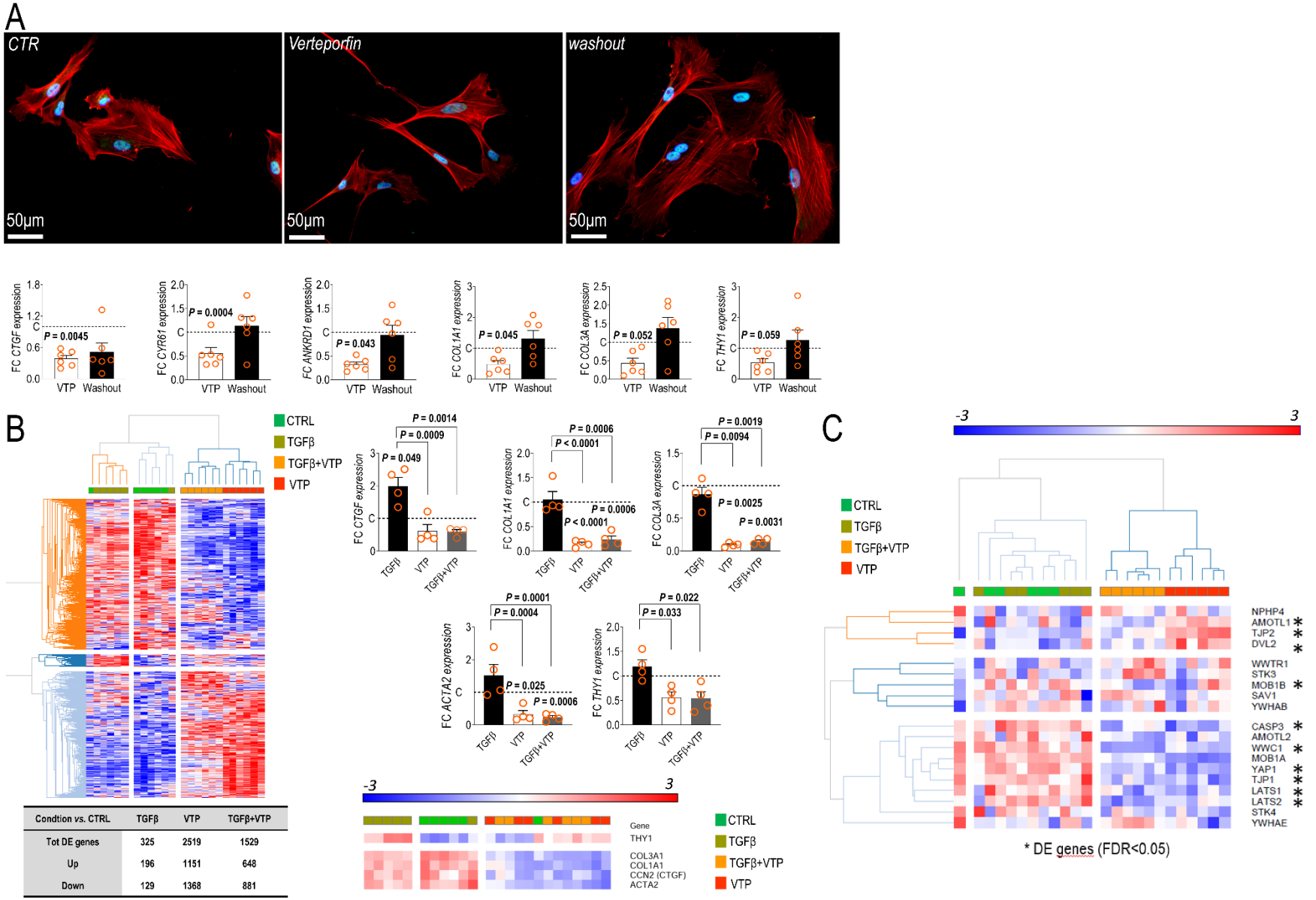

A
58 kPa
17kPa
50µm
a
b
Nuclei Ellipticity
Glass
58 kPa
28kPa
17kPa
B
Nuclei Ellipticity
20µm
Nuclei Ellipticity (3D)
Core
Shell
C
CTRL
Y27632
washout
Blebbistatin
D
% nYAP+ cells (%)
CTRL
Y27632
Washout
Blebb
E
Untreated cells
Y27632
Y27632 washout
Blebbistatin
Blebbistatin washout
Nuclear z-axis (µm)
Nuclear stiffness (kPa)
Inhibitor
RECOVERY
Stiff/strained
Soft/relaxed
F
FC CTGF expression
FC CYR61 expression
FC ANKRD1 expression

Figure 4



Condtion vs. CTRL	TGFβ	VTP	TGFβ+VTP
Tot DE genes	325	2519	1529
Up	196	1151	648
Down	129	1368	881

Figure 5

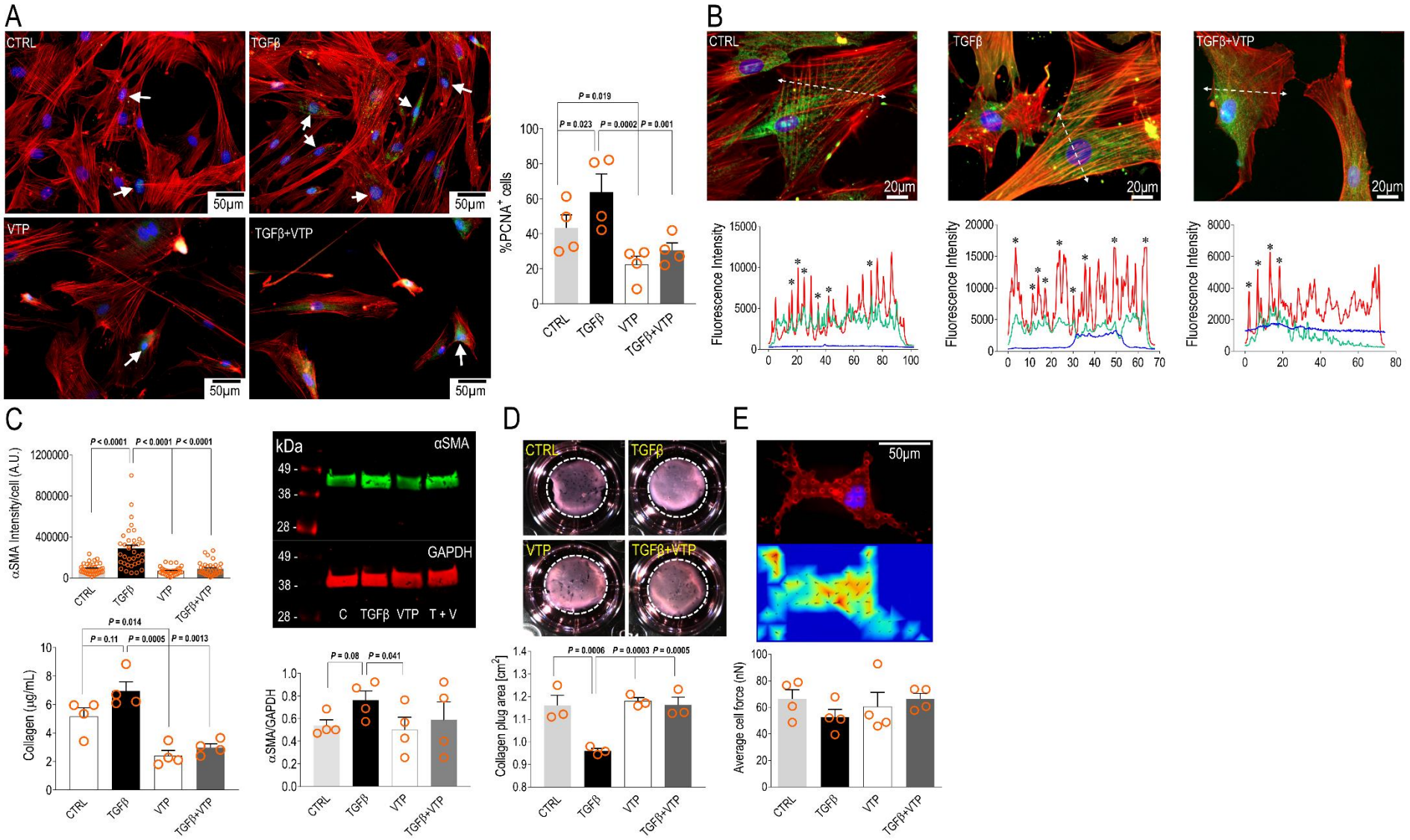

A
CTRL
TGFβ
VTP
TGFβ+VTP
50µm
%PCNA+ cells
P = 0.019
P = 0.023
P = 0.0002
P = 0.001
B
20µm
Fluorescence Intensity
C
αSMA Intensity/cell (A.U.)
P < 0.0001
Collagen (µg/mL)
P = 0.014
P = 0.11
P = 0.0005
P = 0.0013
kDa
αSMA
GAPDH
C TGFβ VTP T + V
αSMA/GAPDH
P = 0.08
P = 0.041
D
Collagen plug area [cm2]
P = 0.0006
P = 0.0003
P = 0.0005
E
Average cell force (nN)

Figure 6

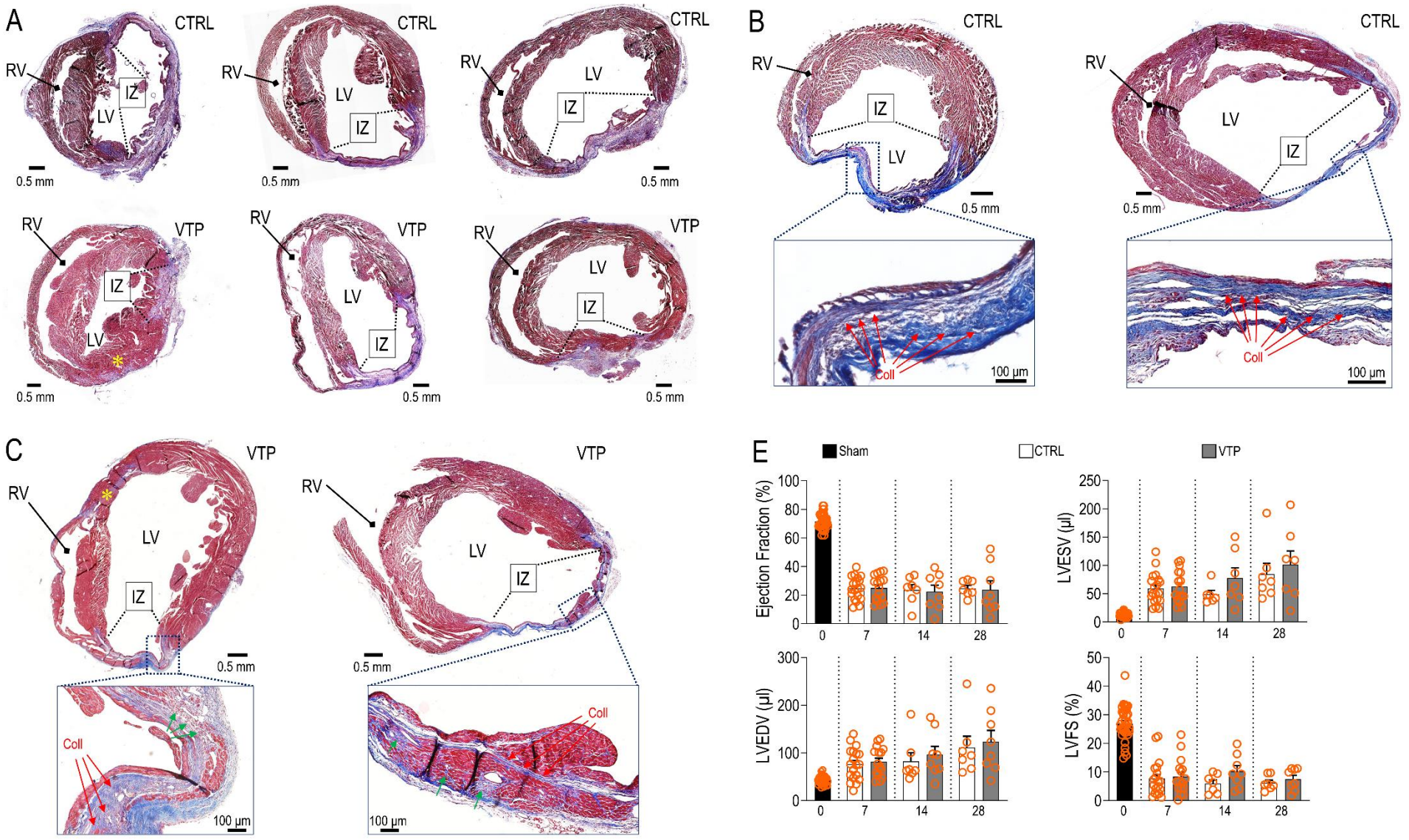

A
CTRL
RV
LV
IZ
0.5 mm
VTP
B
Coll
100 µm
C
E
Sham
CTRL
VTP
Ejection Fraction (%)
LVESV (µl)
LVEDV (µl)
LVFS (%)
0
7
14
28

Figure 7

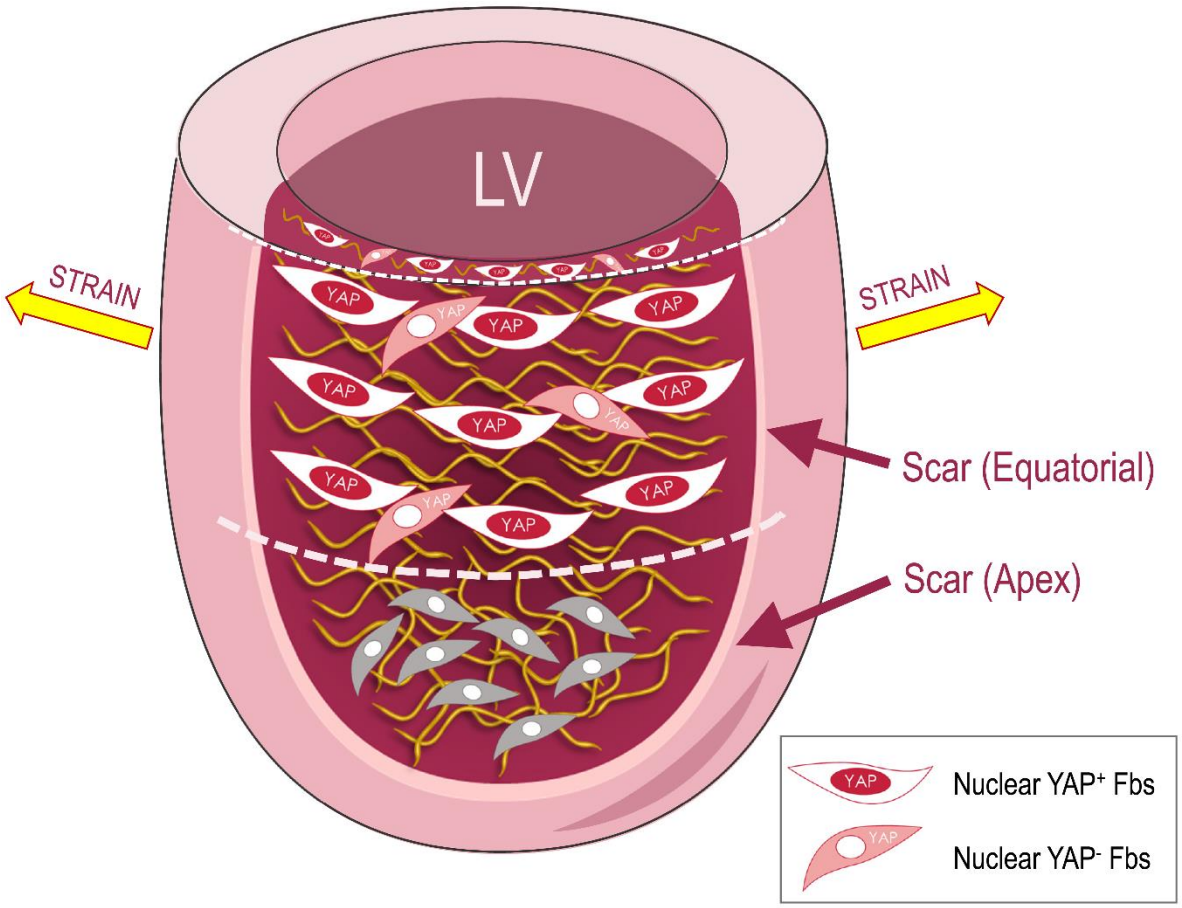

LV
STRAIN
STRAIN
YAP
Scar (Equatorial)
Scar (Apex)
Nuclear YAP+ Fbs
Nuclear YAP- Fbs

Figure 8

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