

Supplementary Figure. S5 Matrix stiffness activates lipolysis in HSCs. (A) Representative images (left) and quantification (right) of lipid level in JS-1 cells cultured on 1kPa, 10kPa and 25kPa polyacrylamide hydrogels. Scale bar: 5μm (one-way ANOVA, n=5 independent experiments). **(B)** Relative mRNA levels of lipolysis related genes in LX-2 (upper) and JS-1 cells (lower) cultured on 1kPa, 10kPa and 25kPa polyacrylamide hydrogels (one-way ANOVA, n=5 independent experiments). **(C)** Protein levels of lipolysis related genes in cultured on 1kPa, 10kPa and 25kPa polyacrylamide hydrogels. **(D)** Protein levels of ATGL in LX-2 (upper) and JS-1 cells (lower) on indicated stiffness after treated with DMSO or 10μM atglistatin (ATGLi) for 24 h. **(E)** FFA content in supernatant from LX-2 (upper) and JS-1 cells (lower) on indicated stiffness after treated with DMSO or 24 h (one-way ANOVA).

n=5 independent experiments). (F) Representative images (left) and quantification (right) of lipid level in JS-1 cells cultured on indicated stiffness after treated with DMSO or 10 μ M atglistatin (ATGLi) for 24 h. Scale bar: 5 μ m (one-way ANOVA, n=5 independent experiments). (G) Representative images (left) and quantification (right) of labelled lipids in MC38 cells after co-culturing with JS-1 cells with indicated treatment. Scale bar: 5 μ m (one-way ANOVA, n=5 independent experiments). (H) Fatty acid oxidation rate of MC38 cells treated with indicated conditioned medium from JS-1 cells (one-way ANOVA, n=5 independent experiments). (I) Protein levels of Cd36, Ppar γ , Acsl1, Cpt1a and Cpt1b in MC38 cells treated with indicated conditioned medium from JS-1 cells. Data are graphed as the mean \pm SD. **P* < 0.05; ***P* < 0.01; ****P* < 0.001.