

Figure S1. Relative levels of **A)** collagen type I (*COL1A1*), **B)** collagen type III (*COL3A1*) mRNA in endometrial fibroblasts treated with 1 μ M, 5 μ M of 5-aza-dC or untreated fibroblasts (control), **C)** collagen type I (*COL1A1*) and **D)** collagen type III (*COL3A1*) mRNA in non-treated (control) endometrial fibroblasts or treated with TGF- β 1 (10ng/mL), (1 μ M), TGF- β 1 (10ng/mL) + 1 μ M 5aza-dC or TGF- β 1 (10ng/mL) + 5 μ M 5aza-dC; $n=4$. Bars represent mean \pm SEM. Asterisks indicate significant differences between treatments (* $P<0.05$, ** $P<0.01$, *** $P<0.001$).

1st Protocol

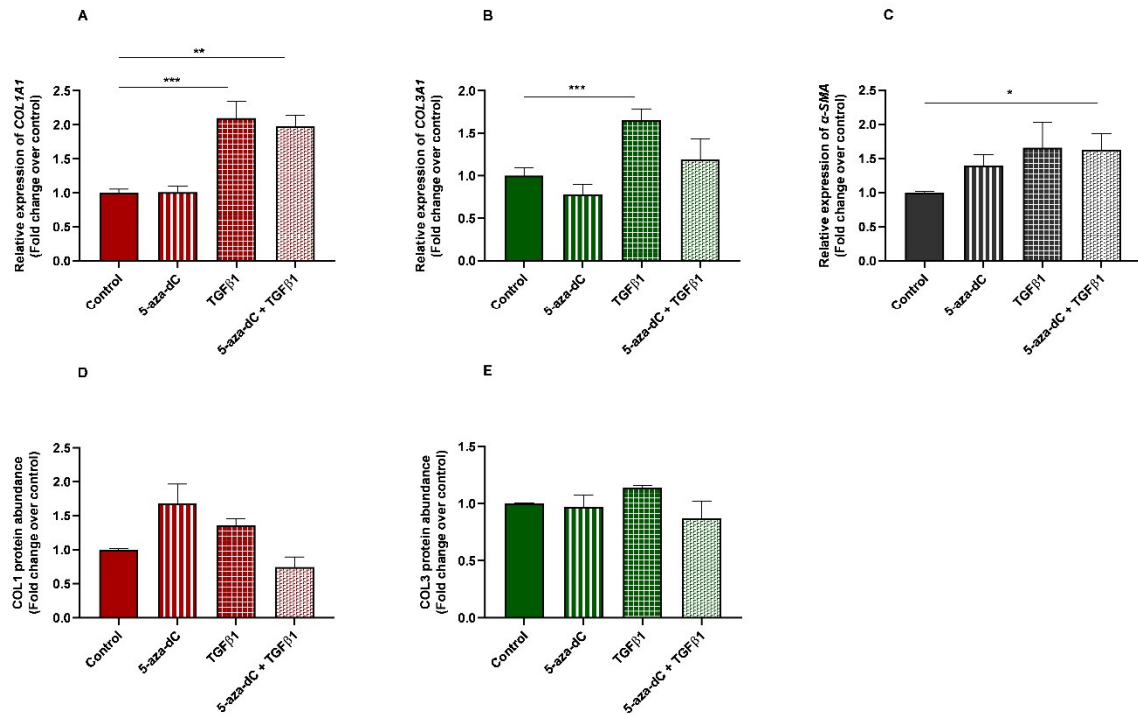


Figure S2. Relative levels of **A)** collagen type I (*COL1A1*), **B)** collagen type III (*COL3A1*), and **C)** α -smooth muscle actin (α -SMA) mRNA, and of **D)** COL1 and **E)** COL3 protein concentrations in non-treated (control) endometrial fibroblasts or treated with 5-aza-dC (1 μM), TGF-β1 (10 ng/mL) or TGF-β1 (10 ng/mL) + 5-aza-dC (1 μM) for 48h. Each treatment was compared to respective control (all groups with control C and TGF-β1+5-aza-dC with TGF-β1); $n=5$. Bars represent mean \pm SEM. Asterisks indicate significant differences between treatments (* $P<0.05$, ** $P<0.01$, *** $P<0.001$).

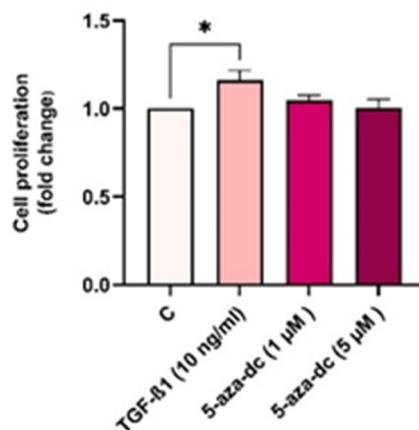


Figure S3. Effect of TGF-β1 at a dose of 10 ng/mL and 5-aza-dC at doses of 1 μM and 5 μM on cell viability, $n=4$. Bars represent mean \pm SEM. Asterisks indicate significant differences between treatment (* $P<0.05$). C-control (no factors added to the culture medium).