

Short title: Genetic targeting of myofibroblasts

Supplementary Figure S1: Reporter activity of deletion constructs of the upstream region of SMA gene in SMA-negative L6 myoblasts, cultivated in 5% FCS in DMEM and in SMA-positive L6 myotubes, induced to differentiate into myotubes by 48 h serum deprivation (0.5 % FCS in DMEM).

Supplementary Figure S2: Mesangial cells myofibroblastically induced by high glucose treatment (4.5 g glucose L⁻¹ culture medium) and transduced with pAd-SMA-GFP driving reporter expression by the artificial SMA-promoter hybrid, shown in figure 2A (a-c). a) DAPI stained, b) SMA immunostained and c) GFP reporter fluorescence.

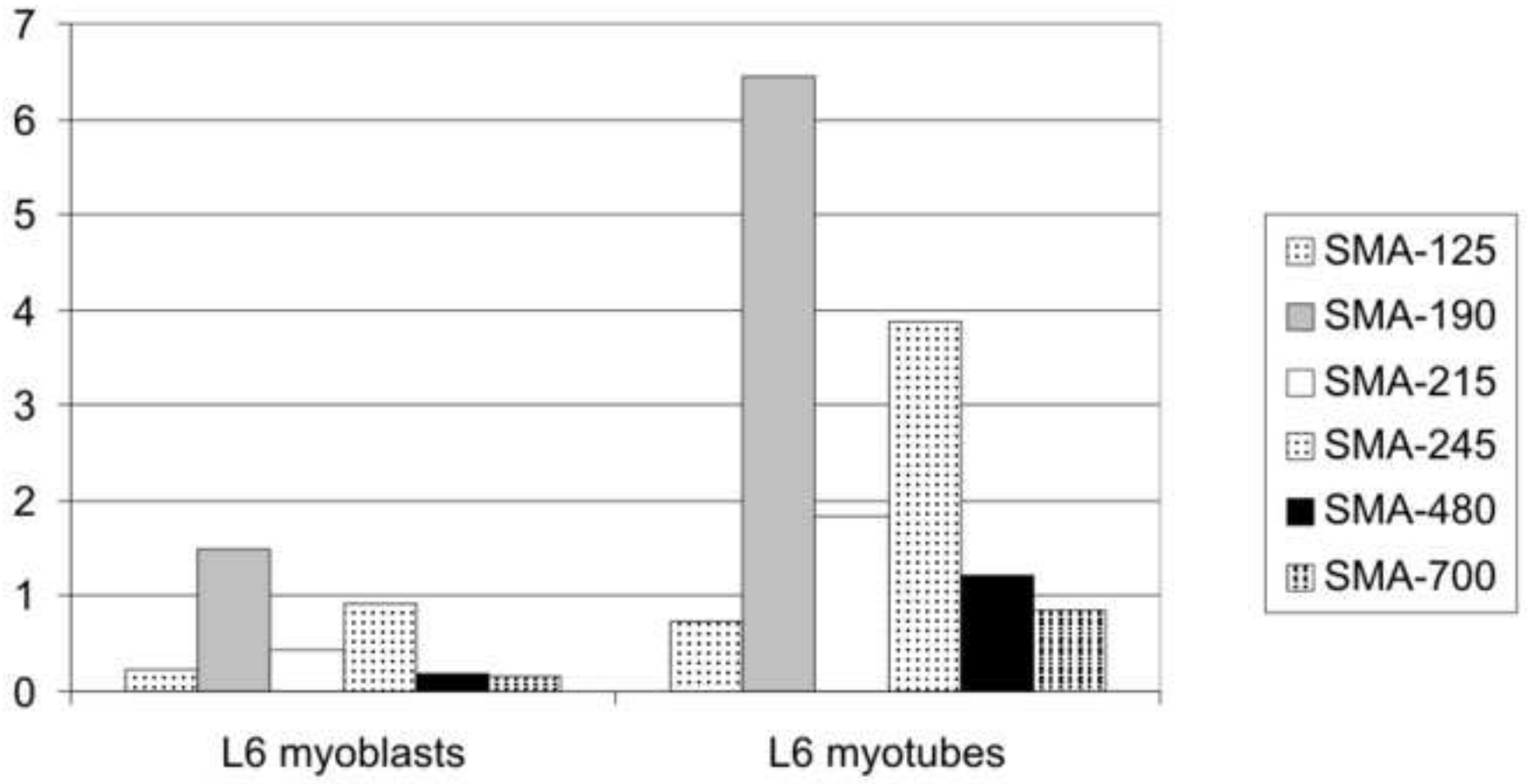


Figure S1

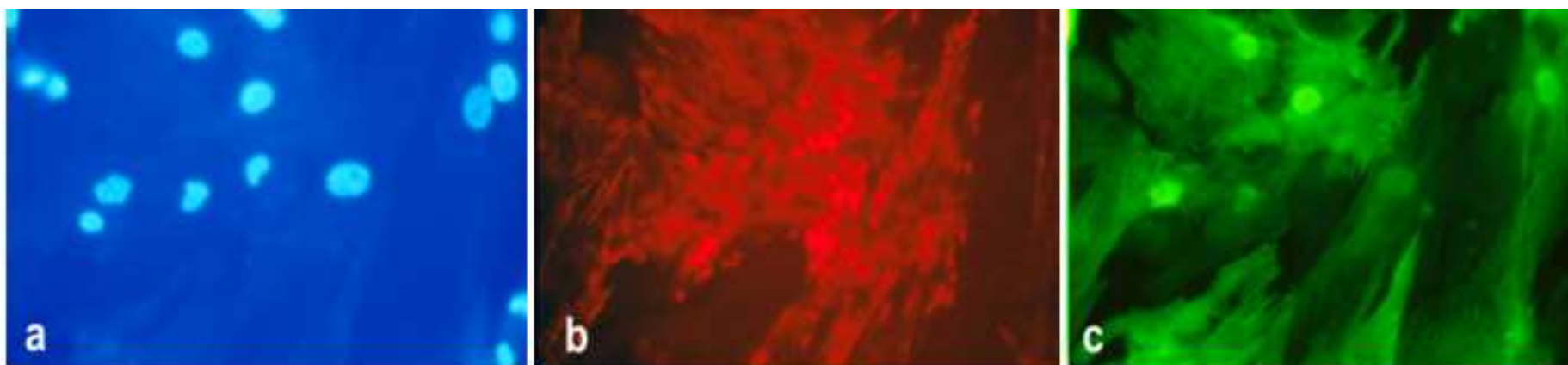


Figure S2