S5 Fig. Examination of gene expression using qRT-PCR. (A) Relative expression of probasin from Gli1-CreER driven GFP expressing cells and epithelial cells isolated from prostates of either R26mTmG/+;Gli1CreER+/+ or R26mTmG/+;ArLY/Gli1CreER+/+ mice. Both Gli1CreER driven GFP expressing cells and prostatic epithelial cells were isolated and sorted by GFP or CD24 antibody, respectively. RNA samples were prepared and used to generate cDNA. The relative expression levels from three individual experiments were shown. (B-C) Fold changes in labeled expression of genes determined by qRT-PCR analysis using FACS-sorted GFP positive cells from either UGM tissues at day E16.5 (B) or prostate tissues at postnatal day 56 (C) isolated from R26mTmG/+;Gli1CreER+/+ and ArLY/Gli1CreER+/+ or R26mTmG/+;ArLY/Gli1CreER+/+ mice. Error bars indicate s.d.; *P < 0.05, ** P < 0.01; analyzed using 2-tailed students’ t test. (n = 3 replicates per data point).