Supplemental Figures

p130Cas IS REQUIRED FOR MAMMARY TUMOR GROWTH AND TRANSFORMING GROWTH FACTOR-β (TGF-β)-MEDIATED METASTASIS THROUGH REGULATION OF SMAD2/3 ACTIVITY

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Supplemental Figures:

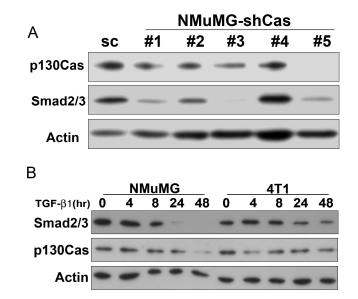


Figure S1: M.K. Wendt, J.A. Smith, and W.P. Schiemann

FIGURE S1. **p130Cas-deficiency decreases Smad2 expression in normal MECs.** *A*, Five unique shRNA sequences targeting p130Cas (shCas #1-5) were stably expressed in NMuMG cells, and differences in p130Cas expression were analyzed by immunoblotting with anti-p130Cas antibodies (sc, scrambled shRNA). β -actin immunoreactivity (Actin) is provided as a loading control. *B*, Quiescent NMuMG and 4T1 cells were stimulated with TGF- β 1 (5 ng/ml) for varying times, and the resulting whole-cell extracts were immunoblotted with Smad2/3 antibodies, followed by p130Cas and β -actin (Actin) as a loading control. Data are representative of 2 independent experiments and show that Smad2/3 are degraded more readily upon prolonged TGF- β treatment in NMuMG cells as compared to 4T1 cells.

Figure S2: M.K. Wendt, J.A. Smith, and W.P. Schiemann

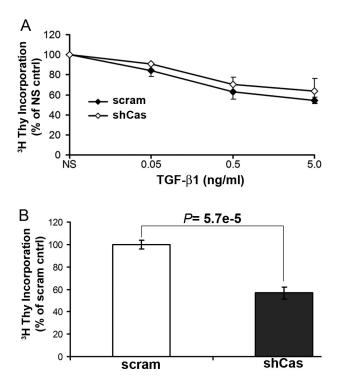


FIGURE S2. **p130Cas-deficiency decreases basal 4T1 proliferation but does not affect TGF-\beta1growth inhibition when cultured on plastic.** *A***, Control (scram) and p130Cas-deficient (shCas) cells were stimulated with the increasing TGF-\beta1 concentrations for 48 h as indicated. Afterward, [³H]thymidine incorporation into cellular DNA was determined. Data are the mean (± SE; n=3) [³H]thymidine incorporation normalized to untreated controls (NS, no stimulation).** *B***, Data are the mean (± SE; n=3) basal [³H]thymidine incorporation normalized to control (scram) 4T1 cells.**