Article

Peroxiredoxin V (PrdxV) negatively regulates EGFR/Stat3-mediated fibrogenesis via a Cys48-dependent interaction between PrdxV and Stat3

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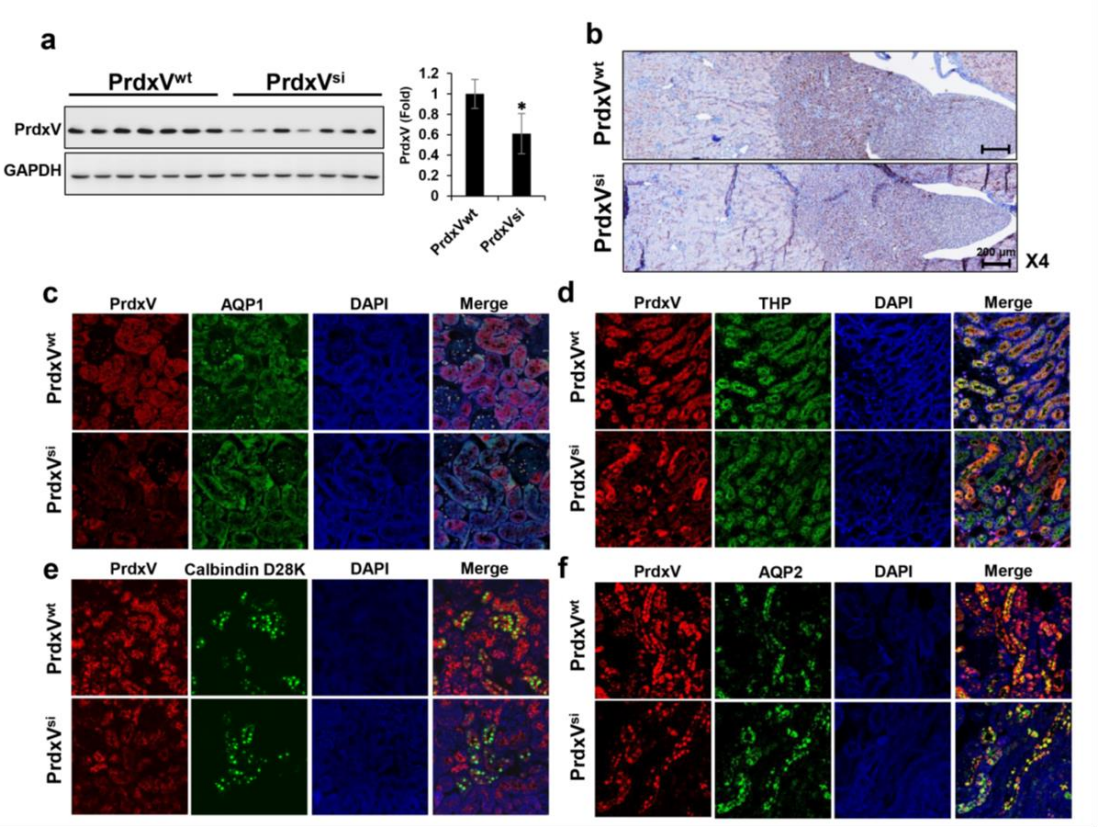
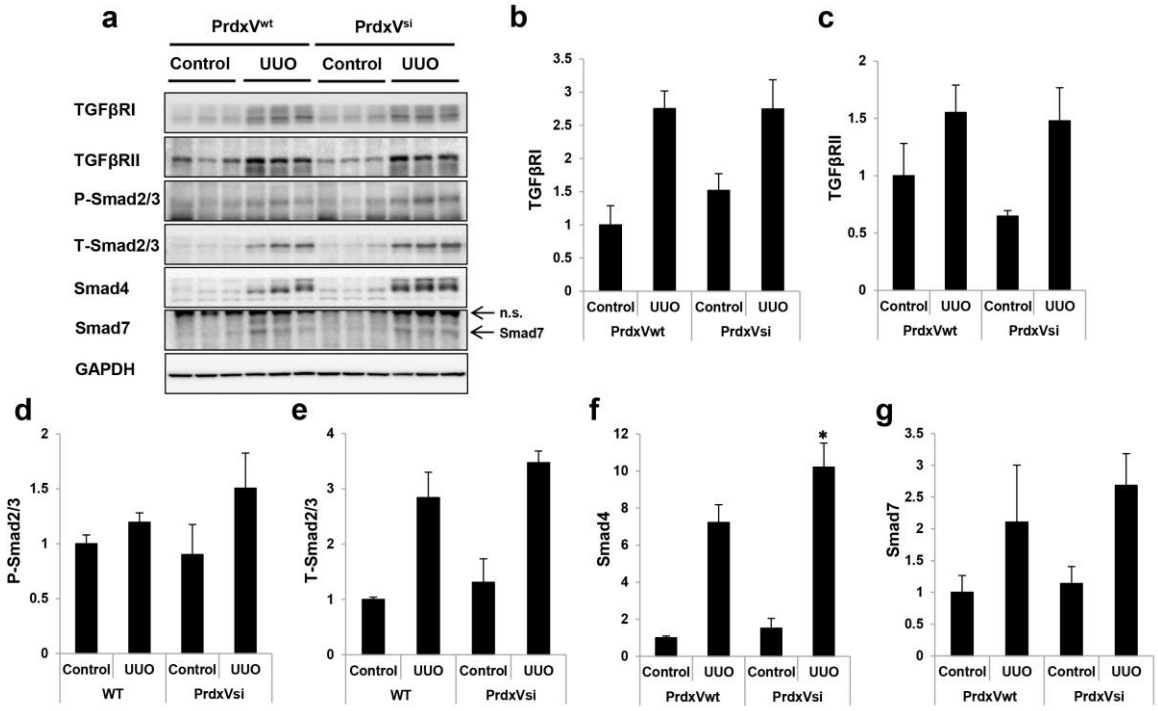
Supplementary Figures.

Supplementary Figure. S1



Comparison of PrdxV expression between PrdxV^{wt} and PrdxV^{si} mouse kidney. **(a)** Total protein level of PrdxV were assessed by western blotting. GAPDH was used as an internal control. Bar graphs show the mean PrdxV protein/GAPDH expression as measured by densitometry. $*p < 0.05$; PrdxV^{wt} *vs.* PrdxV^{si}. **(b)** Light micrographs illustrating immunostaining of PrdxV of PrdxV^{wt} and PrdxV^{si} kidney. Immunolabeling is present from cortex to inner medullar. Image was magnified at x4, Bar = 200 µm. **(c-f)** Tubule specific expression of PrdxV (Red) were assessed by tubule markers (Green), AQP1, THP, Calbindin D28K, and AQP2, using immunofluorescence. Nucleus was stained with DAPI. Image was magnified at x400.

Supplementary Figure. S2



Canonical TGF-β signaling pathway in UUO-induced PrdxV^{si} mouse kidney. To verify the involvement of the canonical TGF-β signaling pathway in renal fibrosis aggravated by knockdown of PrdxV, the expression levels and activation levels of major TGF-β/Smad signaling molecules were assessed by western blotting **(a)**. GAPDH was used as an internal control. **(b-e)** Bar graphs show the mean target protein/GAPDH expression as measured by densitometry (**a**; TGFβRI, **c**; TGFβRI, **d**; Phospho-Smad2/3, **e**; Total-Smad2/3).

Uncropped western blotting data

Figure 2

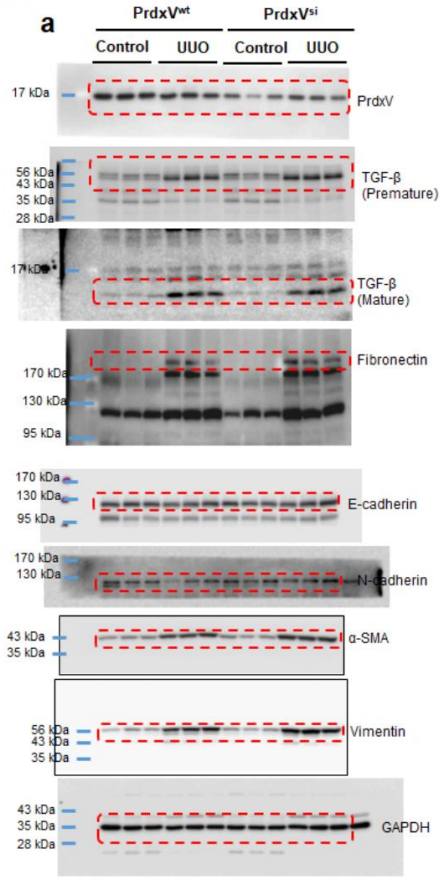


Figure 3

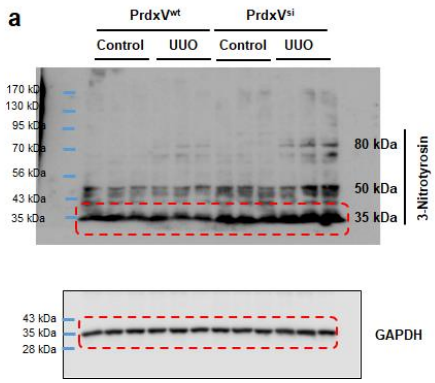


Figure 4

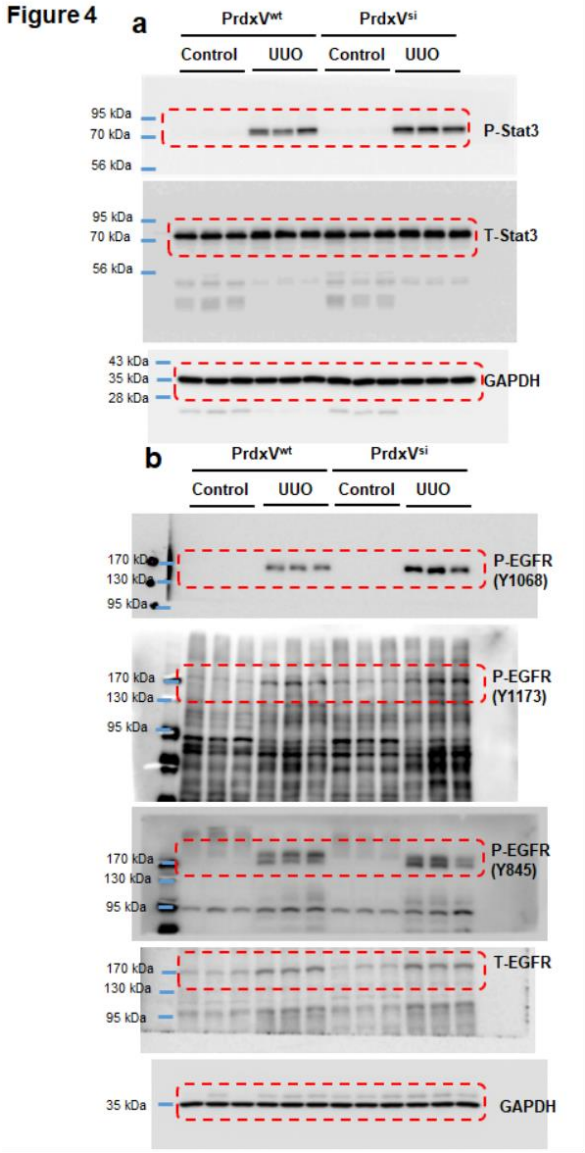


Figure 5

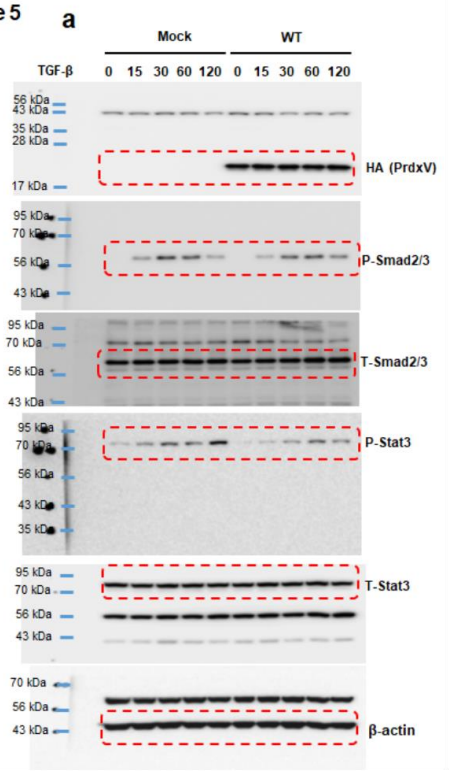


Figure 5

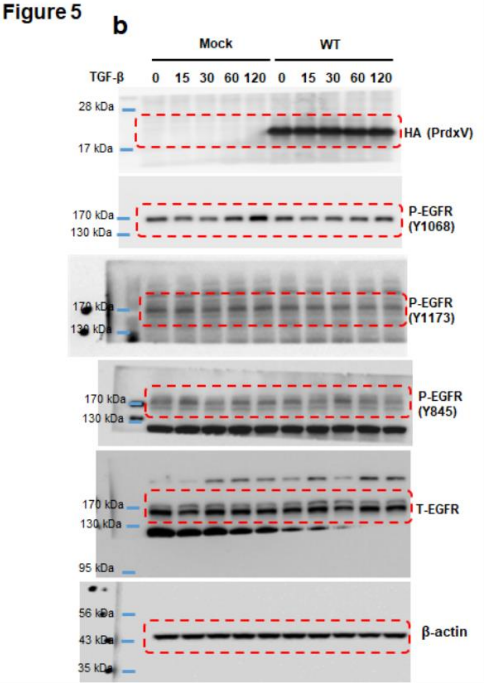


Figure 6

a

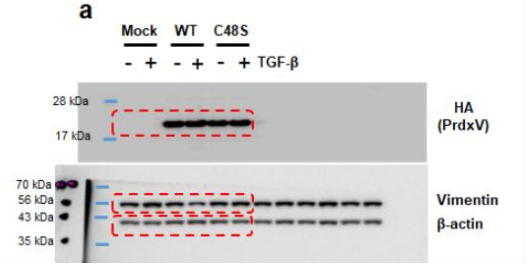
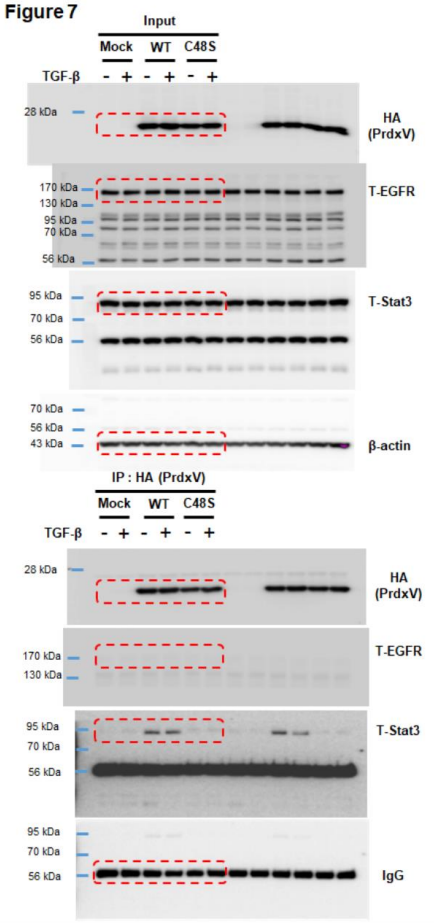
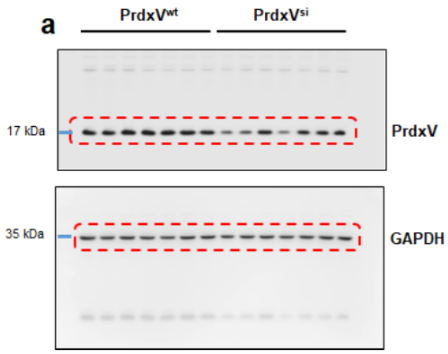


Figure 7



Supplementary Figure S1



Supplementary Figure S2.