



Supplementary Figure S1. Regulation of sodium and potassium homeostasis in 9month-old $Fgf23^{-/}$ /VDR^{Δ/Δ} and $K\Gamma^{/-}$ /VDR^{Δ/Δ} compound mutants. (A) Western blotting quantification of renal membrane expression of the full length β and γ subunits of ENaC in 9month-old male wild-type (WT), VDR^{Δ/Δ}, $Fgf23^{-/-}$ /VDR^{Δ/Δ} and $K\Gamma^{-/-}$ /VDR^{Δ/Δ} double mutant mice on the rescue diet (n=8-10, 1-way ANOVA followed by SNK test, * p < 0.05 vs. WT). (B) Serum Na⁺ concentration, serum potassium (K⁺) concentration, urinary potassium excretion corrected by urinary creatinine (Crea), urinary pH and urinary volume over a 12hour sampling period in 9-month-old male wild-type (WT), VDR^{Δ/Δ}, *Fgf23^{-/-}*/VDR^{Δ/Δ}, or *Kl^{-/-}*/VDR^{Δ/Δ} compound mutant mice (n=8-12, 1-way ANOVA followed by SNK test, * *p* < 0.05 vs. WT). (C) Mean food consumption of 9-month-old male wild-type, VDR^{Δ/Δ}, *Fgf23^{-/-}*/VDR^{Δ/Δ}, and *Kl^{-/-}*/VDR^{Δ/Δ} compound mutant mice measured over a period of 7 days (n=6-8). (D) Western blotting quantification of NCC phosphorylation at Ser71, Ser91 and Thr55 (pNCC S71, S91 and T55) in renal cortical total membrane fractions (n=6-8) and (E) plasma renin activity assay in 9-month-old male wild-type, VDR^{Δ/Δ}, *Fgf23^{-/-}*/VDR^{$\Delta/\Delta}, and$ *Kl^{-/-}* $/VDR^{<math>\Delta/\Delta}, and$ *Kl^{-/-}* $/VDR^{<math>\Delta/\Delta} compound mutant mice (n=7-8, 1-way ANOVA followed by SNK test, *$ *p*< 0.005 vs. WT). Data represent mean ± s.e.m</sup></sup></sup>