Gene	Taqman probe or primer sequences (5' to 3')
Ccl2 (Mcp1)	F: GCTTCTTTGGGACACCTGCTGC
	R: TTAACGCCCCACTCACCTGCTG
Colla2	F: AAGGGGTCTTCCTGGTGAAT
	R: GGGGTACCACGTTCTCCTC
Fnl	F: TGTGACCAGCAACACGGTG
	R: ACAACAGGAGAGTAGGGCGC
Hmox1(HO1)	F:ACATCGACAGCCCCACCAAGTTCAA
	R:CTGACGAAGTGACGCCATCTGTGAG
Mmp2	F: TCTGGGCAGTCTTGAACATTT
	R: AGAGTCAGGTGATGGATGTCG
Tgfb1	F: TGCGTCTGCTGAGGCTCAA
	R: TTGCTGAGGTATCGCCAGGA
Tnf	F: GCCAGCCGATGGGTTGTA
	R: GGCAGCCTTGTCCCTTGA
Atg5	Mm01187303_m1
Atg12	Mm00503201_m1
Becn1	Mm01265461_m1
<i>Emr1</i> (F4/80)	Mm00802529_m1
Hk1	Mm00439344_m1
Hk2	Mm00443385_m1
Illb	Mm00434228_m1
116	Mm01210733_m1
Nos2(iNOS)	Mm00435175_ml
P2rx7	Mm00440578_m1
Rn18S	Mm03928990_g1
Slc2a1	Mm00441480_m1
Slc16a3	Mm00446102_m1
Sqstm1/p62	Mm00448091_m1
Ulk1	Mm00437238_m1

Supplementary Table 1. Taqman probe ID and primer sequences

Supplementary Figure 1.



No effect of trehalose on the renal expression of autophagy markers in wild-type mice. The kidney lysates of 3-month-old wild-type mice were immunoblotted for (A) LC3, (B) p62, and β -actin. Graphs represent mean \pm standard error of the mean (n = 3 per group). WT, wild-type; C, control; S, sucrose; T, trehalose.

Supplementary Figure 2.



Pkd1 miR Tg+Sucrose

Pkd1 miR Tg+Trehalose

Immunofluorescence staining for p62 protein in cystic epithelial cells. Note p62 staining (shown in red, arrows) of the cystic epithelial cells in *Pkd1* miR Tg mice, indicating reduced autophagic degradation. The nuclei are counterstained with DAPI (blue). No significant differences in the fluorescent p62 signal were found among the different treatment groups. Representative merged images are shown (n=3 per group). Scale bar, 50 μ m. WT, wild-type; PKD, polycystic kidney disease. Original magnification, 400×.