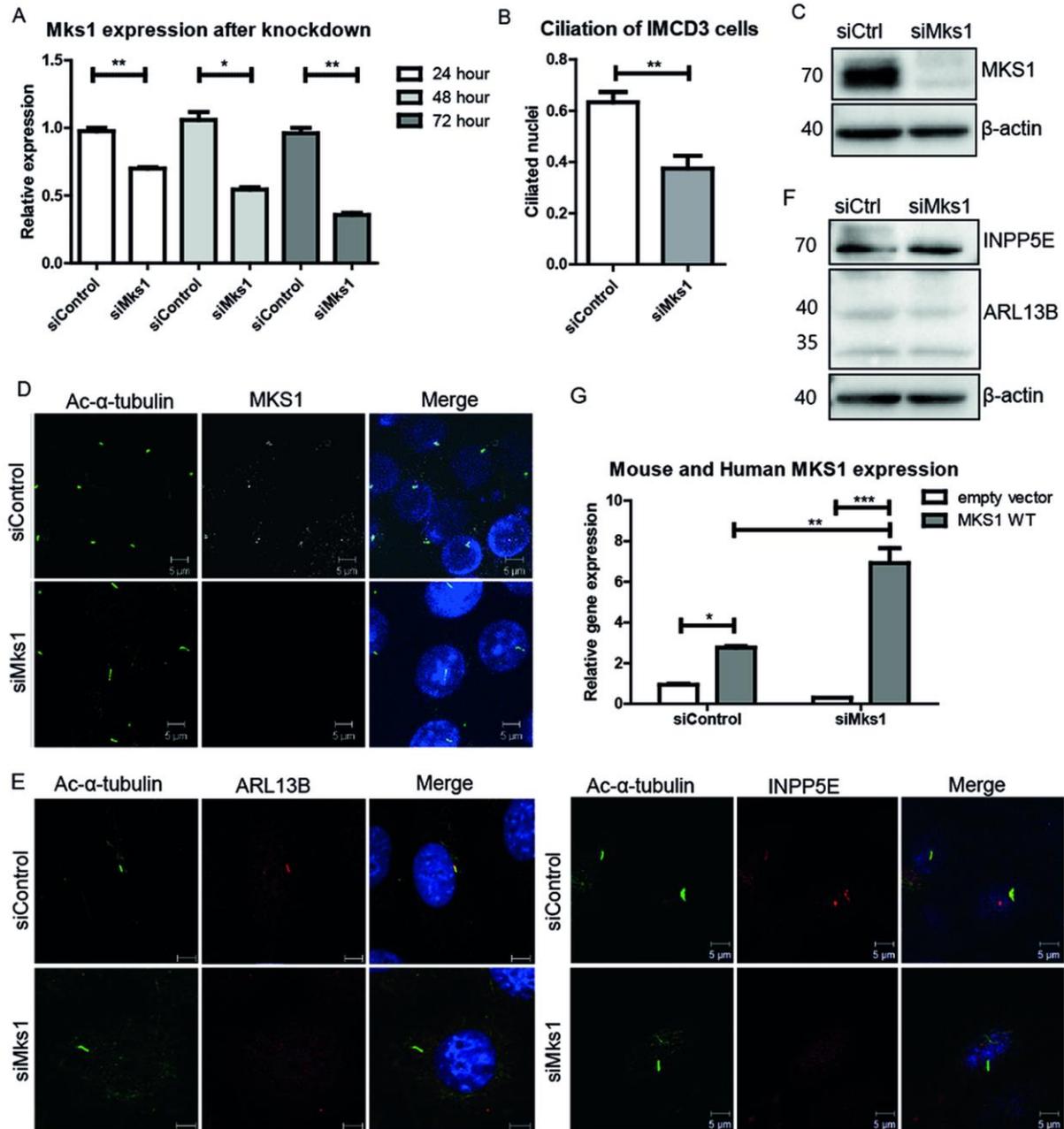
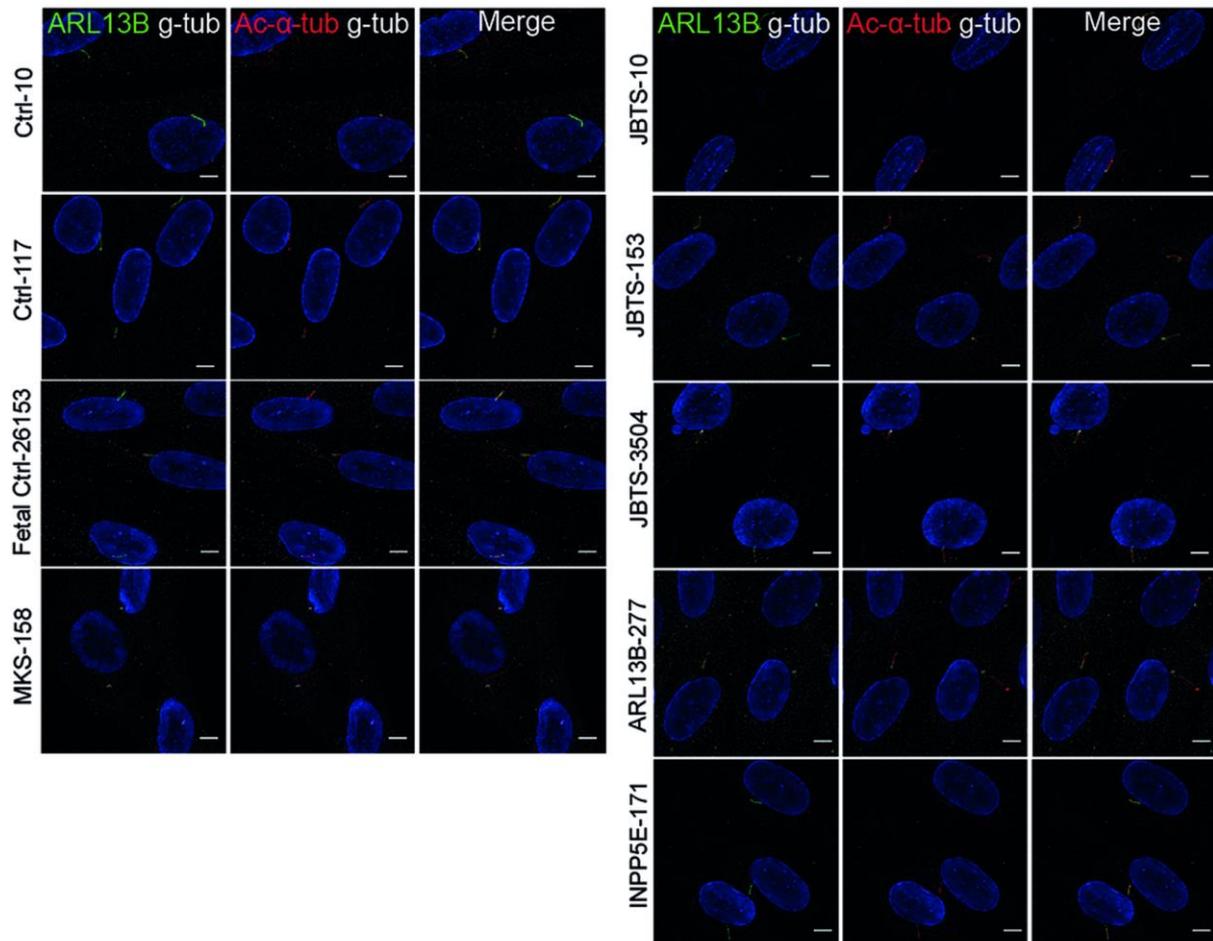


## SUPPORTING INFORMATION LEGENDS

Figure S1. Flow diagram for determining ARL13B and INPP5E protein content in cilia of fibroblasts.

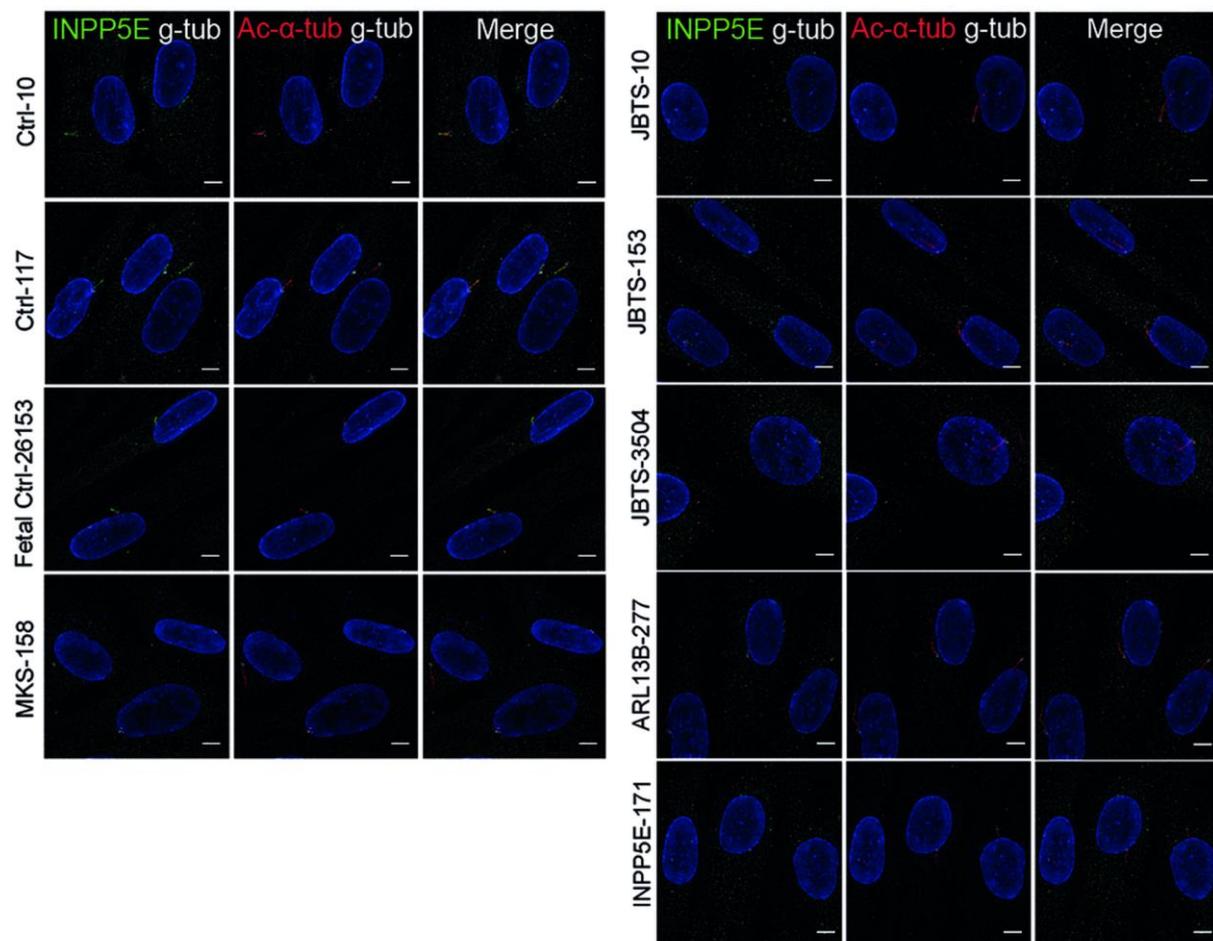


**Figure S2. ARL13B cilia staining of fibroblasts.** Original deconvoluted images of immunostaining of fibroblasts derived from skin biopsies of JBTS-10, JBTS-153, JBTS-3504, MKS-158, INPP5E-171, ARL13B-277 and controls. ARL13B (green), gamma tubulin (g-tub; white) and cilia (acetylated tubulin, red; scale bar 5  $\mu$ m).

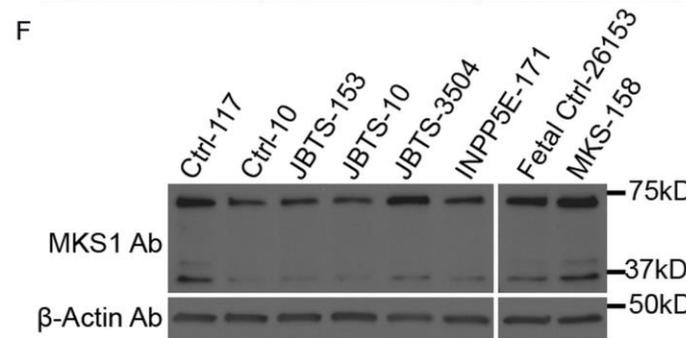
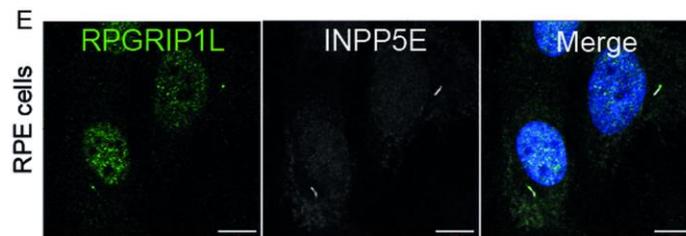
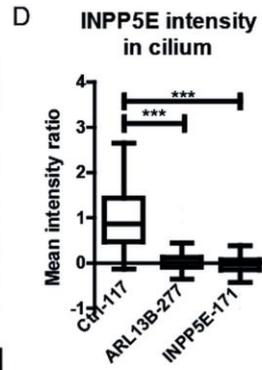
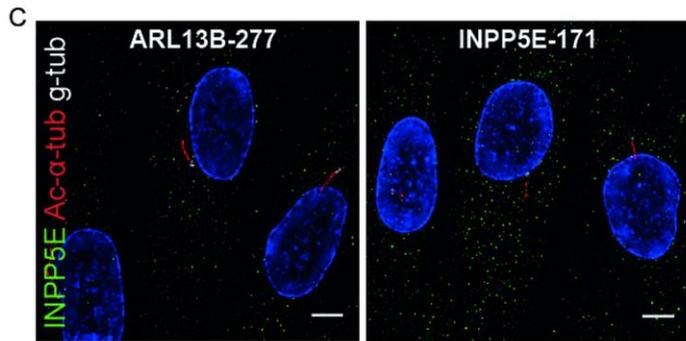
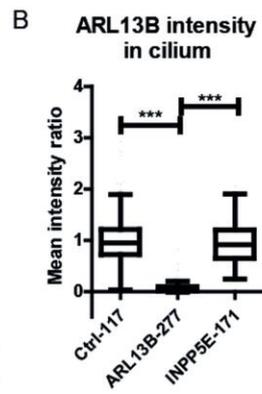
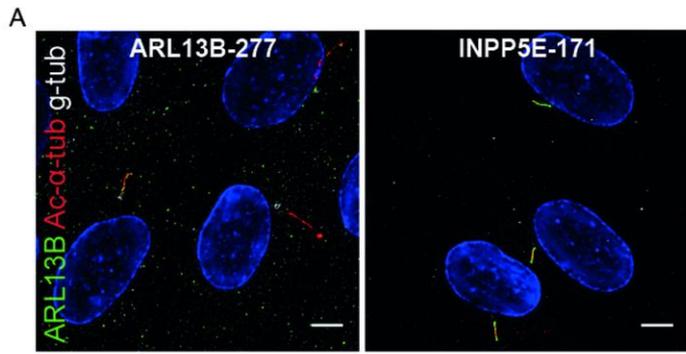


**Figure S3. siRNA knockdown of *Mks1* in IMCD3 cells results in decreased MKS1 protein, fewer cilia and decreased ciliary ARL13B and INPP5E levels.** (A) RT-QPCR detects lower mRNA levels of *Mks1* after siRNA depletion compared to control siRNA transfected IMCD3 cells ( $p < 0.02$ ). Error bars represent SEM ( $n = 3$ ). (B) Quantification of cilia frequency in IMCD3 cells treated with control siRNA or *Mks1* siRNA for 72 hours ( $p < 0.004$ ). Error bars represent SEM ( $n = 3$ ). (C) Immunoblot of MKS1 of IMCD3 lysates transfected with siControl or si*Mks1* oligonucleotides for 56 hours. Less MKS1 protein is

detected in si*Mks1*-treated versus siControl-treated IMCD3 cells.  $\beta$ -actin is used as loading control. **(D)** Immunostaining of IMCD3 cells treated with siControl or si*Mks1* for 48 hours. Basal body (MKS1, white) and cilia (acetylated tubulin, green) staining shows less MKS1 protein at the base of primary cilia in si*Mks1*-treated versus siControl-treated IMCD3 cells. **(E)** Immunostaining of IMCD3 cells treated with siControl or si*Mks1* for 72 hours. ARL13B and INPP5E cilia staining (red) does not colocalize with cilia (acetylated tubulin, green) in si*Mks1* treated cells. **(F)** Immunoblot of ARL13B and INPP5E of IMCD3 lysates transfected with siControl or si*Mks1* oligonucleotides for 56 hours. ARL13B and INPP5E protein levels are unchanged by MKS1 depletion.  $\beta$ -actin is used as loading control. **(G)** RT-QPCR with primers recognizing mouse *Mks1* and human *MKS1* detects higher levels of human *MKS1* expression in siControl and si*Mks1* oligonucleotide (56 hours) treated IMCD3 cells transfected (32 hours) with wild-type MKS1 allele (Two-way ANOVA, Bonferroni test \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ ). Error bars represent SEM (n = 3).



**Figure S4. Reduced ciliary ARL13B and INPP5E in fibroblasts from individuals with ARL13B- and INPP5E-related Joubert syndrome. (A)** Immunostaining of fibroblasts derived from skin biopsies of ARL13B-277 and INPP5E-171. ARL13B (green), gamma tubulin (g-tub; white) and cilia (acetylated tubulin, red; scale bar 5  $\mu$ m). Brightness and contrast were identically adjusted across photos for visualization purposes; original data is in Figure S2. **(B)** Only ARL13B-277 fibroblasts have less ARL13B in the cilium than control (Tukey whiskers). \*\*\* indicates  $p < 0.001$  (Kruskal-Wallis test,  $n > 100$  cilia in 2 batches, see Methods and Fig S1 for details). **(C)** Immunostaining of fibroblasts derived from skin biopsies of ARL13B-277 and INPP5E-171. INPP5E (green), gamma tubulin (white) and cilia (acetylated tubulin, red; scale bar 5  $\mu$ m). Brightness and contrast were identically adjusted across photos for visualization purposes; original data is in Figure S5. **(D)** Both mutant fibroblasts have less INPP5E in the cilium than control (Tukey whiskers). \*\*\* indicates  $p < 0.001$  (Kruskal-Wallis test,  $n > 100$  cilia in 2 batches, see Methods and Fig S1 for details). Ctrl-117 images are included in Figure 2A (Fig S2) for ARL13B and 4A (Fig S5) for INPP5E. **(E)** Co-staining of INPP5E (white) with the TZ marker RPGRIP1L (green) in 48 hour serum starved RPE cells. No significant overlap of RPGRIP1L and INPP5E is observed (scale bar 10  $\mu$ m). **(F)** Immunoblot of MKS1 of lysates of fibroblasts derived from skin biopsies of JBTS-10, JBTS-153, JBTS-3504, MKS-158, INPP5E-171, and controls.  $\beta$ -actin is used as loading control. Densitometry was performed using the Gel Analysis functionality in Fiji ( $n=2$ ).



MKS1	1.00	0.53	0.57	0.50	1.08	0.65	1.04	1.28
Actin	1.00	1.08	1.06	1.19	0.80	1.07	1.16	1.16
MKS1/Actin	1.00	0.49	0.54	0.42	1.34	0.61	0.90	1.11

**Figure S5. INPP5E cilia staining of fibroblasts.** Original deconvoluted images of immunostaining of fibroblasts derived from skin biopsies of JBTS-10, JBTS-153, JBTS-3504, MKS-158, INPP5E-171, ARL13B-277 and controls. INPP5E (green), gamma tubulin (g-tub; white) and cilia (acetylated tubulin, red; scale bar 5  $\mu$ m).

