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 KP****C-6865
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 (R172H)

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MDA-PATC-148

(F134L)

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MDA-PATC-108

(R175H)

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MDA-PATC 108	R175H
MDA-PATC 148	F134L
PANC1	R273H
MiaPaCa-2	R248W
KP ^{wm} C-8865	R172H
KP ^{wm} C-6866	R172H



Supplementary Figure 1

Supplementary Figure 1. Characterization of the KP^{wm/+}C mouse model and derived cell lines. A and B, The overall survival and disease-specific survival between $KP^{Wm/+}C$ and $KP^{fl/+}C$ PDAC models as determined by the log-rank (Mantel-Cox) test. C and D, The overall survival and disease-specific survival between KPwm/+C and KPLSL/+C PDAC models as determined by the log-rank (Mantel-Cox) test. E, The timeline of PDAC progression is histologically similar between $KP^{wm/+}C$ and $KP^{fl/+}C$ PDAC models. Scale bars, 200µM. F, All murine and human PDAC cell lines underwent Sanger sequencing to confirm p53 missense mutations (listed) and to exclude the presence of wildtype p53 alleles. G, Transwell migration/invasion assays of PDAC cell lines derived from pancreas tumors in $KP^{wm/+}C$ mice and human PDAC cell lines (MDA-PATC) following mutant p53 knockdown. **H**, The effects of ectopic mutant p53^{R172H} expression on PDAC cell migration and invasion in p53-null, KP^{fl}C cell lines. I, GSEA analysis of hallmark pathways in $KP^{wm/+}C$ tumors reveals dysregulated pathways similar to human PDAC. (G, H) Data are guantified by 20X fields as mean±s.d. and pooled from at least two independent experiments performed in triplicate. P values determined by unpaired two-tailed t-tests.