



Supplementary Fig. S2. Loss of Kindlin-1 reduces tumour onset, survival associated with reduced integrin activation. (A) Kaplan–Meier analysis of tumor onset in NIC-Kin-1^{fl/fl} ($n = 18$) and NIC-Kin-1^{wt/wt} ($n = 15$) mice ($P < 0.0001$, Log rank test). (B, C) Kaplan-Meier survival curves for MT-Kin-1^{wt/wt} ($n=10$) and MT-Kin-1^{fl/fl} ($n=16$) ($P < 0.0001$, Log-rank test) (B) and NIC-Kin-1^{fl/fl} ($n=18$) and NIC-Kin-1^{wt/wt} ($n=15$) ($P < 0.0001$, Log-rank test) (C). (D) Western blot analysis of Kindlin-1 in MT-Kin-1^{wt/wt} and MT-Kin-1^{fl/fl} tumors and corresponding quantification on right ($P < 0.05$, Student's t -test). (E) Western blot analysis of Kindlin-1 in dissociated MT-Kin-1^{wt/wt} and MT-Kin-1^{fl/fl} tumors prior to tail vein injection. (F) Representative FACS histograms showing active β 1 and total integrin in MT-Kin-1^{wt/wt} and MT-Kin-1^{fl/fl} tumors.