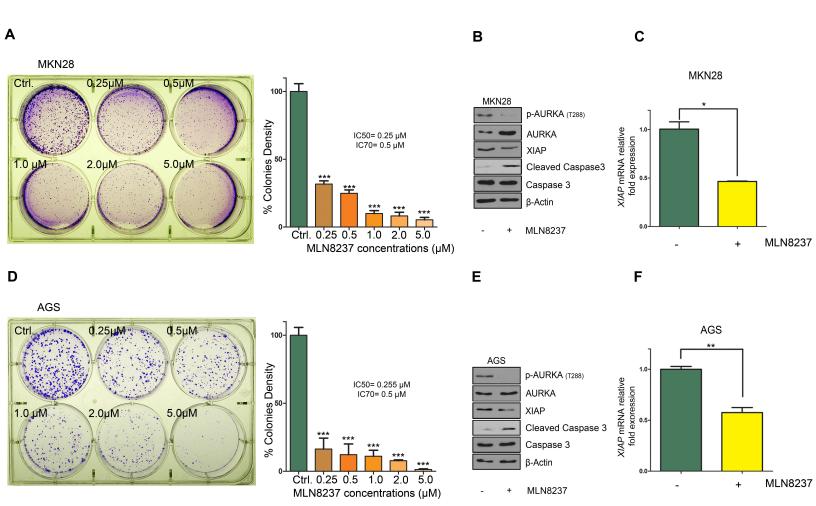
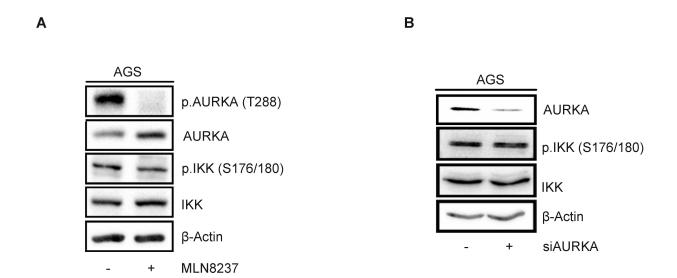


NF- $\kappa$ B reporter assay was performed in AGS (A) and MKN28 (B) cells with TNF- $\alpha$  treatment alone and with MLN8237 or Bay for 6h. The results showed that AURKA inhibition prevented TNF- $\alpha$  activation oof NF- $\kappa$ B.

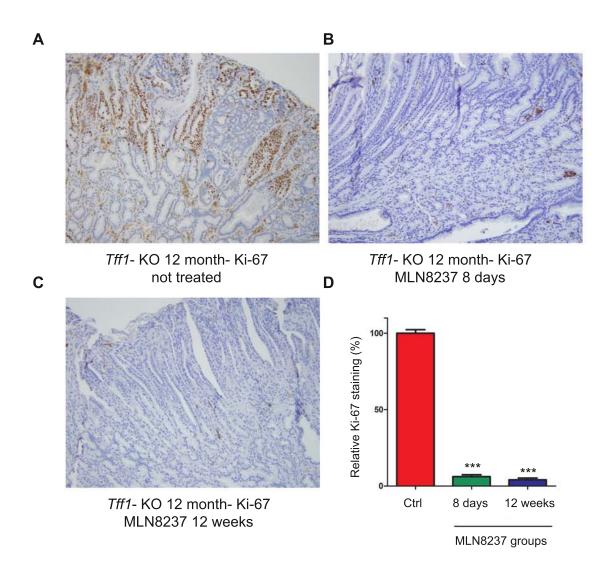
## Supplementary Figure 2: Inhibition of AURKA with MLN8237 reduces survival of gastric cancer cells



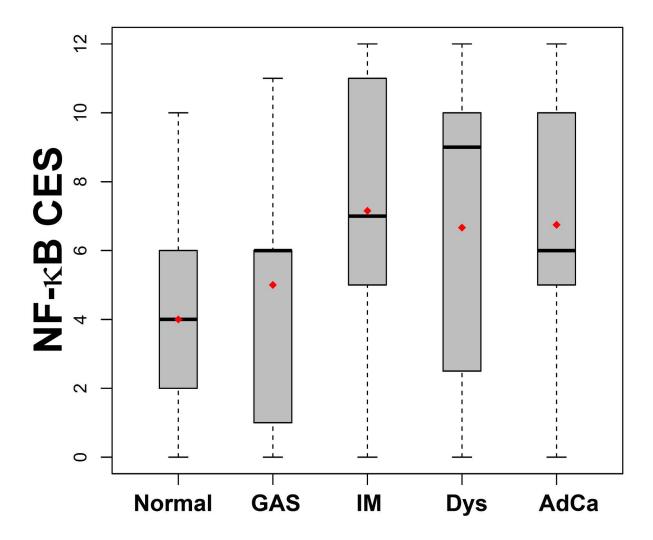
MKN28 (A) and AGS (D) cells were seeded at low density and treated with vehicle or the indicated concentrations of MLN8237 for 24h. Cells were then cultured in complete medium for 10 days.Graphs show the colonies density relative to vehicle-treated controls (right panels). MKN28 (B) and AGS (D) cells were treated with MLN8237 for 48h and proteins were collected and analyzed by Western blotting for XIAP, p-AURKA, AURKA, caspase-3, and cleaved caspase-3. Real-time RT-PCR analysis of *XIAP* in MKN28 (C) and AGS (F) after MLN8237 (0.5  $\mu$ M) treatment for 48h was performed.



A) AGS cells were treated with MLN8237 (0.5  $\mu$ M) and subjected to Westren blotting analysis for the indicated proteins.B) Western blot analysis of the indicated proteins in AGS cells after knocking down AURKA with siRNA.



A) Representative images of immunohistochemical analysis of Ki-67 protein expression in control (A) or MLN8237-treated (B&C) *Tff1*-knockout animals. D) Quantification of Ki-67 expression.



The graph summarizes the NF-kB immunohistochemical staining results on gastric tissues that are shown in Figure 7D. Horizontal bars indicate the median whereas red dots depict the mean. Normal, normal gastric glands; GAS, gastritis; IM, intestinal metaplasia; Dys, dysplasia; AdCa, adenocarcinoma. NF- $\kappa$ B was significantly overexpressed in all stages of gastric tumorigenesis (P<.01).