



Fig. S6. Otub1 deficiency promotes NF- κ B activation and B cell hyperplasia independently of p100 C-terminal phosphorylation. **a,b** Flow cytometric analyses of B220⁺ B cells and TCR β ⁺ T cells in the spleen of *Nfkb2*^{lym1/+} (Lym1/+) or *Otub1*-BKO/*Nfkb2*^{lym1/+} double mutant mice (8-10 week old). Data are presented as a representative plot (**a**) and summary graphs of mean \pm SD values based on multiple mice (**b**, each circle represents a mouse). **c,d** Flow cytometric analyses of immature (Imm, B220⁺CD93⁺) and mature (Mature, B220⁺CD93⁻) B cells as well as follicular (FO, B220⁺CD21^{int}CD23⁺) and marginal zone (MZ, B220⁺CD21^{hi}CD23⁻) B cells in the spleen of *Nfkb2*^{lym1/+} (Lym1/+) or *Otub1*-BKO/*Nfkb2*^{lym1/+} double mutant mice (8-10 week old). Data are presented as a representative plot (**c**) and summary graphs (**d**). **e,f** Immunoblot analysis of the indicated proteins using cytoplasmic or nuclear extracts of BAFF-stimulated (**e**) or anti-IgM-stimulated (**f**) splenic B cells from wildtype (WT), *Otub1*-BKO, *Nfkb2*^{lym1/+} (Lym1/+) or *Otub1*-BKO/*Nfkb2*^{lym1/+} double mutant mice (8-10 week old). Data are representative of 3 independent experiments. P values are determined by two-tailed unpaired t-test. *P<0.05, **P<0.01.