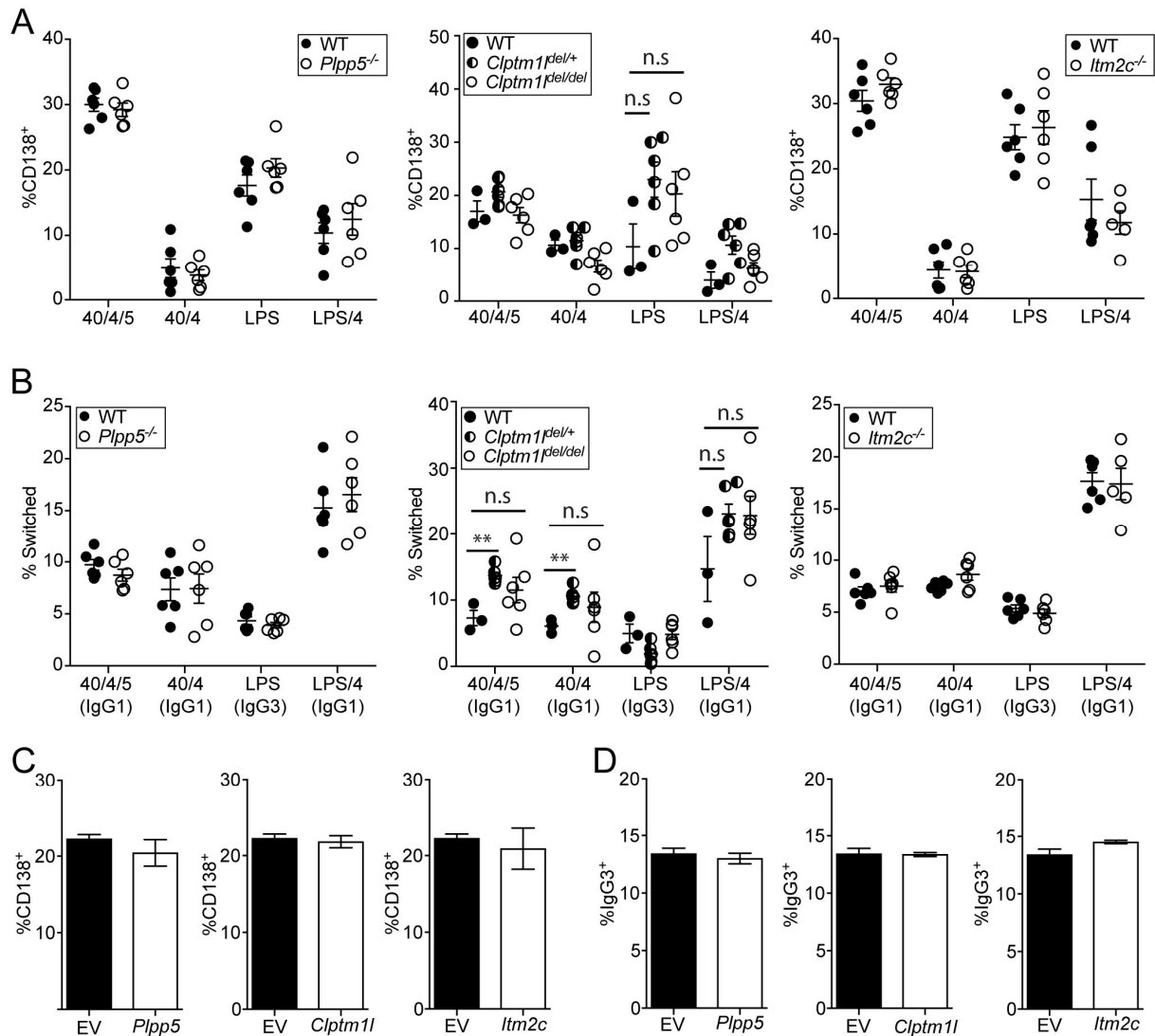


Supplementary Figure 1. *Plpp5*^{-/-}, *Clptm1*^{del/del} and *Itm2c*^{-/-} mice have normal B cell development and maturation. (A) Frequency and; (B) Number of precursor (B220⁺IgM⁻), immature (B220^{lo}IgM⁺) and recirculating (B220^{hi}IgM⁺) B cells in the bone marrow of *Plpp5*^{-/-}, *Clptm1*^{del/del}, *Itm2c*^{-/-} and age-matched WT mice. (C) Frequency and; (D) Number of mature (IgD^{hi}IgM⁺) and immature (IgD^{lo}IgM^{hi}) B cells in the spleen of *Plpp5*^{-/-}, *Clptm1*^{del/del}, *Itm2c*^{-/-} and age-matched WT mice. Results are combined from 2 (*Clptm1*) or 3 (*Plpp5*, *Itm2c*) independent experiments. Horizontal line shows the mean ± SEM. Each dot represents an individual mouse.



Supplementary Figure 2. *Plpp5*^{-/-}, *Clptm1*^{del/del} and *Itm2c*^{-/-} B cells respond normally to *in vitro* stimulation. **(A-B)** Naïve splenic B cells were isolated from *Plpp5*^{-/-}, *Clptm1*^{del/+}, *Clptm1*^{del/del}, *Itm2c*^{-/-} and age-matched WT mice and cultured for 4 days in CD40L, IL-4, IL-5 (40/4/5), CD40L, IL-4 (40/4), LPS or LPS/IL-4 (LPS/4) before analysis by flow cytometry. The proportion of B cells that had undergone **(A)** differentiation to ASCs (CD138⁺) or **(B)** Immunoglobulin class-switch recombination is shown. Each dot represents an individual mouse. **(C-D)** Naïve splenic B cells were isolated and stimulated for 24 hours with LPS before retroviral transduction with full length *Plpp5*, *Clptm1l*, *Itm2c* or empty vector (EV) control. After a further 3 days of LPS stimulation, cells were analyzed by flow cytometry for the proportion of transduced (GFP⁺) cells that had undergone **(C)** ASC differentiation (CD138⁺) and **(D)** Immunoglobulin class-switch recombination. Results are the mean \pm SEM from 2 independent experiments. Statistical significance was analyzed using unpaired t-test, correcting for multiple comparisons. ** P<0.01 for the indicated comparison. n.s, not significant (P>0.05).