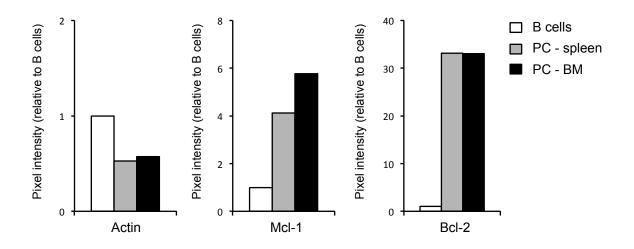
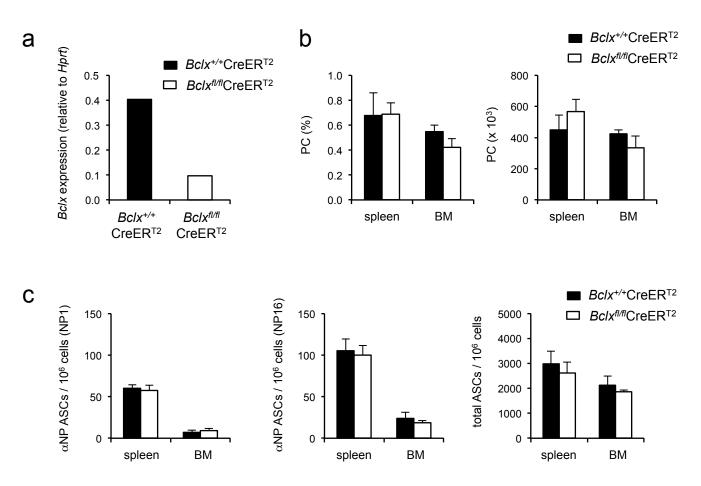


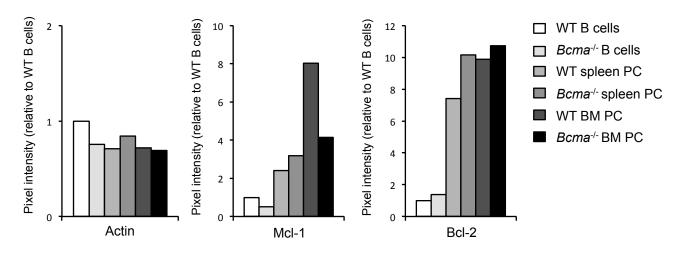
Supplementary Figure 1 | Expression of pro-survival Bcl-2 family members in Blimp-1-intermediate PBs and Blimp-1-high PCs. Gene transcription of prosurvival Bcl-2 family members in FACS sorted PBs (CD138+GFP<sup>int</sup>) or PCs (CD138+GFP<sup>hi</sup>) from the spleen or bone marrow (BM) and GC B cells (CD19+PNA+) from  $Prdm1^{GFP/+}$  mice, corrected for expression of the household gene Hprt and relative to expression in BM PCs (CD138+GFP<sup>hi</sup>, arbitrarily set to 1). Values ± SEM are indicated when relative expression is lower than 0.1. The average from 4 individual mice is shown; error bars, SEM.



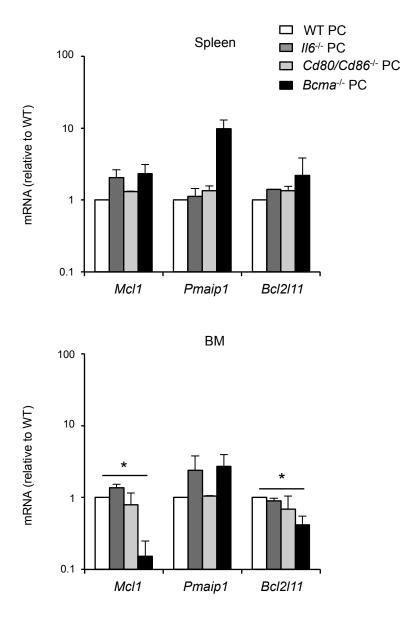
Supplementary Figure 2 | Quantification of Mcl-1 and Bcl-2 protein expression in PCs. Pixel intensity of the immunoblots shown in Fig. 1a measured using Photoshop software and relative to the pixel intensity as measured in B cells.



**Supplementary Figure 3 | Survival of existing PCs is not perturbed after deletion of** *Bclx*. (a) Gene transcription of *Bclx* in flow cytometry sorted PCs (B220<sup>-</sup>CD138<sup>+</sup>) from the bone marrow (BM) of *Bclx<sup>+/+</sup>*CreER<sup>T2</sup> or *Bclx<sup>fl/fl</sup>*CreER<sup>T2</sup> mice 2 days after treatment with Tamoxifen. Data are presented relative to expression of the household gene *Hprt* and the average of 4 mice pooled per group. (b) Percentage and absolute number of total PCs (B220<sup>-</sup>CD138<sup>+</sup>) in spleens and BM of *Bclx*<sup>+/+</sup>CreER<sup>T2</sup> or *Bclx<sup>fl/fl</sup>*CreER<sup>T2</sup> mice at 18 days after immunization with NP-KLH and 4 days after treatment with Tamoxifen. (c) ELISpot analysis of total (NP16), or high affinity (NP1), NP-specific or total antibody secreting cells (ASC) 4 days after deletion of *Bclx* and 18 days after immunization with NP-KLH as in (b). Data represent the average of 4 animals per group; error bars, SEM. The results are representative of 2 independent experiments.

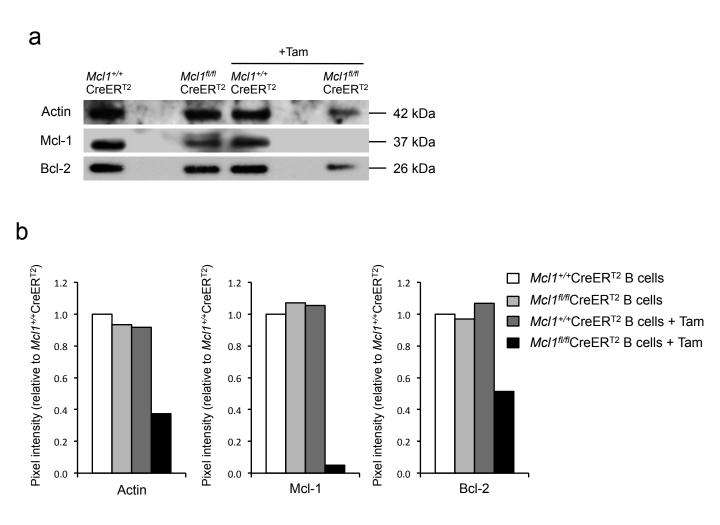


**Supplementary Figure 4 | Quantification of Mcl-1 and Bcl-2 protein expression in** *Bcma<sup>-/-</sup>* **PCs.** Pixel intensity of the immunoblots shown in **Fig. 2d** measured using Photoshop software and relative to the pixel intensity as measured in WT B cells.



Supplementary Figure 5 | Bim expression is reduced in *Bcma*-deficient BM PCs. Gene transcription of *Mcl1* and

pro-apoptotic Bcl-2 family members *Noxa* (*Pmaip1*) and *Bim* (*Bcl2111*) in PCs (B220<sup>-</sup>CD138<sup>+</sup>) isolated from the spleen or BM of wild type (WT), *Il6<sup>-/-</sup>*, *Cd80<sup>-/-</sup>Cd86<sup>-/-</sup>* and *Bcma<sup>-/-</sup>* mice. Values are presented as the expression relative to that found in WT PCs (arbitrarily set to 1) and represent the average of 2 experiments with PCs from 4 mice pooled per group; error bars, SEM. \*p≤0.05



Supplementary Figure 6 | Mcl-1 protein expression after Tamoxifen-mediated *Mcl1* deletion. (a) Immunoblot analysis for Mcl-1 and Bcl-2 of B cells purified from  $Mcl1^{fl/+}$ CreER<sup>T2</sup> or  $Mcl1^{fl/+}$ CreER<sup>T2</sup> mice and cultured for 4 days in the presence or absence of Tamoxifen added on day 3. (b) Pixel intensity of the immunoblots shown in (a), measured using Photoshop software and relative to the pixel intensity as measured in  $Mcl1^{+/+}$ CreER<sup>T2</sup> B cells.