



Published in final edited form as:

Nat Immunol. 2014 April ; 15(4): 354–364. doi:10.1038/ni.2830.

Innate lymphoid cells integrate stromal and immune signals to enhance antibody production by splenic marginal zone B cells

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Abstract

Innate lymphoid cells (ILCs) regulate stromal, epithelial and immune cells, but their impact on B cells remains unclear. We identified ROR γ ⁺ ILCs nearby the marginal zone (MZ), a splenic compartment containing innate-like B cells that respond to circulating T cell-independent (TI) antigens. Splenic ILCs established a bidirectional crosstalk with MAdCAM-1⁺ marginal reticular cells by providing tumor necrosis factor (TNF) and lymphotoxin, and activated MZ B cells via BAFF, CD40 ligand and the Notch ligand, Delta-like 1. Splenic ILCs further helped MZ B cells and their plasma cell progeny by co-opting neutrophils through the release of GM-CSF.

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AUTHOR CONTRIBUTIONS

G.M. and M. Miyajima designed and performed research, discussed data and wrote the paper; S.B., A.M., I.P. and A. Chudnovskiy designed and performed research; C.M.B., L. Cassis, L. Comerma, M.G., M.C. and D.L. performed research; S.S., J.I.A., M.J. and J.Y. provided blood and tissue samples and discussed data; S.F. and M. Merad designed research, provided reagents and discussed data; A.C. designed research, discussed data and wrote the paper.

COMPETING INTEREST STATEMENT

The authors declare that they have no competing financial interests.

Consequently, ILC depletion impaired both pre- and post-immune TI antibody responses. Thus, ILCs integrate stromal and myeloid signals to orchestrate innate-like antibody production at the interface between the immune and circulatory systems.

INTRODUCTION

The spleen is a highly perfused organ specialized in host defense against blood-borne pathogens. Interposed between the follicles of the splenic white pulp and the circulation, the marginal zone (MZ) contains B cells enmeshed with macrophages and dendritic cells (DCs) in a stromal reticular cell network¹⁻³. All of these cells provide an efficient immunosurveillance of the circulatory system by readily interacting with circulating antigens from commensal or pathogenic microbes owing to the slow flow rate of the blood passing through the MZ⁴. Following antigen capture, macrophages, DCs and possibly neutrophils of the innate immune system expose antigen to MZ B cells, a unique subset of antibody-producing lymphocytes that develop from transitional B cells in response to NOTCH2 signals⁵.

Lymphoid sites positioned between the host and the environment contain innate-like B and T cells that belong to the adaptive immune system, but share several properties with effector cells of the innate immune system. Mucosal and serosal membranes include innate-like B-1 cells that generate a first line of protection through early production of low-affinity immunoglobulin M (IgM) to bacteria⁶. When microbes breach the mucosal barrier and enter the general circulation, innate-like MZ B cells provide a second line of protection via low-affinity IgM and IgG that bridge the temporal gap required for the slower production of high-affinity IgG by follicular (FO) B cells⁴.

Similar to B-1 cells, MZ B cells express clonally distributed and somatically recombined but rather unspecific B cell receptor (BCR) molecules encoded by poorly diversified immunoglobulin (Ig) genes^{4, 6}. MZ B cells also express non-clonally distributed and germline-encoded Toll-like receptors (TLRs)⁷, a subfamily of nonspecific microbial sensors generally known as pattern recognition receptors. Typically expressed by effector cells of the innate immune system, TLRs activate MZ B cells after recognizing conserved microbial molecular signatures in cooperation with BCRs⁸. The activation of MZ B cells is further enhanced by B cell-stimulating cytokines released by DCs, macrophages and neutrophils^{9, 10}.

Besides innate-like lymphocytes, mucosal surfaces include innate lymphoid cells (ILCs) that express neither somatically recombined antigen receptors nor conventional surface lineage molecules¹¹. These ILCs require the transcriptional repressor inhibitor of DNA 2 (Id2) and the cytokine interleukin-7 (IL-7) for their development and generate cytokine secretion patterns that mirror those of T helper (T_H) cells of the adaptive immune system^{12, 13}. Similar to pro-inflammatory T_H1 cells, group 1 ILCs (ILC1) release interferon- γ (IFN- γ) and require the transcription factor T-bet for their development as do natural killer (NK) cells of the innate immune system¹⁴. ILC2, which include natural helper cells and nuocytes, secrete IL-5 and IL-13 and require the transcription factor GATA-3, thus resembling pro-inflammatory T_H2 cells¹⁵⁻¹⁷. Finally, ILC3 require the transcription factors retinoic acid

receptor-related orphan receptor- γ t (ROR γ t) and aryl hydrocarbon receptor (AhR) and include mucosal NK-22 cells, which secrete IL-22 and thus mimic non-inflammatory T_H22 cells¹⁸⁻²¹, as well as fetal and mucosal lymphoid tissue inducer (LTi) cells, which produce IL-22 and IL-17 and thus resemble pro-inflammatory T_H17 cells²²⁻²⁴. While NK-22 cells express natural cytotoxicity receptors (NCRs) usually associated with NK cells and mediate mucosal homeostasis by targeting epithelial cells via IL-22 (refs. 25-27), LTi cells lack NCRs and promote fetal lymphoid organogenesis and post-natal mucosal immunity by targeting stromal cells via lymphotoxin (LT) and tumor necrosis factor (TNF)²⁸⁻³⁰.

Mucosal NK-22 cells, also defined as NCR⁺ ILC3 to distinguish them from inflammatory NCR⁻ ILC3 characterized by constitutive IL-17, IL-22 and activation-induced IFN- γ production^{31, 32}, express B cell-activating factor of the TNF family (BAFF)²⁰, a cytokine used by DCs, macrophages and neutrophils to help MZ B cells and plasma cells in a T cell-independent (TI) manner^{1, 9, 10}. BAFF and its homologue a proliferation-inducing ligand (APRIL) are related to CD40 ligand (CD40L), a TNF family member used by T follicular helper (T_{FH}) cells to activate FO B cells³³. Given their involvement in mucosal TI antibody production^{29, 34}, ILCs could regulate humoral immunity also in the MZ, a lymphoid area that is continually exposed to antigen as are mucosal membranes.

Here we identified ILCs with mucosa-like properties in the MZ and perifollicular zone of the spleen. These ILCs required survival signals from marginal reticular cells (MRCs), a MZ subset of stromal cells that responded to TNF and LT from ILCs. In addition to stimulating MZ B cells and plasma cells via BAFF, APRIL, CD40L and the NOTCH2 ligand Delta-like 1 (DLL1), splenic ILCs co-opted MZ B cell-helper neutrophils via granulocyte monocyte-colony stimulating factor (GM-CSF). Consequently, ILC depletion hampered MZ B cell production of antibodies to TI antigens.

RESULTS

Splenic ILCs have type-3 mucosal features

The MZ is continually exposed to blood-borne antigens and may thus require homeostatic signals from ILCs as mucosal membranes do⁴. Flow cytometry showed that histologically normal adult human spleens contained cells that lacked common lymphoid and myeloid lineage (Lin)-associated molecules, but expressed the IL-7 receptor CD127 and the stem cell growth factor receptor CD117 (**Fig. 1a** and **Supplementary Fig. 1a**). These ILCs resembled mucosal NCR⁺ ILC3s^{19, 20}, because they expressed the NK cell molecules CD56, CD96, CD161, NKp44 and NKp46, the CCL20 chemokine receptor CCR6, the activation molecules CD25 and CD69, and NOTCH2 (**Fig. 1a** and **Supplementary Figs. 1b**). Splenic ILCs expressed neither the CD8 co-receptor, which is associated with cytotoxic T cells, nor the integrin CD103, which is expressed by some mucosal lymphocytes, nor the CD4 co-receptor and the prostaglandin D₂ receptor CRTH2 (**Fig. 1a** and **Supplementary Fig. 1b**), which are expressed by mouse LTi cells and human ILC2, respectively^{16, 22}. Albeit expressing as much CD161 as NK cells, splenic ILCs showed more NKp44, CCR6 and CD96 but less NKp46 than NK cells did (**Supplementary Fig. 1c**).

Gene expression studies performed through quantitative reverse transcription-polymerase chain reaction (qRT-PCR) showed that splenic ILCs expressed the transcription factors ROR γ t, AhR and Id2, the cytokines IL-22, IL-26, LT- α , LT- β and TNF and the IL-23 receptor, whereas splenic NK cells, T cells and macrophages did not, except B cells that expressed abundant LT- β (**Fig. 1b** and **Supplementary Fig. 1d**). Compared to mucosal ILCs from tonsils, splenic ILCs expressed comparable ROR γ t and AhR, but less IL-22 and more LT- α , LT- β and TNF (**Fig. 1b**). These latter cytokines are also hallmarks of LTi cells together with IL-17 (refs. 23, 30). However, splenic ILCs lacked IL-17, which was instead detected in T cells (**Supplementary Fig. 1d**). Of note, splenic ILCs included CD56⁺ and CD56⁻ subsets with an overlapping ILC3 gene expression profile that did not include IFN- γ and Perforin-1, which were instead abundant in NK cells (**Fig. 1b** and **Supplementary Fig. 1e,f**). Flow cytometry showed that splenic ROR γ t⁺ ILCs lacked T-bet, a transcription factor that was instead expressed by ROR γ t⁻ NK cells (**Fig. 1c**).

Flow cytometry and enzyme-linked immunosorbent assay (ELISA) demonstrated that splenic ILCs were more abundant than mucosal ILCs (**Fig. 1d**) and displayed NCR⁺ ILC3 properties^{19, 20}. Indeed, splenic ILCs received anti-apoptotic signals from the stromal cytokine IL-7 and the innate cytokines IL-1 β and IL-23 and secreted IL-22 in response to IL-1 β and IL-23 as mucosal ILCs did (**Fig. 1e,f**). Finally, tissue immunofluorescence analysis (IFA) and immunohistochemistry (IHC) showed that ROR γ t⁺NKp44⁺CD117⁺tryptase⁻ ILCs occupied MZ and perifollicular zone (PFZ) areas and were distinct from CD117⁺tryptase⁺ mast cells and CD3⁺ROR γ t⁻NKp44⁻ T cells (**Fig. 1g,h** and **Supplementary Fig. 2a-d**). Some ILCs were also found in the periarteriolar lymphoid sheath (not shown). Thus, the human spleen contains mucosa-like ILCs different from NK cells and positioned nearby MZ B cells.

Splenic ILCs receive survival signals from MRCs

Given that LTi cells interact with stromal cells via LT and TNF^{22, 29, 30}, we sought to identify stromal cells in the human MZ by tissue IFA, IHC and flow cytometry. Fibroblast-like cells expressing mucosal addressing cell adhesion molecule-1 (MAdCAM-1) were detected in the MZ nearby IgD⁺ B cells and some ROR γ t⁺ ILCs (**Fig. 2a** and **Supplementary Fig. 3a**). These MAdCAM-1⁺ cells did not express the coagulation protein von Willebrand factor (vWF), which characterizes endothelial cells, whereas MAdCAM-1⁻ sinusoid-lining cells (SLCs) from the red pulp did (**Fig. 2a**). MAdCAM-1⁺ cells also lacked the endothelial molecules platelet endothelial cell adhesion molecule-1 (PECAM-1) and CD34 as well as the leukocyte molecule CD45, but expressed the stromal molecules Thy-1 (or CD90), CD141 (thrombomodulin), intercellular adhesion molecule-1 (ICAM-1), vascular cell adhesion molecule-1 (VCAM-1) and α -smooth muscle actin (**Fig. 2b** and **Supplementary Fig. 3b,c**), and were thus considered equivalent to mouse MRCs³.

Human MRCs expressed TLR3, which binds double-stranded RNA, TLR4, which binds lipopolysaccharide (LPS), and TLR9, which binds unmethylated CpG-rich DNA (**Fig. 2b**), pointing to a role of microbial products in the regulation of MRCs. Laser capture microdissection followed by qRT-PCR demonstrated that MRC-enriched MAdCAM-1⁺ tissue contained more ILC survival factors such as IL-1 β , IL-7 and IL-23 and more ILC-

recruiting factors such as CCL20 (a CCR6 ligand) than SLC-enriched mannose receptor (MR, an endocytic receptor)⁺ tissue or control MR⁻ tissue (**Fig. 2c**).

As shown by flow cytometry, splenic ILCs up-regulated the MAdCAM-1-binding integrin $\alpha_4\beta_7$ in response to IL-7 and IL-1 β , pointing to a role of MAdCAM-1- $\alpha_4\beta_7$ interaction in the perifollicular positioning of ILCs. Conversely, MRCs up-regulated ICAM-1 and VCAM-1 in response to splenic ILCs or a combination of LT and TNF or LPS (**Fig. 2d** and **Supplementary Fig. 3d,e**). Blocking experiments and qRT-PCR indicated that ILCs activated MRCs via LT and TNF, whereas MRCs up-regulated IL-7, CCL20 and MAdCAM-1 in response to LT and TNF (**Fig. 2e** and **Supplementary Fig. 3f,g**). Of note, transwell assays showed that MRCs supported ILC survival via contact-dependent and contact-independent signals that included IL-7 (**Fig. 2g,h**). Finally, splenic macrophages and DCs expressed more IL-1 β and IL-23 than MRCs (**Supplementary Fig. 3h**), suggesting that ILCs establish a bi-directional interplay with MRCs in splenic niches that may also entail innate immune cells.

Splenic ILCs and MRCs deliver B cell-helper signals

Given their MZ and perifollicular location, human splenic ILCs may regulate MZ B cells. As shown by tissue IFA and qRT-PCR, splenic ROR γ ⁺ ILCs were positioned nearby CD20⁺ B cells and, compared to other splenic leukocyte subsets or mucosal ILCs, expressed more BAFF, CD40L and DLL1 (**Fig. 3a,b**), three major B cell and plasma cell helper factors^{5, 9, 10, 35}. Unlike splenic macrophages, splenic ILCs expressed little or no APRIL (**Fig. 3b**), a BAFF-related plasma cell survival factor³⁶.

As shown by flow cytometry and ELISA, splenic ILCs expressed surface BAFF, CD40L and DLL1 and secreted BAFF, whereas APRIL was mainly secreted by macrophages (**Fig. 3c,d**). Accordingly, splenic ILCs stimulated the survival, proliferation, IgM secretion and plasmablast differentiation of IgD^{lo}CD27⁺ MZ but not IgD^{hi}CD27⁻ FO B cells (**Fig. 3e-g**). Having shown that MRCs provide survival and activation signals to ILCs, we verified whether MRCs augmented the B cell-helper function of ILCs. Compared to MZ B cells exposed to ILCs or MRCs alone, MZ B cells exposed to both ILCs and MRCs not only underwent more IgM secretion and plasmablast differentiation, but also acquired the ability to release IgG and IgA, particularly in the presence of CpG DNA (**Fig. 3f,g**). Thus, human splenic ILCs may integrate stromal and innate immune signals to orchestrate antibody production.

Splenic ILCs help MZ B cells via BAFF, CD40L and DLL1

Next, we used flow cytometry and ELISA to determine the mechanism by which human splenic ILCs help MZ B cells. BAFF-R-Ig, a soluble decoy receptor that blocks the binding of BAFF to B cells, attenuated the survival of MZ B cells exposed to ILCs (**Fig. 4a**). Of note, MZ B cells underwent plasmablast differentiation and IgM secretion in response to ILCs but not ILC-conditioned medium (**Fig. 4b,c**), suggesting the involvement of contact-dependent signals from membrane-bound BAFF, CD40L and/or DLL1. Accordingly, BAFF-R-Ig, CD40-Ig or a NOTCH signaling inhibitor mitigated ILC-induced IgM secretion, particularly when these inhibitors were used in combination (**Fig. 4d**).

Although required for the early development of MZ B cells, NOTCH2 remains expressed on mature MZ B cells and may thus modulate their function⁵. Indeed, when combined with BAFF, DLL1-expressing OP9 stromal cells stimulated plasma cell differentiation and IgM secretion through NOTCH signals in MZ B cells (**Fig. 4e-g**). In contrast, BAFF alone or combined with DLL1-negative OP9 cells had little or no stimulatory effect (**Fig. 4e-g**). Thus, besides eliciting MZ B cell survival and activation, human splenic ILCs promote plasmablast differentiation and survival via BAFF, CD40L and DLL1.

Splenic ILCs activate perfollicular neutrophils via GM-CSF

In human spleens, MZ B cells interact with neutrophils (termed N_{BH} cells) possibly originating from circulating precursors (termed N_C cells)¹⁰. Given the involvement of ROR γ t in GM-CSF expression by T_H17 cells and the important role of GM-CSF in neutrophil activation and survival³⁷, we wondered whether splenic ILCs expressed GM-CSF. Tissue IHC showed CD117⁺ ILCs nearby CD66 (CEACAM-1)⁺ N_{BH} cells positioned around the MZ (**Fig. 5a**). qRT-PCR, ELISA and flow cytometry demonstrated the expression of abundant GM-CSF in splenic ILCs, but not other leukocyte subsets (**Fig. 5b,c**). Splenic ILCs further expressed the neutrophil-recruiting chemokine IL-8 (or CXCL8), which was also produced by macrophages and NK cells, albeit in lesser amounts (**Fig. 5b,c**).

Given that GM-CSF enhances the B cell-helper function of neutrophils¹⁰, we used flow cytometry, ELISA, IFA and qRT-PCR to establish whether GM-CSF-producing splenic ILCs induce N_{BH} cell-like properties to N_C cells. In the presence of splenic ILCs, N_C cells not only acquired phenotypic traits typical of N_{BH} cells¹⁰, such as up-regulation of the activation molecules CD69, CD11b and CD24 and down-regulation of the adhesion molecule CD62 ligand (CD62L), but also received survival signals from GMCSF (**Fig. 5d,e**). When incubated with splenic ILCs or their conditioned medium, N_C cells up-regulated APRIL and thus stimulated more IgA production in MZ B cells (**Fig. 5f,g**). ILC-conditioned medium also augmented the capacity of N_{BH} cells to induce IgM, IgG and IgA in MZ B cells (**Fig. 5h**). Similar to GMCSF or LPS, ILC-conditioned medium stimulated N_C cells to form neutrophil extracellular traps (NETs) (**Fig. 5i**), which are antigen-trapping projections often emanating from N_{BH} cells¹⁰. Thus, human splenic ILCs may amplify MZ B cell responses by stimulating perfollicular neutrophils via GM-CSF.

Splenic ILCs enhance TI antibody production

To elucidate the *in vivo* contribution of splenic ILCs to humoral immunity, we initially took advantage of *Rorc*^{-/-} mice, which lack the ROR γ t transcription factor required for the development of ILC3 (refs. 22, 28). *Rorc*^{-/-} mice also lack T_H17 cells, a subset of CD4⁺ T cells that regulate neutrophils via IL-17 and GM-CSF³⁷⁻³⁹. Flow cytometry showed that *Rorc*^{-/-} mice had fewer splenic Lin⁻CD117⁺CD127⁺ ILCs than *Rorc*^{+/+} mice (**Fig. 6a**). In these wild type mice, flow cytometry and qRT-PCR demonstrated that splenic ILCs included both CD4⁺ and CD4⁻ subsets that expressed more ROR γ t, IL-22, LT- α and TNF than did other splenic leukocytes (**Fig. 6b** and **Supplementary Fig. 4a**). In contrast to human splenic ILCs, mouse splenic ILCs lacked BAFF and CD40L, but expressed APRIL and DLL1 (**Fig. 6b**).

We then verified whether *Rorc*^{-/-} mice had an altered MZ. As shown by tissue IFA, *Rorc*^{-/-} mice had no perturbations of either MOMA1⁺ metallophilic macrophages, ER-TR9⁺ MZ macrophages and ICAM-1⁺ marginal sinus cells from the MZ or IgM⁺ B cells from the MZ and white pulp or F4/80⁺ macrophages and VCAM-1⁺ endothelial cells from the red pulp (**Supplementary Fig. 4b**). Flow cytometry, tissue IFA and ELISA demonstrated that spleens from non-immunized *Rorc*^{-/-} mice contained conserved B cells, but fewer plasmablasts and plasma cells expressing IgG3 (**Fig. 6c,d** and **Supplementary Fig. 5a**), an antibody produced by MZ B cells through a TI pathway under pre-immune conditions or after immunization^{40, 41}. Besides less total serum IgG3, *Rorc*^{-/-} mice had less serum IgG3 to the bacterial TI antigen phosphorylcholine (PC) (**Fig. 6e,f**). To ascertain the relative contribution of ILCs and T cells to pre-immune IgG3 responses, *Rorc*^{+/-} and *Rorc*^{-/-} mice were crossed with T cell-deficient *Cd3e*^{-/-} mice. In agreement with the TI nature of pre-immune IgG3 production^{40, 41}, non-immunized *Rorc*^{+/-}*Cd3e*^{-/-} mice had normal pools of splenic B cells, plasma cells and plasmablasts expressing IgG3 (**Fig. 6g**). Compared to these mice, *Rorc*^{-/-}*Cd3e*^{-/-} mice showed conserved B cells, but fewer plasmablasts and plasma cells expressing IgG3 and less pre-immune serum IgG3 (**Fig. 6g,h**).

Unlike pre-immune serum IgG3, pre-immune serum IgM was augmented in *Rorc*^{-/-} mice together with splenic IgM^{hi}B220^{int/lo}CD5^{int} B-1 cells but not B220⁺CD21⁺CD23⁺ FO B cells and B220⁺CD21^{hi}CD23⁻ MZ B cells (**Supplementary Fig. 5b,c**). The expansion of IgM-producing B-1 cells in *Rorc*^{-/-} mice may result from enhanced migration of B-1 cells from the peritoneal cavity to the spleen due to the lack of gut-associated lymphoid tissue and increased systemic translocation of gut bacteria^{22, 28, 34, 42}. Nonetheless, *Rorc*^{-/-} mice had less pre-immune serum IgM to PC (**Supplementary Fig. 5d**), suggesting a role of ILCs in at least some TI IgM responses.

To dissect the MZ B cell-helper function of ILCs in a more physiological model, an anti-Thy-1.2 antibody was used to deplete ILCs in Thy-1-disparate *Rag1*^{-/-} chimeric mice generated as described in a recently published study⁴³. Compared to mice treated with a control antibody, mice treated with anti-Thy-1.2 showed fewer splenic ILCs, fewer IgG3-expressing plasmablasts and plasma cells, and reduced pre-immune serum IgG3 (**Fig. 7a-c** and **Supplementary Fig. 5e**). Furthermore, anti-Thy-1.2-treated mice had impaired post-immune IgG3 responses to 2,4,6-trinitrophenyl (TNP)-Ficoll (**Fig. 7d**), a TI antigen that induces antibody production in innate-like B cells, including MZ B cells⁴⁰.

The involvement of splenic ILCs in TI antibody production was also demonstrated by reconstituting lethally irradiated *Rorc*^{-/-} mice with a mix of bone marrow cells from *Rag1*^{-/-} mice – which generate ILCs but not T and B cells – and *Rorc*^{-/-} mice – which generate T and B cells but not ILCs. ILC-sufficient (ILC⁺) chimeric mice developed splenic ILCs, whereas control ILC⁻ mice generated by reconstituting irradiated *Rorc*^{-/-} mice with bone marrow cells from *Rorc*^{-/-} mice did not (**Fig. 7e**). Compared to ILC⁺ mice, ILC⁻ mice showed fewer splenic plasmablasts and plasma cells expressing IgG3 and less pre-immune serum IgG3 (**Fig. 7f,g**). In all mouse models, IgM responses were only marginally affected (not shown), indicating that splenic ILCs predominantly help IgG3-expressing plasmablasts and plasma cells emerging from MZ B cell responses to TI antigens.

Splenic ILCs regulate neutrophil homeostasis

We next determined the role of mouse splenic ILCs in the homeostasis of N_{BH} cells. Tissue IFA and flow cytometry showed that ILC⁻ bone marrow chimeric mice had fewer splenic Ly6G⁺CD11b⁺ neutrophils and extrafollicular IgG3-producing plasmacytes than control ILC⁺ mice (**Fig. 8a,b**). Splenic neutrophils were also decreased in *Rorc*^{-/-} mice with either a T cell-sufficient or T cell-deficient background (**Supplementary Fig. 6a-c**), indicating that splenic ILCs provide helper signals to neutrophils independently of T cells. To ascertain the contribution of splenic neutrophils to pre-immune IgG3 production, ILC-reconstituted (ILC⁺) bone marrow chimeric mice were depleted of splenic neutrophils with an anti-Ly6G antibody (**Fig. 8c**). These mice had normal splenic ILCs and IgG3-expressing B cells and plasmablasts, but fewer splenic IgG3-expressing plasma cells and decreased pre-immune serum IgG3 compared to ILC⁺ mice treated with a control antibody (**Fig. 8d-f**). Thus, unlike ILCs, neutrophils may predominantly control terminally differentiated IgG3-secreting cells.

Finally, we determined whether splenic ILCs regulate neutrophils via GM-CSF. As shown by qRT-PCR, splenic ILCs from wild-type mice expressed more GM-CSF than other leukocyte subsets did, including macrophages and NK cells (**Fig. 8g**). Compared to controls, GM-CSF (also known as colony-stimulating factor 2)-deficient *Csf2*^{-/-} mice had reduced pre-immune serum antibodies to bacterial PC and fewer splenic neutrophils, which increased after inoculation of GM-CSF-expressing B16 melanoma cells (**Fig. 8h** and **Supplementary Fig. 6d**). Also *Rag1*^{-/-}*Il2rg*^{-/-} mice, which lack the IL-2 receptor common γ chain mandatory for IL-7-dependent ILC development¹³, had fewer splenic neutrophils compared to *Rag1*^{-/-}*Il2rg*^{+/+} controls (**Fig. 8i**). Moreover, adoptive transfer of ILCs from *Csf2*^{+/+} but not *Csf2*^{-/-} mice increased splenic neutrophils in *Rag1*^{-/-}*Il2rg*^{-/-} mice (**Fig. 8i**). Thus, in addition to helping MZ B cells and plasma cells through BAFF (or APRIL), CD40L and DLL1, splenic ILCs co-opt neutrophils via GM-CSF to enhance TI antibody production (**Supplementary Fig. 7a,b**).

DISCUSSION

We have shown that the MZ and PFZ of the spleen contained group-3 mucosa-like ILCs that released LT and TNF to establish a bi-directional crosstalk with stromal MRCs. In addition to stimulating MZ B cells and plasma cells through BAFF, APRIL, CD40L and DLL1, splenic ILCs co-opted neutrophils with MZ B cell- and plasma cell-helper functions through GM-CSF, thereby sustaining TI antibody production.

Mucosal and serosal membranes contain evolutionarily primitive lymphocyte subsets that include B-1 cells, $\gamma\delta$ T cells, invariant NKT cells and mucosa-associated invariant T cells⁴⁴. By sensing conserved microbial signatures through somatically recombined and germline-encoded receptors, these innate-like lymphocytes rapidly activate protective programs that crossover the conventional boundaries between the innate and adaptive immune systems⁴⁴. Mucosal membranes also contain ILCs that lack somatically recombined receptors and yet mount prompt T cell-like responses¹³. Here, we found that splenic ILCs enhanced antibody production to TI antigens by activating innate-like B cells positioned at the interface between the immune and circulatory systems.

Similar to mucosal ILC3 (refs. 19, 20), human splenic ILCs contained both ROR γ t and AhR transcription factors, expressed NCRs along with various activation molecules, and secreted IL-22 in response to IL-23. In addition, human splenic ILCs activated MZ-based stromal MRCs via LT and TNF and thus also resembled fetal or mucosal LTi cells^{28-30, 45}. However, unlike LTi cells^{23, 30}, human splenic ILCs did not express IL-17, raising questions as to the ontogenetic relationship between these ILC subsets. Mouse splenic ILCs had a phenotype and gene expression profile similar to those of human splenic ILCs, but expressed CD4, thus showing a closer affiliation with LTi cells.

Human splenic ILCs not only expressed the gut homing receptor $\alpha_4\beta_7$, but also interacted with MRCs equipped with MAdCAM-1, an $\alpha_4\beta_7$ ligand usually associated with gut endothelial cells³⁴. Given their additional expression of CCL20 and the presence of the CCL20 receptor CCR6 on splenic ILCs, MRCs might be involved in the positioning of ILCs within and around the MZ. Consistent with their stroma phenotype and reticular morphology, MRCs lacked endothelial molecules and exhibited strong responsiveness to LT and TNF from ILCs, thus resembling MAdCAM-1⁺ stromal cells lining the mouse marginal sinus⁴⁶. Conversely, MRCs delivered both contact-dependent and contact-independent survival signals to ILCs, including IL-7. Remarkably, MRCs increased IL-7 expression in response to LT and TNF, pointing to the presence a bi-directional functional crosstalk between MRCs and ILCs in the MZ and PFZ of the spleen. Additional ILC survival factors such as IL-1 β and IL-23 were predominantly expressed by splenic DCs and macrophages, suggesting that ILCs inhabit a splenic niche that entails survival signals from both stromal and immune cells. Of note, MRCs received further activation signals from TLR ligands such as LPS, lending support to the possibility that commensal antigens from the intestinal mucosa help the post-natal development and function of the splenic MZ^{4, 10, 47, 48}.

In humans, splenic ILCs enhanced the survival of MZ B cells through a contact-independent mechanism involving soluble BAFF, but triggered the differentiation of MZ B cells into plasmablasts through contact-dependent signals provided by membrane-bound BAFF, DLL1 and CD40L. Of note, DLL1 synergized with soluble BAFF to induce IgM secretion, possibly owing to the pro-survival effect of soluble BAFF on plasmablasts³⁵. Given that DLL1 also cooperates with BAFF to stimulate the early development of MZ B cells⁵, there is the possibility that DLL1 on ILCs operates at both early and late stages of MZ B cell differentiation with distinct signaling programs. Of note, splenic ILCs further helped MZ B cells by co-opting N_{BH} cells through GM-CSF. This cytokine may also enhance the MZ B cell-stimulating function of DCs and possibly macrophages^{1, 4}, which may explain why a relative small number of ILCs is sufficient to enhance TI antibody responses.

Despite expressing CD40L, a class switch-inducing factor typically detected on T_{FH} cells³³, human splenic ILCs did not trigger IgG or IgA production in MZ B cells. Nonetheless, MZ B cells induced IgA and IgG when exposed to ILCs in the presence of MRCs and microbial TLR ligands. Accordingly, small amounts of these bacterial components can be detected in splenic perifollicular areas together with plasma cells releasing IgM, IgG or IgA¹⁰. Thus, similar to gut LTi cells²⁹, splenic ILCs may integrate stromal and innate immune signals to rapidly generate antibodies for antimicrobial protection.

Although lacking both CD40L and BAFF, mouse splenic ILCs expressed APRIL, a BAFF-related molecule that enhances plasma cell survival³⁶. By also expressing DLL1, mouse splenic ILCs would sustain the differentiation and/or survival of plasmablasts and plasma cells emerging during IgG3 responses to TI antigens. In mice, ILC depletion did not impair IgM production, suggesting that ILCs predominantly enhance MZ B cell-derived IgG3 rather than B-1 cell-derived IgM responses. Nonetheless, splenic ILCs sustained pre-immune production of IgM to bacterial PC, which points to the involvement of ILCs in IgM responses to some but not all TI antigens.

In ILC-deficient mice, impaired TI IgG3 production was associated with a reduction of N_{BH} cells, which help MZ B cells via BAFF and APRIL¹⁰. Accordingly, human splenic ILCs enhanced the survival and B cell-helper activity of N_{BH} cells through GM-CSF. This cytokine also induced NET-like structures, which might enable N_{BH} cells to capture blood-borne antigens transiting through the MZ^{4, 10}. Human splenic ILCs also released IL-8, which may contribute to the recruitment of N_{BH} cells. Mice do not express IL-8, but their splenic ILCs may attract N_{BH} cells via alternative factors, including GM-CSF. Remarkably also splenic innate response activator (IRA) B cells express GM-CSF⁴⁹, which could explain the persistence of some N_{BH} cells in mice lacking ILCs.

GM-CSF further derives from T_H17 cells³⁷, but the analysis of T cell-deficient mice suggested that ILCs can regulate the homeostasis of N_{BH} cells independently of T cells. In mice, splenic ILCs sustained both IgG3⁺ plasmablasts and IgG3⁺ plasma cells, whereas mouse N_{BH} cells predominantly helped IgG3⁺ plasma cells, suggesting that multiple subsets of innate immune cells orchestrate TI IgG3 responses in a hierarchical manner. Of note, IgG3 may provide protection against certain pathogens following its interaction with ficolins, a family of soluble pattern recognition receptors released by hepatocytes and various cells of the innate immune system⁴¹. In summary, our studies indicate that splenic ILCs orchestrate a stromal-innate cell network that fosters TI antibody production by MZ B cells. Harnessing this network with adjuvants may enhance protective humoral responses to poorly immunogenic TI antigens^{4, 50}, including viral glycoproteins and bacterial carbohydrates.

ONLINE METHODS

Human samples

Mononuclear cells and neutrophils were isolated from the peripheral blood of healthy volunteers or histologically normal spleens from deceased organ donors or splenectomized trauma patients without clinical signs of infection or inflammation¹⁰. Tonsils were from individuals with follicular hyperplasia¹⁰. The use of blood and tissues was approved by the Ethical Committee for clinical investigation of Institut Hospital del Mar d'Investigacions Mèdiques (CEIC-IMIM 2011/4494/I). Tissue samples were collected by the department of Pathology of Hospital del Mar. Prior to collection, signed informed consent was obtained from the patient or his/her parent or guardians. All the blood and tissue samples were coded and relevant clinical information remained anonymous.

Mice

Rorc^{-/-} (*Rorc*^{gfp/gfp}), *Cd3e*^{-/-}, *Rag1*^{-/-}, *Rag1*^{-/-}*Il2rg*^{-/-} and *Csf2*^{-/-} mice on the C57BL/6 background have been described previously^{29,51-54}. C57BL/6 CD90.1 mice were purchased from the Jackson Laboratory. All mice were housed in specific pathogen-free conditions. Male and female mice were used at age 8-12 weeks unless specified in the text. All the experiments involving mice were performed in accordance with approved protocols from the Institutional Animal Care at RIKEN and Icahn School of Medicine at Mount Sinai.

Mouse chimeras and neutrophil depletion

4×10^7 bone marrow cells from *Rag1*^{-/-} mice were mixed with 1×10^7 bone marrow cells from *Rorc*^{-/-} mice and injected intravenously into 10-Gy irradiated 2-month old *Rorc*^{-/-} mice. After transplantation, mice were given 500 mg/L ampicillin (Sigma) and 1 g/L neomycin (Sigma) in drinking water for 2 weeks. To deplete neutrophils, control RTK2758 monoclonal antibody (mAb) or neutrophil-depleting 1A8 mAb to Ly6G (BioXCell) were administered intraperitoneally every 5 days at a dose of 250 μ g/mouse for a month. Thy-1 disparate *Rag1*^{-/-} chimeras were generated and depleted of ILCs with a 30-H12 mAb to Thy-1.2 (BioXCell) injected intraperitoneally every 3 days at a dose of 500 μ g/mouse for a month as previously published⁴³. LTF-2 mAb was used as control.

Adoptive transfers

$2-3 \times 10^4$ ILCs from the small intestine of either *Csf2*^{+/+} or *Csf2*^{-/-} mice were adoptively transferred into *Rag1*^{-/-}*Il2rg*^{-/-} mice via retro orbital injection as previously described and splenic neutrophils were analyzed 21 days later. 1×10^5 B16^{Csf2} melanoma cells overexpressing GM-CSF (from G. Dranoff) were injected into *Csf2*^{-/-} mice and splenic neutrophils were analyzed 12-14 days later.

Immunization

Serum TNP-specific antibody titers were determined by ELISA 7 days after intraperitoneal injection of 50 μ g TNP-Ficoll.

Cells

Human splenocytes or tonsillar mononuclear cells were obtained from fresh tissue samples as reported in published studies¹⁰. Human ILCs, NK cells, naïve B cells, MZ B cells, macrophages, T cells, N_C cells and N_{BH} cells were sorted by flow cytometry as described previously¹⁰ or reported below. To isolate human MRCs, splenic tissues were enzymatically digested in Hank's balanced salt solution (Lonza) containing 1 mg/ml collagenase-IV (Invitrogen) and 50 ng/ml DNase-I (50 ng/ml) at 37 °C for 45 min, followed by separation on Ficoll-Hypaque gradient. Splenocytes were resuspended in EGM medium (Lonza) and cultured in 0.1% gelatin (Sigma) pre-coated plate. 72 h later, adherent cells were obtained with Accutase cell detachment solution as instructed by the manufacturer (Millipore). CD45⁻ CD31⁻ adherent cells were sorted by flow cytometry and expanded up to three passages in EGM medium. Murine OP9 and OP9-DLL1 stromal cell lines (from R. Gimeno) were tested for mycoplasma and grown in essential medium- α (Invitrogen) supplemented with 20% Fetalclone I (Hyclone).

Cultures and reagents

Human splenic ILCs (5×10^4 /well) were plated in 96 U-bottom plates and cultured in complete RPMI medium with or without 50 ng/mL IL-7, 50 ng/mL IL-1 β and/or 50 ng/mL IL-23 (Peprotech). Conditioned medium was obtained by culturing splenic ILCs in complete medium for 24 h. In co-culture experiments, splenic ILCs were first expanded with IL-7 and IL-1 β for 5-8 days as previously published. ILCs were co-cultured with confluent human MRCs in flat-bottom plates for 72 h. To avoid cell-to-cell contact, some co-cultures were performed in a 24-transwell system (Corning). Human ILCs were also co-cultured with human MZ B cells, N_C cells and/or N_{BH} cells at a 1:10 ratio. Some co-cultures were supplemented with 500 ng/ml BAFF (Alexis), 0.25 μ g/ml CpG ODN-2006 (InvivoGen), 30 μ g/ml BAFF-R-Ig, 5 μ g/ml CD40-Ig, 5 μ g/ml, 10 μ g/ml neutralizing MAB215 mAb to human GM-CSF, or 10 μ g/ml isotype control IgG1 mAb (R&D system). To inhibit NOTCH2, human MZ B cells were incubated with 25, 5 or 2.5 μ M N-[N-(3,5-difluorophenacetyl)-L-alanyl]-S-phenylglycine t-butyl ester (DAPT) (Sigma), which interferes with the cleavage of NOTCH proteins by γ -secretase. In some experiments, human MRCs were stimulated with 100 ng/ml LT $\alpha_1\beta_2$ and 10 ng/ml TNF (R&D Systems) or with ILCs in the presence or absence of 10 μ g/ml mAbs MAB2101 to TNF or MAB1370 to LT $\alpha_2\beta_1/\alpha_1\beta_2$ (R&D Systems). Human N_C cells were stimulated with 50 ng/ml GM-CSF (Peprotech) or 100 ng/ml LPS (InvivoGen).

Flow cytometry

Cells were incubated with Fc blocking reagent (Miltenyi Biotec) at 4 °C before adding appropriate cocktails of fluorochrome-labeled mAbs (**Supplementary Tables 1 and 2**). Dead cells were excluded with 4'-6-diamidino-2'-phenylindole (DAPI) as indicated by the manufacturer (Boehringer Mannheim). To stain intracellular transcription factors, human ILCs or NK cells were stained with mAbs to cell-specific surface molecules, permeabilized, fixed with a Transcription Factor Buffer Set as indicated by the manufacturer (eBioscience), and labeled with mAbs to human ROR γ t or T-bet (**Supplementary Table 1**). For intracellular cytokine staining, human ILCs were cultured for 4 h in complete RPMI medium with GolgiStop (BD Pharmingen) in the presence or absence of 10^{-7} M phorbol myristate acetate and 0.5 μ g/ml ionomycin (Sigma). These cells were stained with mAbs to specific surface molecules, fixed, permeabilized using the Cytotfix/Cytoperm kit (BD Pharmingen), and finally incubated with mAbs to human GM-CSF or IL-8 (**Supplementary Table 1**). To stain intracellular IgG3, mouse splenocytes were stained with mAbs to specific surface molecules, permeabilized, fixed with the Cytotfix/Cytoperm kit (BD Pharmingen), and labeled with R40-82 mAb to mouse IgG3 (**Supplementary Table 2**). Cells were acquired using FACS calibur, FACS Aria II, LSRII or LSRFortessa (BD Biosciences) and further analyzed using the FlowJo software (Tree Star).

FACSsorting

Human CD3⁻CD14⁻CD19⁻CD117⁺CD127⁺ ILCs, CD3⁻CD14⁻CD19⁻CD117⁻CD127⁻CD56⁺ NK cells, CD19⁺IgD^{hi}CD27⁻ naïve B cells, CD19⁺IgD^{lo}CD27⁺ MZ B cells, CD14⁺ macrophages and CD3⁺ T cells as well as mouse CD3⁺ T cells, B220⁺ B cells, CD11c⁺ DCs, CD11b⁺ macrophages, CD49b⁺ NK cells, Ly6G⁺ neutrophils,

CD4⁺CD45⁺CD117⁺CD127⁺Lin⁻ (CD3⁻B220⁻CD11b⁻CD11c⁻ Ly6G⁻) ILCs and CD4⁻CD45⁺CD117⁺CD127⁺Lin⁻ ILCs were stained with appropriate cocktails of fluorochrome-labeled mAbs (**Supplementary Tables 1 and 2**) and sorted using FACSAria II (BD Biosciences) after exclusion of dead cells through DAPI staining. The purity of FACSsorted cells was consistently > 95%.

Viability and proliferation assays

Cell survival was measured using the Annexin-V Apoptosis Detection Kit II (BD Pharmingen). Gates and quadrants were drawn to give 1% total positive cells in samples incubated with isotype control mAbs. Cell proliferation was assessed by CFSE staining using the CellTrace CFSE Cell Proliferation Kit (Invitrogen).

Laser capture microdissection

Microdissections were performed on fresh human splenic tissues frozen in optimal cutting temperature (OCT) compound using the ArcturusXT Laser Capture Microdissection System following the manufacturer's instructions (Life Technologies). In brief, 10- μ m tissue slices were isolated on membrane slides (Life Technologies) and fixed in cold acetone for 10 s. Immunohistochemistry with specific mAbs was followed by tissue dehydration and fixation with xylene. Groups of cells from completely dried tissues were acquired by microdissection and RNA was extracted using the Arcturus PicoPure RNA Extraction Kit (Life Technologies).

Immunofluorescence analysis

Frozen tissues and cells from humans or mice were fixed as previously published¹⁰ and stained with various combinations of Abs (**Supplementary Tables 1 and 2**). Biotinylated Abs were detected using horseradish peroxidase (HRP)-conjugated streptavidin followed by tetramethylrhodamine from a Tyramide Signal Amplification Kit (PerkinElmer Life Sciences) or streptavidin-Alexa Fluor conjugates. Nuclear DNA was stained with DAPI. Coverslips were applied with FluorSave reagent (Calbiochem). Images were obtained using Axioplan2 (Carl Zeiss) microscopy and further analyzed using Axiovision (Carl Zeiss) software. To perform NET formation assays, human circulating neutrophils were seeded on poly D-lysine coated glass cover slips and stimulated for 3 h with either human GM-CSF or human splenic ILC-conditioned medium. Cells were fixed with 4% paraformaldehyde and stained with mAbs NP57 or ab21595 to human elastase and DAPI.

Immunohistochemistry

5- μ m thick formalin-fixed and paraffin-embedded tissue sections were stained with various mAbs (**Supplementary Table 1**) using EnVisionTM + Dual Link System-HRP (DAB+) for single stainings and EnVision G/2 Doublestain System Rabbit/Mouse (DAB+/Permanent Red) for double stainings (Dako). Sections were counterstained with hematoxylin.

ELISA

Total human IgM, IgG and IgA were measured as previously published¹⁰. Soluble human BAFF ELISA Kit (Adipogen), LEGEND MAX Human APRIL/TNFSF13 ELISA Kit

(Biolegend), Human GMCSF ELISA Development Kit, and Human IL-8 ELISA Development Kit (Peprotech) were used to measure human BAFF, APRIL, GM-CSF and IL-8. Human Th1/Th2/Th9/Th17/Th22 13plex FlowCytomix Multiplex (eBioscience) was used to measure human IL-17 and IL-22. The concentration of mouse serum IgM, IgG3 and IgA were determined with Mouse ELISA Quantification Set (Bethyl Laboratories). To detect mouse TNP-specific IgG3, ELISA plates coated with 5 µg/ml bovine serum albumin (BSA)-conjugated TNP (BioSearch Technologies) were sequentially incubated with mouse serum and HRP-conjugated goat polyclonal antibody A90-111P to mouse IgG3 (Bethyl Laboratories). To detect mouse phosphorylcholine-specific IgM, ELISA plates coated with 5 µg/ml BSA-conjugated phosphorylcholine were sequentially incubated with serum and an HRP-conjugated goat polyclonal antibody A90-101P to IgM (Bethyl Laboratories). BSA-conjugated phosphorylcholine was synthesized as follows. 1 mg of BSA was coupled to 1 mg *p*-aminophenylphosphorylcholine in 0.1 M 2-(*N*-morpholino)ethanesulfonic buffer (Sigma) at pH 4.5 supplemented with 2 mg 1-ethyl-3-[3-dimethylaminopropyl]carbodiimide hydrochloride in a 1.6 ml final volume. This mixture was incubated for 2 h at 20 °C. Conjugated BSA was dialyzed against PBS at pH 7.4 and 4 °C.

qRT-PCR

To quantify human gene products, total RNA was extracted and reverse transcribed into cDNA as reported in previously published studies¹⁰. qRT-PCRs were performed as reported previously¹⁰ using specific primer pairs (**Supplementary Table 3**). To quantify mouse gene products, total RNA was extracted with TRIzol reagent (Gibco BRL). After DNase I (Invitrogen) treatment, random hexamers (Invitrogen) were used for first-strand cDNA synthesis. qRT-PCRs were performed in 96-well plates with a LightCycler 480 real-time PCR instrument (Roche Diagnostics) using the LightCycler 480 SYBR Green I Master kit (Roche Diagnostics) and specific primer pairs (**Supplementary Table 4**). Gene expression was normalized to that of glyceraldehyde 3-phosphate dehydrogenase in each sample.

Statistical analysis

Values were expressed as mean ± standard error of the mean (s.e.m.) or standard deviation (s.d.). Statistical significance was assessed with two-tailed or one-tailed unpaired Student's *t*-test unless specified otherwise. Mann-Whitney *U*-test or Wilcoxon matched-pairs signed rank test were performed on non-parametric data. Results were analyzed with Prism software (Graph Pad) and *P* values less than 0.05 were considered significant. In animal experiments, littermates (minimum 3 mice/group) were randomly distributed to the treatment groups so that all groups were age-matched and sex-matched. No specific randomization or blinding protocol was used and no animals were excluded from the analysis.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

ACKNOWLEDGMENTS

We thank A. Chorny, S. Casas (Icahn School of Medicine at Mount Sinai), Gregory F. Sonnenberg (University of Pennsylvania), G. Dranoff (Dana Farber Cancer Institute), Anna Mensa (Hospital Clínic of Barcelona), R. Gimeno, A. Bigas and M. López-Botet (Institut Hospital del Mar d'Investigacions Mèdiques) for providing reagents and samples and helping with discussions; E. Ramirez, E. Julià and O. Fornas (Barcelona Biomedical Research Park) for helping with cell sorting. Supported by European Advanced Grant ERC-2011-ADG-20110310, Ministerio de Ciencia e Innovación grant SAF2011-25241, and Marie Curie reintegration grant PIRG-08-GA-2010-276928 to A.C.; Juan de la Cierva post-doctoral fellowships to I.P. and G.M.; pre-doctoral fellowships from the Instituto de Salud Carlos III and Ministerio de Ciencia e Innovación to C.B, M.G and S.B.G.; an EUROPADnet HEALTH-F2-2008-201549 grant to A.C.; US National Institutes of Health grants R01 AI74378, R01 AI57653, U01 AI95613, U01 AI95776 IOF, P01 AI61093 and U19 096187 to A.C.; Grant-in-Aid for Scientific Research in Priority Areas to S.F.; and Grant-in-Aid for Scientific Research (B) to S.F.

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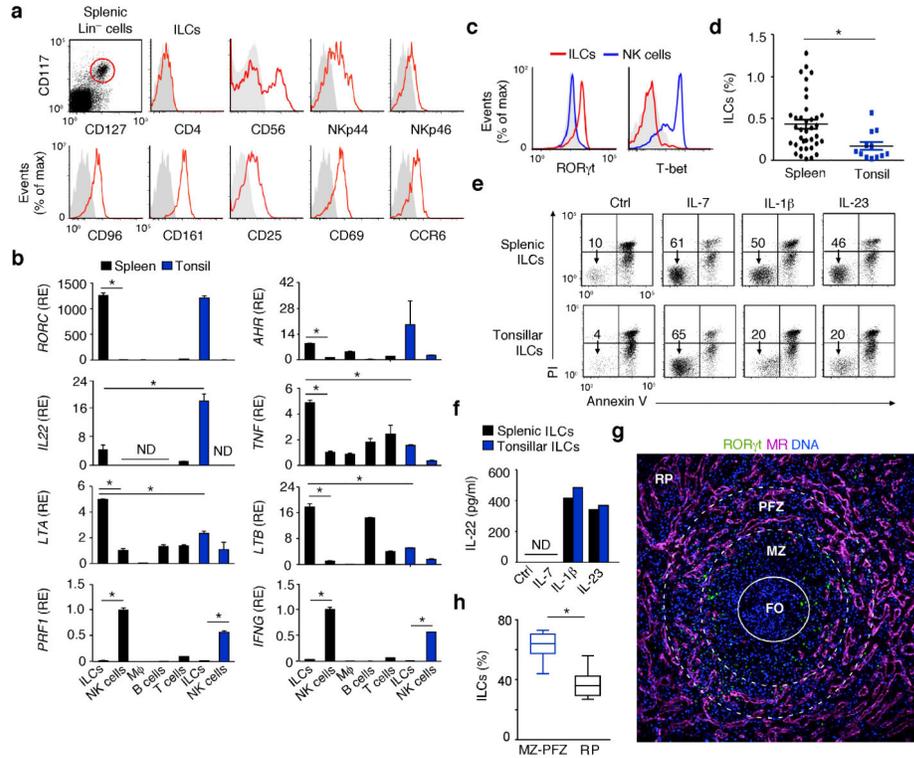


Figure 1. Human splenic ILCs express a mucosa-like group-3 phenotype and occupy MZ and perifollicular areas

(a) Flow cytometry of CD4, CD25, CD56, CD69, CD96, CD161, NKp44, NKp46 and CCR6 on splenic Lin⁻CD117⁺CD127⁺ ILCs (red gate). Gray shading, negative control. (b) qRT-PCR of *RORC* (*RORγt*), *AHR* (*AhR*), *IL22* (*IL-22*), *TNF* (*TNF*), *LTA* (*LT-α*), *LTB* (*LT-β*), *PRF1* (*Perforin-1*) and *IFNG* (*IFN-γ*) mRNAs from splenic or tonsillar ILCs and NK cells and from splenic macrophages (*Mφ*), B cells and T cells. Results are normalized to *ACTB* (*β-actin*) mRNA and presented as relative expression (RE) compared with that of fresh splenic NK cells. Error bars, s.e.m.; **P* < 0.05 (two-tailed unpaired Student's *t* test). (c) Flow cytometry of intracellular *RORγt* and T-bet in splenic ILCs (red lines) and NK cells (blue lines). (d) Flow cytometry of Lin⁻CD117⁺CD127⁺ ILCs in splenic and tonsillar lymphocytes. **P* < 0.05 (Mann-Whitney U-test). (e) Flow cytometric analysis of frequency of viable Annexin-V⁻propidium iodide (PI)⁻ ILCs after culture of splenic and tonsillar ILCs with medium alone (Ctrl), IL-1β, IL-7 or IL-23 for 72 h. (f) ELISA of IL-22 from splenic and tonsillar ILCs cultured as in (e). (g) IFA of spleen stained for MR (purple), *RORγt* (green) and DNA-binding 4'-6-diamidino-2'-phenylindole (DAPI; blue). RP, red pulp. Original magnification, ×10. FO: center of the follicle (h) Immunohistochemical quantification of CD117⁺Tryptase⁻ ILCs from MZ-PFZ and red pulp (RP) areas in nine microscopic ×20 fields from two spleens. **P* < 0.05 (Wilcoxon Matched-pairs signed Rank test). Data summarize three measurements from three pooled experiments with 1 donor in each (spleen and tonsil) (b), display values from thirty seven spleens and twelve tonsils (d), or show one of four experiments with similar results (a,c,e,f,g).

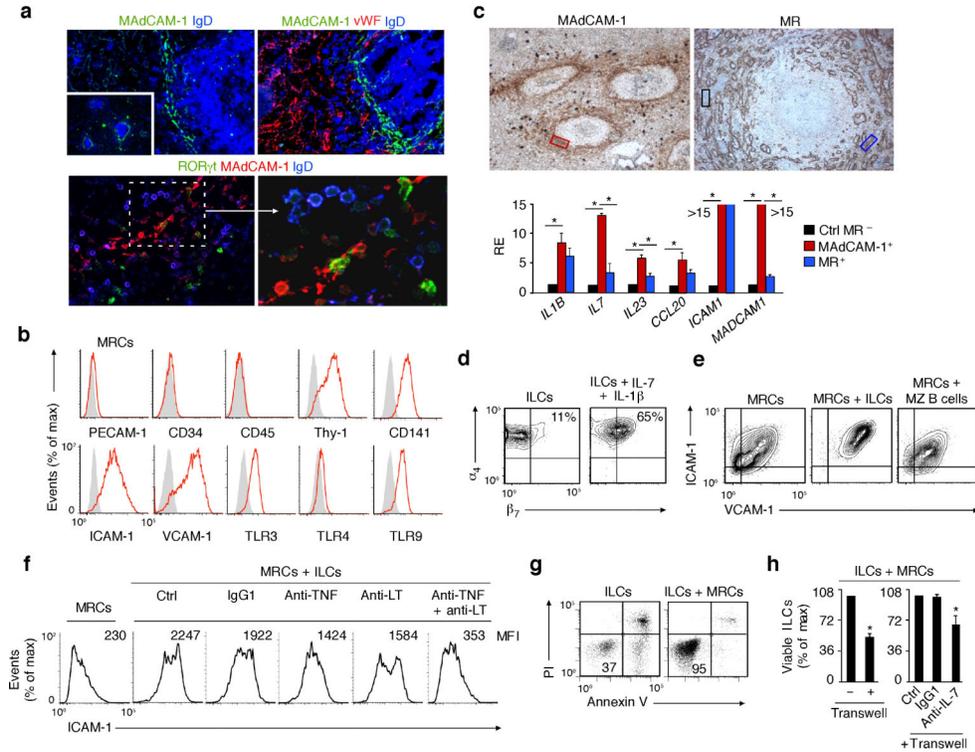


Figure 2. Human splenic ILCs establish a bidirectional crosstalk with MRCs

(a) IFA of spleens stained for MAdCAM-1 (green or red), IgD (blue), vWF (red) and/or ROR γ t (green). Original magnification, $\times 40$ (upper panels), $\times 10$ (upper inset) and $\times 63$ (bottom-left panel) and zoom $\times 2$ (bottom-right panel). (b) Flow cytometry of PECAM-1, CD34, CD45, Thy-1, CD141, ICAM-1, VCAM-1, TLR3, TLR4 and TLR9 expression by *ex vivo* expanded MRCs. Gray shading, negative control. (c) qRT-PCR of *IL1B* (IL-1 β), *IL7* (IL-7), *IL23* (IL-23), *CCL20* (CCL20), *ICAM1* (ICAM-1) and *MADCAM1* (MadCAM-1) mRNAs in MadCAM-1⁺ MRC-enriched (red box), MR⁺ SLC-enriched (blue box) and control (ctrl) MR⁻ (black box) areas microdissected from immunohistochemically stained splenic tissue (original magnification, $\times 20$). qRT-PCR results are normalized to *ACTB* (β -actin) mRNA and presented as relative expression (RE) compared with that of MR⁻ areas. (d) Flow cytometry of integrins α_4 and β_7 on freshly isolated splenic ILCs (left) or ILCs cultured with IL-1 β and IL-7 for 72 h (right). (e,f) Flow cytometry of ICAM-1 and VCAM-1 on MRCs incubated for 72 h with or without ILCs in the presence or absence of control (Ctrl), anti-TNF, anti-LT α or anti-TNF plus anti-LT α β antibodies. (g,h) Flow cytometric analysis of frequency of Annexin-V⁻propidium iodide⁻ ILCs from splenic ILCs cultured for 72 h with or without MRCs in the presence or absence of a transwell, a control (Ctrl) antibody or anti-IL-7. Error bars, s.e.m.; **P* < 0.05 (two-tailed unpaired Student's *t* test). Data summarize three experiments with one donor in each (c,h) or show one of four experiments with similar results (a,b,d,e,f,g).

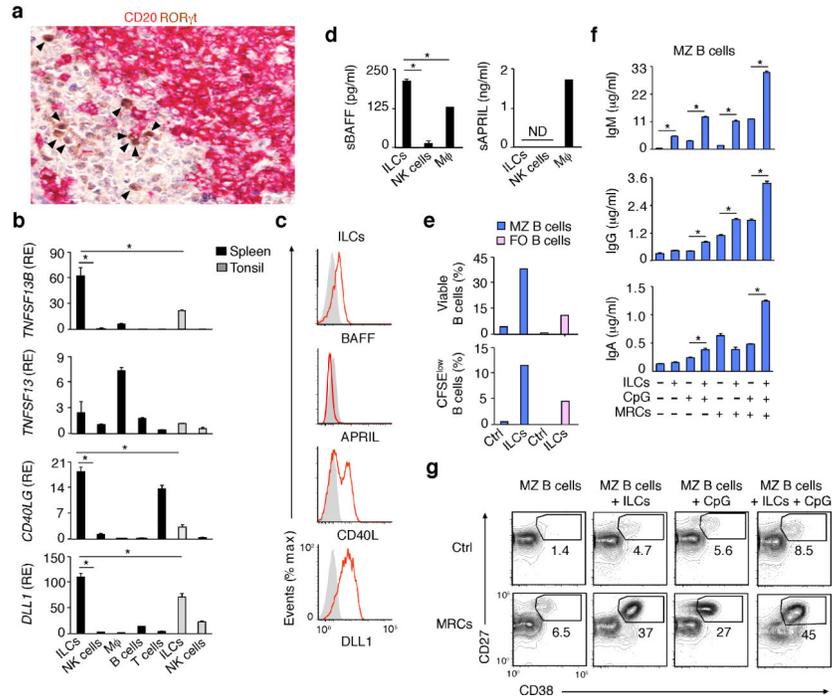


Figure 3. Human splenic ILCs express the MZ B cell-helper factors BAFF, CD40L and DLL1 and activate MZ B cells in cooperation with MRCs

(a) IHC of spleen stained for ROR γ t (brown) and CD20 (purple). Arrowheads point to ROR γ t⁺ cells. Original magnification, \times 40. (b) qRT-PCR of *TNFSF13B* (BAFF), *TNFSF13* (APRIL), *CD40LG* (CD40L) and *DLL1* (DLL1) mRNAs from splenic or tonsillar ILCs and NK cells and from splenic macrophages (M ϕ), B cells and T cells. Results are normalized to *ACTB* (β -actin) mRNA and presented as relative expression (RE) compared with that of fresh splenic NK cells. (c) Flow cytometry of BAFF, APRIL, CD40L and DLL1 on splenic ILCs exposed to IL-1 β plus IL-7 for 72 h. Gray shading, negative control. (d) ELISA of soluble BAFF and APRIL from splenic ILCs and NK cells cultured as in (c) as well as M ϕ . (e) Flow cytometric analysis of the frequency of viable Annexin-V⁻propidium iodide (PI)⁻ and divided carboxyfluorescein diacetate succinimidyl ester (CFSE)^{low} splenic MZ (blue bars) and FO (pink bars) B cells after incubation with medium alone (ctrl) or ILCs for 5 d. (f) ELISA of IgM, IgG and IgA from splenic MZ B cells incubated for 5 d with medium alone (Ctrl), ILCs, MRCs and/or CpG. (g) Frequency of CD27^{hi}CD38^{hi} plasmablasts in splenic MZ B cells cultured for 5 d as in (f). Error bars, s.e.m.; **P* < 0.05 (two-tailed unpaired Student's *t* test). Data summarize 3 measurements from 3 pooled experiments with one donor in each (b,d) or show one of three experiments with similar results (a,c,e,f with s.e.m of triplicate, g).

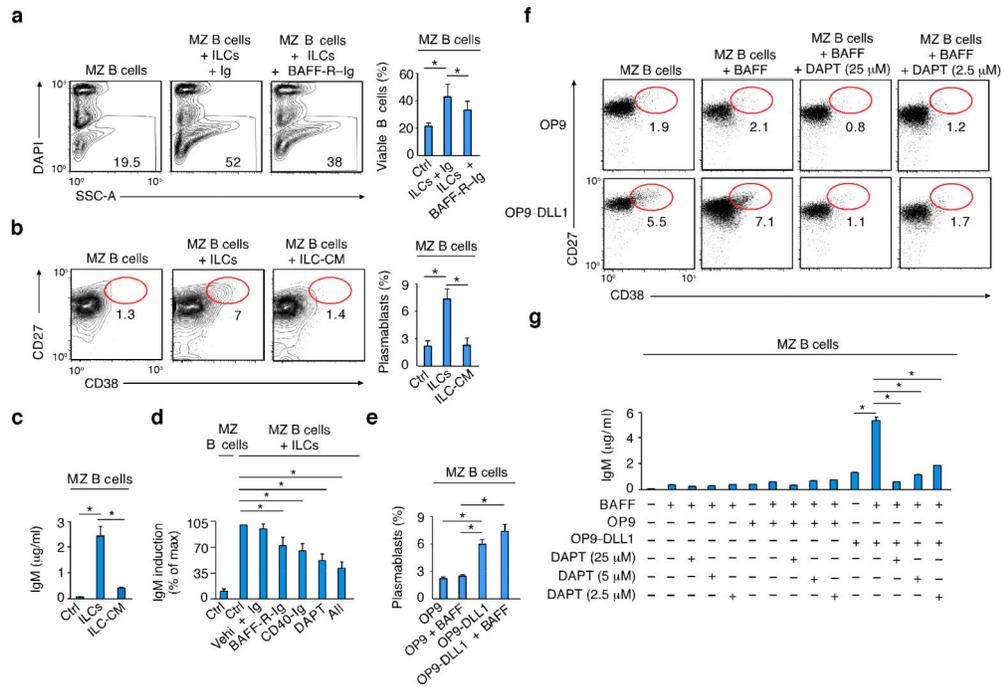


Figure 4. Human splenic ILCs help MZ B cells via BAFF, CD40L and DLL1

(a,b) Flow cytometric analysis of the frequency of viable DAPI⁻ MZ B cells from MZ B cells cultured for 5 d with or without splenic ILCs in the presence or absence of control Ig or BAFF-R-Ig. SSC-A, side scatter area. (b) Frequency of CD27^{hi}CD38^{hi} plasmablasts from splenic MZ B cells cultured for 5 d with or without splenic ILCs or ILC conditioned medium (ILC-CM). (c) ELISA of IgM from splenic MZ B cells cultured as in (b). (d) ELISA of IgM from splenic MZ B cells cultured for 5 days with medium alone (Ctrl) or ILCs in the presence or absence of control Ig and dimethyl sulfoxide vehicle (Vehi), BAFF-R-Ig, CD40-Ig, N-[N-(3,5-difluorophenacetyl)-L-alanyl]-S-phenylglycine t-butyl ester (DAPT, a NOTCH inhibitor) or a combination of BAFF-R-Ig plus CD40-R-Ig plus DAPT (All). Summary of data from 7 donors shown as % of maximal induction (100%) after MZ B cell stimulation with ILCs and control vehicle (Vehi). (e) Frequency of CD27^{hi}CD38^{hi} plasmablasts from splenic MZ B cells cultured for 5 days with control OP9 cells or DLL1-OP9 cells in the presence or absence of BAFF. (f) Frequency of CD27^{hi}CD38^{hi} plasmablasts from splenic MZ B cells cultured for 5 d with or without control (ctrl) OP9 cells or DLL1-OP9 cells in the presence or absence of BAFF and/or DAPT. (g) ELISA of IgM from splenic MZ B cells cultured as in (e). Error bars, s.e.m.; **P* < 0.05 (two-tailed unpaired Student's *t* test). Data summarize four (a), three (b,c,e) and seven (d) experiments with one donor in each or show one of three experiments with similar results (f, g with s.e.m of triplicate).

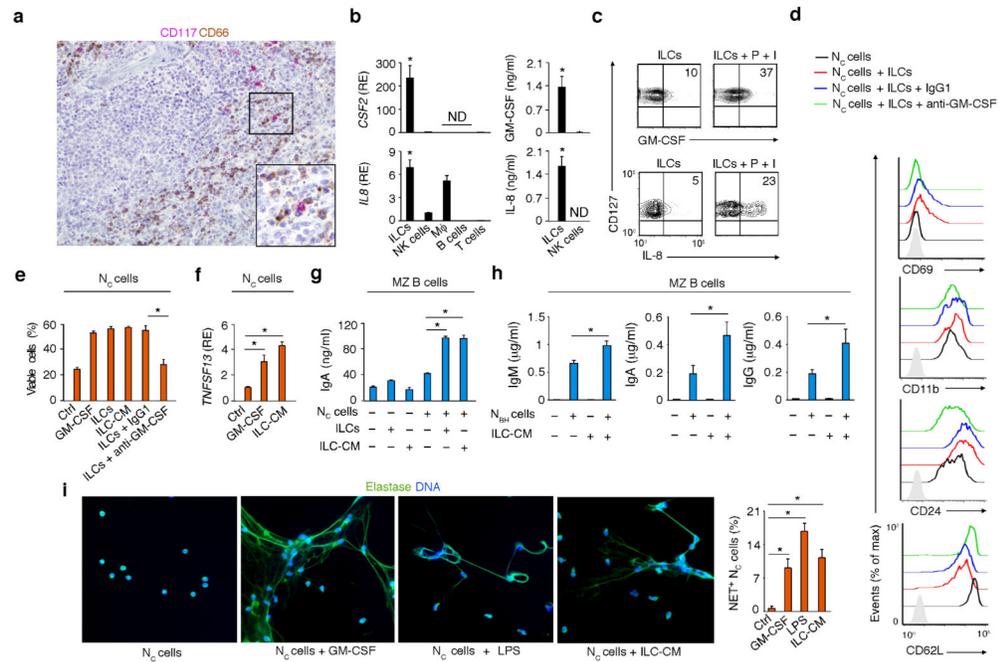


Figure 5. Human splenic ILCs activate MZ B cell-helper neutrophils via GM-CSF

(a) IHC of spleen stained for CD117 (red) and CD66 (brown). Original magnification, $\times 20$ or $\times 40$ (inset). (b) Left: qRT-PCR of *CSF2* (GM-CSF) and *IL8* (IL-8) mRNAs from splenic ILCs, NK cells, macrophages (M ϕ), B cells and T cells. *CSF2* and *IL8* data are normalized to *ACTB* (β -actin) mRNA and presented as relative expression (RE) compared with that of fresh splenic NK cells. Right: ELISA of GM-CSF and IL-8 from splenic ILCs and NK cells cultured for 48 h with IL-1 β and IL-7. (c) Flow cytometry of intracellular GMCSF and IL-8 in splenic CD127⁺ (Lin⁻CD117⁺) ILCs activated with or without PMA (P) plus ionomycin (I) for 4 h. (d) Flow cytometry of CD69, CD11b, CD24 and CD62L on Nc cells cultured for 24 h with or without splenic ILCs in the presence or absence of control IgG1 or blocking anti-GM-CSF antibodies. Gray shading: isotype control. (e) Flow cytometry of viable DAPI⁻ Nc cells cultured with medium alone (Ctrl), GM-CSF, ILCs or ILC-derived conditioned medium (ILC-CM) in the presence or absence of IgG1 or anti-GM-CSF antibodies. (f) qRT-PCR of *TNFSF13* (APRIL) mRNA from Nc cells cultured for 24 h with medium alone (Ctrl), GM-CSF or ILC-CM. Results are normalized to *ACTB* mRNA and presented as RE compared with that of Nc cells cultured with medium alone. (g) ELISA of IgA from splenic MZ B cells cultured for 5 d with or without Nc cells, ILCs and/or ILC-CM. (h) ELISA of IgM, IgA and IgG from splenic MZ B cells cultured for 5 d with or without N_{BH} cells, ILC-CM and/or N_{BH} cells preconditioned with ILC-CM. (i) IFA of elastase (green) and DAPI-stained DNA (blue) in Nc cells cultured for 3 h either with medium alone (Ctrl), GM-CSF, ILC-CM or LPS (original magnification, $\times 63$) and quantification of NET-forming Nc cells from nine $\times 10$ fields from two independent experiments. Error bars, s.e.m.; * $P < 0.05$ (two-tailed unpaired Student's *t* test). Data summarize 2-3 experiments with one donor in each (b,e,f,g,h,i) or show one of four experiments with similar results (a,c,d).

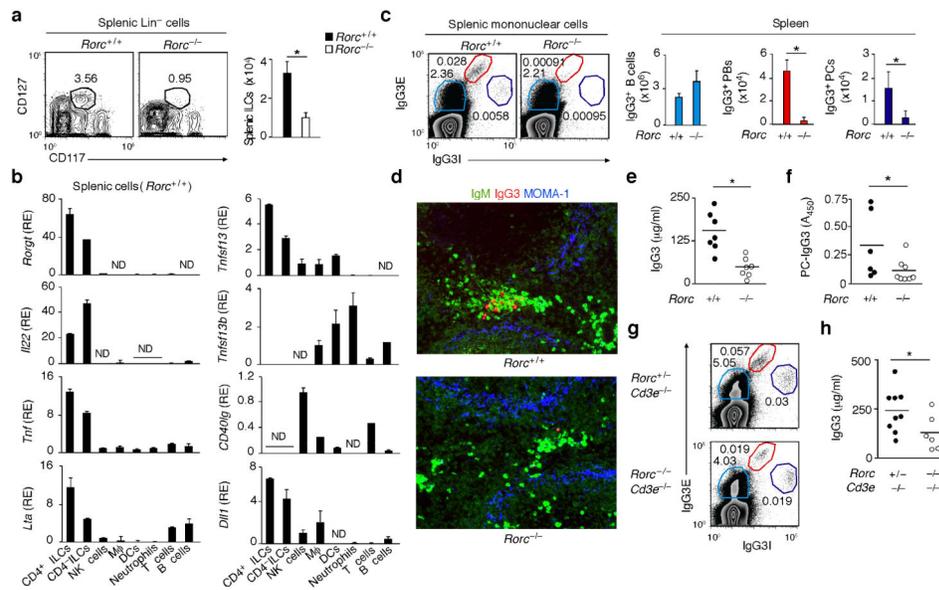


Figure 6. Mouse splenic ILCs express the plasma cell-helper factors APRIL and DLL1 and enhance TI IgG3 responses

(a) Flow cytometric analysis of frequency and absolute numbers of splenic Lin⁻ CD117⁺CD127⁺ ILCs from *Rorc*^{+/+} ($n = 3$) and *Rorc*^{-/-} mice ($n = 3$). (b) qRT-PCR of *Rorc* (ROR γ t) *Il22* (IL-22) *Tnf* (TNF), *Lta* (LT- α), *Tnfsf13* (APRIL), *Tnfsf13b* (BAFF), *CD40lg* (CD40L) and *Dll1* (DLL1) mRNAs from splenic ILCs, macrophages (M ϕ), DCs, neutrophils, NK cells, T cells or B cells. Results are normalized to *Gapdh* (glyceraldehyde 3-phosphate dehydrogenase) mRNA and presented as relative expression (RE) compared to the expression level in NK cells. (c) Flow cytometric analysis of frequency and absolute numbers of splenic IgG3E⁺IgG3I^{lo} B cells, IgG3E^{hi}IgG3I⁺ plasmablasts (PBs) and IgG3E⁺IgG3I^{hi} plasma cells (PCs) from *Rorc*^{+/+} ($n = 3$) and *Rorc*^{-/-} mice. E, extracellular; I, intracellular. (d) IFA of IgM (green), IgG3 (red) and MOMA-1 (blue) in spleens from *Rorc*^{+/+} and *Rorc*^{-/-} mice. Original magnification, $\times 20$. (e,f) ELISA of total (e) and PC-reactive (f) serum IgG3 from *Rorc*^{+/+} ($n = 6-7$) and *Rorc*^{-/-} ($n = 7-8$) mice. (g) Frequency of IgG3-expressing B cells, PBs and PCs from spleens of *Rorc*^{+/-}*Cd3e*^{-/-} and *Rorc*^{-/-}*Cd3e*^{-/-} mice. (h) Serum IgG3 from *Rorc*^{+/-}*Cd3e*^{-/-} ($n = 9$) and *Rorc*^{-/-}*Cd3e*^{-/-} ($n = 6$) mice. Error bars, s.e.m. (b), s.d. (a,c); * $P < 0.05$ (one-tailed unpaired Student's t test). Data show one of four experiments with similar results (a,d,g; c: cytograms) or summarize results from 3-8 animals per each group (a,c; bars; b,e,f,h).

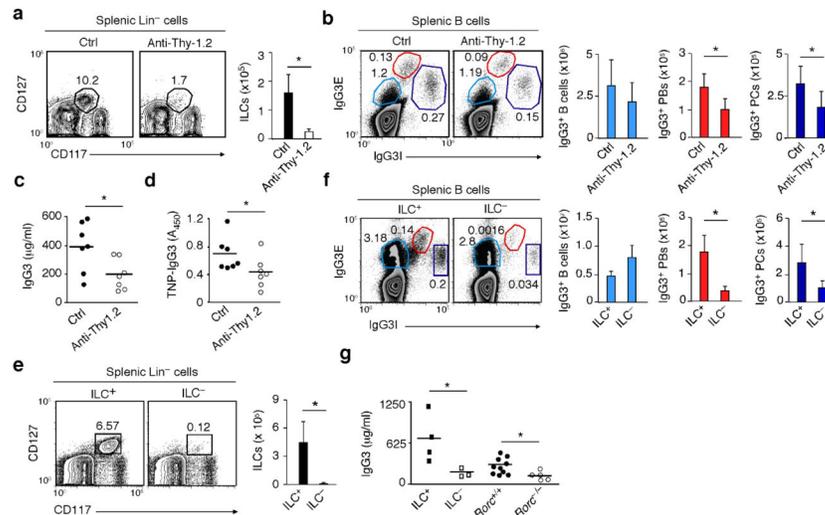


Figure 7. Mouse splenic ILCs help plasmablasts and plasma cells emerging from TI IgG3 responses

(a) Flow cytometric analysis of frequency and absolute numbers of splenic Lin⁻CD117⁺CD127⁺ ILCs from Thy-1-disparate chimeric *Rag1*^{-/-} mice treated with control (ctrl) ($n = 7$) or anti-Thy.1.2 antibodies ($n = 7$). (b) Frequency and absolute numbers of splenic IgG3E⁺IgG3I^{lo} B cells, IgG3E^{hi}IgG3I⁺ plasmablasts (PBs) and IgG3E-IgG3I^{hi} plasma cells (PCs) from Ctrl or ILC-depleted mice obtained as in (a). E, extracellular; I, intracellular. (c,d) ELISA of pre-immune total (c) and PC-reactive (d) serum IgG3 from Ctrl or ILC-depleted mice obtained as in (a). (e,f) Frequency and absolute numbers of splenic Lin⁻CD117⁺CD127⁺ ILCs (e) and IgG3-expressing splenic B cells, PBs and PCs (f) from ILC⁺ ($n = 3$) or ILC⁻ ($n = 3$) bone marrow chimeric mice. (g) Pre-immune total serum IgG3 from ILC⁺ ($n = 4$) and ILC⁻ ($n = 3$) mice, age-matched *Rorc*^{+/+} mice ($n = 10$), and *Rorc*^{-/-} mice ($n = 5$). Error bars, s.d.; * $P < 0.05$ (one-tailed unpaired Student's *t* test). Data show one of four experiments with similar results (a,b,e,f: cytograms) or summarize results from 3-10 animals per each group (a,b,e,f: bars; c,d,g).

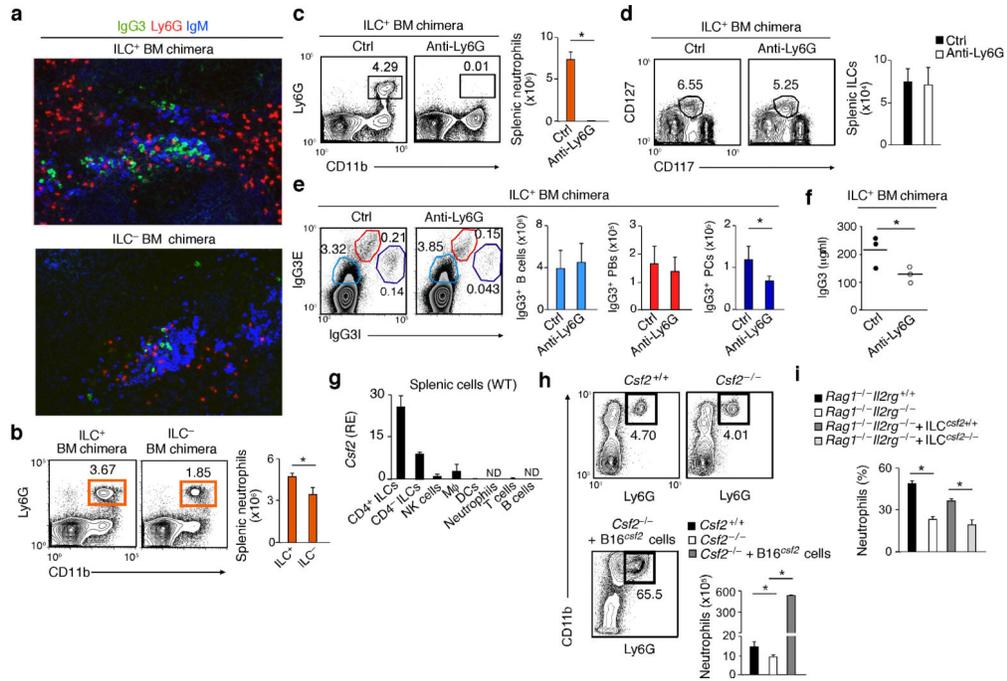


Figure 8. Mouse splenic neutrophils help plasma cells emerging from TI IgG3 responses and receive homeostatic signals from ILCs via GM-CSF

(a) IFA of IgG3 (green), Ly6G (red) and IgM (blue) in spleens from ILC⁺ or ILC⁻ bone marrow chimeric mice. Original magnification, ×20. (b) Flow cytometric analysis of the frequency of splenic Ly6G⁺CD11b⁺ neutrophils from ILC⁺ ($n = 3$) or ILC⁻ ($n = 3$) mice. (c) Frequency and absolute numbers of splenic Ly6G⁺CD11b⁺ neutrophils from ILC⁺ mice treated with control (Ctrl) ($n = 4$) or anti-Ly6G antibodies ($n = 4$). (d,e) Frequency and absolute numbers of splenic Lin⁻CD117⁺CD127⁺ (d) as well as splenic IgG3E⁺IgG3I^{lo} B cells, IgG3E^{hi}IgG3I^{hi} plasmablasts (PBs) and IgG3E⁺IgG3I^{hi} plasma cells (PCs) from ILC⁺ mice treated as in (c). E, extracellular; I, intracellular. (f) ELISA of total serum IgG3 from ILC⁺ mice treated as in (c). (g) qRT-PCR of *Csf2* (GM-CSF) mRNA from splenic ILCs, macrophages (Mφ), DCs, neutrophils, NK cells, T cells or B cells. Results are normalized to *Gapdh* (glyceraldehyde 3-phosphate dehydrogenase) mRNA and presented as relative expression (RE) compared to the expression level in NK cells. (h) Frequency and absolute numbers of splenic Ly6G⁺CD11b⁺ neutrophils from *Csf2*^{+/+} ($n = 8$) and *Csf2*^{-/-} ($n = 12$) mice before and after transfer of B16^{Csf2} cells overexpressing GM-CSF ($n = 3$). (i) Frequency of splenic neutrophils in ILC-sufficient *Rag1*^{-/-}*Il2rg*^{+/+} mice or ILC-insufficient *Rag1*^{-/-}*Il2rg*^{-/-} mice reconstituted or not with gut ILCs from *Csf2*^{+/+} or *Csf2*^{-/-} mice ($n = 3$). Error bars, s.e.m. (g-i), s.d. (b-e); * $P < 0.05$ (one-tailed unpaired Student's t test). Data summarize results from at least two experiments with 3 mice in each group (b,c,d,e,h: bars; f,g,i) or show one of at least three experiments with similar results (a and b,c,d,e,h: cytograms).