Intracellular sterol sensing controls intestinal B cell differentiation

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Summary
Intestinal B cell response is critical for the maintenance of gut homeostasis, yet environmental signals that control B cell metabolism and effector function remain poorly characterized. Here, we show that in Peyer's patches germinal center (GC) B cells are sensitive to 25-hydorchycholesterol (25-HC), an oxidized metabolite of cholesterol produced by the enzyme cholesterol 25-hydroxylase (CH25H). In mice lacking CH25H, antigen-specific GC B cells show an increased cholesterol metabolic signature and preferentially differentiate in plasma cells (PCs), therefore inducing a stronger intestinal IgA response upon immunization or infection. GC B cells express the sterol sensor SREBP2 and use it to sense 25-HC. Deletion of SREBP2 from GC B cells prevents PC differentiation and forces the maintenance of GC identity. GC localized oxysterol production by follicular dendritic cells is central in dictating GC metabolism and imposing B cell fate. Our findings show that the 25-HC-SREBP2 axis shapes B cell effector function in intestinal lymphoid organs and indicate that dietary cholesterol can instruct local B cell response.

Introduction
In the small intestine, most of the T cell-dependent IgA response originates in Peyer's patches (PPs), the lymphoid organs closest to the intestinal lumen. The anatomical position of PPs place them at the primary site of absorption for both dietary and bacterial products, therefore exposing them to a variety of metabolic cues. While metabolites absorbed from the intestinal lumen can shape PP organogenesis[1] and migration of adaptive immune cells[2], mainly via retinoic acid receptor signaling, a precise, mechanistic understanding of how diet-derived metabolites impact B cell differentiation and IgA production in PPs is lacking.
In humans and mice, cholesterol absorption is restricted to the small intestine and cholesterol byproducts[3] possess known modulatory activities on lymphocytes. Diets high in cholesterol are shown to impact adaptive immune responses, including B cell activity[4]. Among the cholesterol byproducts with immunomodulatory properties, 25-hydroxycholesterol (25-HC) plays a multifaceted role in immune responses. 25-HC, the oxysterol produced by the enzyme cholesterol 25-hydroxylase (CH25H), is the substrate necessary for the generation of 7α,25-HC, the strongest ligand for the orphan G protein-coupled receptor 183 (GPR183, also known as EBI2). Despite 7α,25-HC and 25-HC differing only in the hydroxylated group at the 7α position, 25-HC does not bind and activate EBI2. EBI2 drives migration of several immune cell types[5-7,8,9,10-12,] and CH25H is required for this process. 25-HC can also work in a EBI2-independent role, primarily as an innate immune modulator[13,14,15,16,17]. Whether 25-HC is also involved in adaptive immune responses per se remains to be determined.
25-HC was originally identified as a potent inhibitor of sterol biosynthesis, with higher suppressive potency than cholesterol itself[18]. 25-HC prevents sterol biosynthesis by maintaining inactive sterol response element-binding proteins 2 (SREBP2) in the endoplasmic reticulum (ER)[18]. When the intracellular level of 25-HC decreases, SREBP2 moves into the Golgi, where it is cleaved by site 1 and site 2 proteases and subsequently activated as a transcription factor [19]. Despite its critical role in maintaining cellular cholesterol homeostasis, the role of the SREBP2 pathway in immune cells has been poorly characterized[20], and its impact on B cell fate, especially during intestinal responses, is unknown.

B cell response in the PPs, and the subsequent IgA production, are driven by continual exposure to microbial-derived molecules[21,22,23,24], as seen by the presence of chronic germinal center (GC) composed of discrete clusters of GC B cells, T follicular helper cells (Tfh cells) and follicular dendritic cells (FDCs) in the center of B cell follicles. PP GC B cells give rise to plasma cells (PCs) and memory B cells (MBCs): however, the impact of intestinal-derived metabolites on GC B cell differentiation remains uninvestigated. In this study, we sought to further define how 25-HC controls intestinal IgA response and to understand how dietary cholesterol restrains PC generation during enteric B cell response.
Here, we report that B cell differentiation to IgA PCs in intestinal lymphoid organs is controlled by the oxysterol 25-HC independently of the GPR183/EBI2 pathway. Mice lacking *Ch25h* have normal IgA class switch recombination *in vivo* but show increased antigen (Ag)-specific IgA response to toxin and enteric pathogen. In PPs, FDCs are required for the generation of the 25-HC niche that mediates suppression of cholesterol metabolic genes in GC B cells. Mechanistically, 25-HC inhibits SREBP2 activation, nuclear translocation and transcriptional activity: in the absence of 25-HC, SREBP2 target genes are upregulated in GC B cells and ectopic expression of *Srebf2* in GC B cells drives increased PC differentiation. In contrast, GC B cells lacking SREBP2 maintain a GC phenotype and have reduced propensity to become PCs both *in vitro* and *in vivo*. Our results establish a role for a dietary-derived sterol in controlling B cell fate in the mucosal draining lymph node during T cell-dependent IgA response.

Results

CH25H controls antigen-specific intestinal IgA response independently of EBI2.

To measure the impact of the 25-HC producing enzyme CH25H on mucosal B cell responses, we immunized *Ch25h*$^{-/-}$ mice and co-housed, littermate controls (LMCs) with the intestinal antigen cholera toxin (CT) and quantified the IgA response in the intestinal lamina propria (LP). While the IgA^{+} PC fraction (**Fig.1A**) was identical between genotypes, CT-specific IgA-secreting cells (CT-IgA-ASC) were present in higher numbers in the duodenum of mice lacking CH25H (**Fig.S1A**): the total number of IgA-ASC was not different between genotypes (**Fig.S1B**), leading to a higher frequency of CT-IgA-ASC in *Ch25h*$^{-/-}$ mice (**Fig.1B**). In line with the overall distribution of IgA-ASC in the small intestine (**Fig.S1C**), we only observed a consistent frequency of IgA-ASC in the duodenum, with very little IgA or CT-IgA-ASC in the ileum of *Ch25h*$^{-/-}$ mice (**Fig.S1D,E**). Since CH25H is required for the generation of 7α,25-HC, the most potent ligand for EBI2, a GPCR that is expressed on B cells and controls B cell positioning and differentiation, we tested whether CH25H deficiency might impact Ag-specific IgA response through EBI2 by immunizing *Ebi2*$^{-/-}$ and LMCs. *Ebi2*$^{-/-}$ mice showed unchanged number of CT-IgA-ASC, IgA-ASC (**Fig.S1F, G**), and overall frequency of CT-IgA-ASC (**Fig.1C**) suggesting that CH25H controls the intestinal humoral response independently of EBI2 migration. CT-specific IgA titers were increased in the small intestinal lavage (**Fig.1D**), but not in the serum (**Fig.1E**), highlighting that CH25H is predominantly affecting intestinal, but not systemic, antigen-specific IgA responses.

To further investigate the role of CH25H on intestinal IgA response, we infected *Ch25h*$^{-/-}$ mice and LMCs with *aroA*-deficient *(ΔAroA*) *Salmonella enterica Serovar Typhimurium* (*ST*), a defective strain of ST that has an intact route of entry into PPs but reduced infectivity[25], allowing for the measurement of IgA response in B6 mice[26]. In absence of CH25H, we observed an increased number of ST-IgA-ASC in the LP (**Fig.1F**).

Overall our results showed that mice lacking CH25H are characterized by an enhanced Ag-specific IgA response against toxin and enteric bacteria, suggesting that 25-HC can control T cell-dependent IgA responses.

PC-generation, not class switch recombination, is controlled by 25-HC in PPs.

IgA-secreting PCs represent the terminal differentiation step of B cells stimulated by intestinal antigen in the PPs. Upon Ag stimulation and GC entry, GC B cells differentiate to IgA PC that egress from the PP and migrate to LP using a combination of integrin and chemokine receptors. We therefore asked whether CH25H's effect on Ag-specific IgA ASC generation was anatomically restricted and initiated in PPs. To this end we took advantage of FTY720, a sphingosine 1-phosphate receptor (S1PR1) antagonist that can prevent IgA PC egress from PPs[27].

We immunized *Ch25h*$^{-/-}$ mice and LMCs with CT and administered FTY720 to trap newly generated, Ag-specific PCs in PPs. As expected, FTY720 treatment increased the number of IgA-ASC in the PPs (**Fig.2A, Fig.S2A**), but only *Ch25h*–deficient mice showed an enhanced Ag-specific PC response (**Fig.2B**, **Fig.S2B**). In contrast, mice lacking EBI2 showed no increase in CT-response, highlighting how CH25H, but not EBI2, is involved in controlling specific IgA response in PPs (**Fig.2C, Fig.S2C**). In line with the data from LP, blocking of S1PR1-dependent ASC egress with FTY720 during ST infection also enhanced IgA-ASC response in PPs of *Ch25h*$^{-/-}$ mice. (**Fig.2D**). While *Ebi2*$^{-/-}$ mice have showed systemic decrease of PC response, we did not observe an effect on PC generation in PPs (**Fig.S2D**), and *Ch25h*$^{-/-}$ mice, but not *Ebi2*$^{-/-}$ mice, were able to generate more Ag-specific PCs. These results suggest the existence of a process dependent on CH25H, but independent of EBI2, that shapes IgA specific response in PPs.

Previous work has suggested that 25-HC could block IgA class switching[28] however ex-vivo FACS analysis of B cell subsets from PPs and mesenteric LN (mLN) in *Ch25h*$^{+/-}$ and *Ch25h*$^{-/-}$ showed that lack of 25-HC has no impact on the ability of B cells to undergo IgA class switch recombination (**Fig.2E, Fig.S2E**).

To assess whether 25-HC could directly restrain PC generation from GC B cells, we used NB21 culture assay, in which the feeder cell layer expresses CD40L, BAFF and IL-21[29]. Sorted GC B cells were cocultured with NB21 and then treated with different oxysterols: 25-HC, but not 7α,25-HC or cholesterol, decreased the Ab-secreting cell generation (**Fig.2F**). Similar results were obtained with GC B cells stimulated with anti-CD40 and cultured with NB21 supernatant, ruling out a direct effect of 25-HC on NB21 cells (**Fig. S2F**), and suggests that 25-HC can act directly on GC B cells and restrain their ability to differentiate in PCs. Overall our data suggest that intestinal and systemic antibody responses are differentially dependent on oxysterols, with 25-HC controlling local Ag-specific PC response in an EBI2-independent fashion during enteric response.

FDCs contribute to 25-HC PP levels and dietary cholesterol reduces IgA response via CH25H.

25-HC can be secreted[13] and diffuse to the surrounding tissue[3,7] to generate a 25-HC niche, a microanatomical area rich in 25-HC that can regulate nearby immune cells in a EBI2-independent way. *Ch25h* expression has been reported in several cells, both of hematopoietic and non-hematopoietic origin, mainly in the context of EBI2 ligand generation: however, the contribution of *Ch25h*-expressing cells to the 25-HC niche *in vivo* is less clear. While the EBI2 ligand 7α,25-HC can be

easily measured in a transwell assay based on EBI2-mediated migration, 25-HC does not bind EBI2 and therefore remains undetected in that setting. To unambiguously measure 25-HC in tissue extracts, we designed a modified transwell assay and verified it using synthetic oxysterols (**Fig.S3A**). We first inactivated 7α,25-HC by incubating it with 293T cells expressing HSD3B7, a 7α-oxidized cholesterol dehydrogenase enzyme that oxidizes all existing EBI2 ligand; subsequently we harvested and incubated 25-HC-containing extract with 293T cells expressing CYP7B1. This second incubation converted 25-HC to 7α,25-HC, which was now measured in the transwell using EBI2-expressing M12 cells (**Fig.S3B**). Analysis of intestine and intestinal secondary lymphoid organs (SLO) showed that 25-HC can be measured in all of these tissues, and requires the activity of CH25H enzyme: thus, intestinal cells producing 25-HC that is not converted into EBI2 ligand, exist *in vivo* in unmanipulated mice (**Fig.S3C**).
PC differentiation largely takes place in the GC, so we reasoned that a 25-HC niche anatomically restricted to the GC could influence activated B cell fate decision. In GC, Follicular dendritic cells (FDCs), stromal cells (SCs) that display antigen in the light zone for BCR testing, have high levels of *Hsd3b7* transcripts[7] and can inactivate EBI2 ligand, thereby maintaining EBI2 expressing follicular B cells, outside of the GC, while allowing GC B cells, which do not express EBI2, to reach and remain inside the GC structure[3]. As 25-HC is not sensitive to HSD3B7 activity, the GC environment is likely to contain higher concentrations of 25-HC compared to the follicles: since EBI2 ligand regulates naïve B cell positioning, we posit that most of 25-HC in the follicles is not freely available as it is converted to 7α,25-HC by CYP7B1 (**Fig.S3D**). FDCs express *Ch25h*, but whether FDCs are critical to establish GC 25-HC is unknown.
We employed two different strategies to assess whether FDCs are the primary source of 25-HC in PPs. First, we used CD21-DTR mice, where *Cd21*-Cre mice are crossed with mice carrying simian diphtheria toxin receptor (DTR) downstream of a floxed stop element in the ROSA26 locus. We generated bone marrow (BM) chimeras using CD21-DTR mice as hosts: in this setting the DTR is expressed only in non-hematopoietic CD21-expressing cells, i.e. FDCs [30],[7]. Short term DT injection led to FDC ablation: while we only observed a small reduction in EBI2, 25-HC was largely reduced in PPs (**Fig.3A**), suggesting that cells other than FDCs[31], might contribute to the follicular production of EBI2 ligand[32]. As a second test of the FDC requirement in 25-HC niche generation, we depleted FDCs by blocking lymphotoxin (LT) signaling through LT-beta receptor (LTβR) on FDCs using soluble LTβR fused to human IgG Fc (LTβR-Fc). The blockade prevents the interaction between lymphotoxin-$\alpha_2\beta_1$-expressing GC B cells and LTβR-expressing FDCs, leading to FDC death[54]. FDC depletion only marginally impacted EBI2 ligand generation but led to a dramatic reduction of 25-HC (**Fig.3B**), strongly suggesting that FDCs primarily contribute to the 25-HC niche in the GC.
Since CH25H requires cholesterol to generate 25-HC we asked whether changes in dietary cholesterol content could shape the intestinal 25-HC niche. We fed mice with normal food (NF) or normal food with the exclusive addition of 0.15% cholesterol found in the Western diet (high cholesterol food, HCF) for 1 week and measured 25-HC in PPs. We found that 25-HC was increased in animal fed HCF (**Fig.3C**), suggesting that altering the intestinal absorption of dietary cholesterol can shape the 25-HC niche. EBI2 ligand was also increased in PPs, highlighting a previously unreported link between dietary cholesterol and immunomodulatory sterols.
We then asked whether diet-dependent changes in the PP 25-HC niche could influence B cell fate: mice fed with different diets were immunized with CT and treated with FTY720. WT mice fed HCF showed decreased generation of CT-IgA-ASC compared to WT mice fed with NF. In contrast, *Ch25h*$^{-/-}$ mice showed only a modest reduction in CT-IgA-ASC (**Fig.3D**).
We were able to observe a similar effect of HCF and CH25H deficiency on CT-IgA-ASC in the mLN (**Fig.S3E,F),** possibly extending our 25-HC dependent restraint on PC generation to other intestinal SLOs.

SREBP2 activation is induced by BCR signaling and is controlled in vivo by 25-HC.
Due to the role of 25-HC in controlling activation and nuclear translocation of sterol response element-binding proteins (SREBP2)[18], [19]we analyzed the SREBP2 transcriptional activity in B cells. Since SREBP2 regulates its own transcription, we initially measured *Srebf2* transcript in PP B cell subsets (**Fig.4A**). GC B cells and PC showed the highest level of *Srebf2*, followed by MBCs, while follicular B cells showed the lowest *Srebf2* transcription. Similar dynamics of *Srebf2* transcription was observed in B cell subsets from the mLN (**Fig.S4A**). Transcripts for *Srebf1a*, an isoform of SREBP1 protein than can control certain aspects of lipid metabolism, were detected at a considerably lower levels in both PPs and the mLN, with both GC and PC transcribing *Srebf1a*. (**Fig.4B, Fig.S4B**). To understand whether 25-HC can control SREBP2 levels *in vivo* we analyzed *Srebf2* transcript and SREBP2 target gene transcripts in *Ch25h*$^{-/-}$ mice. *Srebf2* expression is increased in GC B cells and PCs in both PPs and mLN in absence of 25-HC (**Fig. 4C, Fig.S4C**) while no difference was observed in follicular B cells and a minimal increase was present in MBCs. These results demonstrate that *in vivo* 25-HC levels can control SREBP2 activation in some antigen-experienced B cell subsets.

SREBP2 induction is generally driven by low levels of intracellular sterols, including selected oxysterol concentrations, but whether other signals can control SREBP2 activation is unclear. All activated B cell subsets, but not naïve B cells, upregulate *Srebf2* transcripts to some extent, thus we reasoned that BCR stimulation and T cell help could possibly induce SREBP2

activation. Follicular B cells stimulated with anti-IgM for 16h showed *Srebf2* induction and increase in SREBP2 target genes *Hmgcr*, *Hmgcs1* and *Sqle* (**Fig.4D**). In contrast anti-CD40 stimulation, that mimics T cell help, had little effect on SREBP2-target *Hmgcr*, *Hmgcs1* genes at a later time point compared to stimulation with both anti-IgM and anti-CD40, but it was able to upregulate *Sqle* transcription (**Fig.S4D**). Transcription of SREBP2 target genes suggest that SREBP2 abundance is increased in BCR-stimulated B cells. We first analyzed the amount of SREBP2 by staining stimulated B cells intracellularly with an antibody that recognizes a site in SREBP2 that is only present in the ER-resident, unprocessed SREBP2. In line with the qPCR data, BCR stimulation drove SREBP2 upregulation (**Fig.S4E**). SREBP2 function is mediated by its nuclear translocation. Therefore we sought to investigate the cellular dynamics of SREBP2 in stimulated B cells by flow cytometry analysis of transcription factor (TF) in isolated cell nuclei[33]. We validated the nuclear isolation by analyzing the B cell line WEHI expressing cytosolic GFP, nuclear RFP (H2B-RFP) and stained with fluorescent ER tracker: this strategy allowed for the quantification of nuclear TFs at single cell level without ER contamination, which is critical since inactive SREBP2 is sequestered in the ER (**Fig.S4F**). Using an antibody directed against the N-terminus of SREBP2, that functions as TF, B cells stimulated through the BCR induced SREBP2 activation and its nuclear translocation (**Fig.4E).** IRF4, a transcription factor that is upregulated upon BCR stimulation and is critical for PC formation, also translocated to the nucleus in a similar fashion (**Fig.S4G**).
Since SREBP2 induction is controlled by BCR stimulation, and the 25-HC niche requires FDCs, cells responsible for displaying antigen to GC B cells for BCR testing, we hypothesized that modulation of SREBP2 by 25-HC would be restricted to the GC in the light zone (LZ). Sorted LZ and DZ GC B cells from *Ch25h*$^{-/-}$ mice and LMCs clearly indicated that both *Srebf2* and SREBP2 target genes are induced to a greater extent in LZ compared to DZ and restrained by 25-HCs (**Fig.4F**, **Fig.S4H**). Since CH25H deficiency leads to increased GC-derived PC, we analyzed *Prdm1*, the gene that encodes for Blimp-1, the master transcription factor required for PC differentiation. We observed that LZ and DZ GC B cells from PPs of *Ch25h*$^{-/-}$ mice upregulate *Prdm1* compared to LMCs GC B cells (**Fig.4G**).
Together, our data demonstrate that *in vivo*, 25-HC restrains cholesterolgenic genes transcription in anatomically distinct subsets of GC B cells and highlights a process for how 25-HC could restrain initiation of PC transcriptional program in GC B cells, possibly via SREBP2 inhibition.

Ectopic SREBP2 activation is sufficient to drive PC differentiation.
To test whether SREBP2 activation could drive B cell differentiation into PCs per se, we sought to manipulate SREBP2 cellular localization, and therefore activity, in PP GC B cells. We retrovirally transduced BM cells expressing Cre in the *Aicda* locus and tdTom in the ROSA26 locus(*Aicda*$^{Cre/+}$ *Rosa26*$^{tdTom/+}$) with retrovirus encoding constitutively active nuclear (n) form of SREBP2 downstream cassette of *lox*P site–stop sequence– *lox*P site cassette (followed by an internal ribosomal entry site and a CD90.1 (Thy-1.1) reporter), which results in restricted expression of nSREBP2 to GC B cells and post-GC B cells (MBCs and PCs) (**Fig. 5A)** Overexpression of nSREBP1a was used as control. Since nSREBPs can translocate into the nucleus despite high intracellular sterol concentration, this approach uncouples intracellular metabolism and SREBP2 activation in the context of B cell differentiation. Active SREBP2 led to a striking increase in PC generation in PPs, while active SREBP1a did not result in preferential B cell fate compared to empty vector (EV) transduced cells (**Fig. 5B, C**). The increased PC generation in B cells with active nSREBP2 and the unaffected B cell fate in presence of nSREBP1a is in line with the notion that 25-HC specifically controls SREBP2, but not SREBP1 processing[13]. Untransduced B cells showed no preferential differentiation, highlighting how the alterations in cholesterol metabolism affect activated B cell identity in a strictly intrinsic fashion (**Fig.S5A**). In LP, (**Fig. 5D)** SREBP-2 transduced cells were present in higher frequency compared to both EV and SREBP1a transduced cells, showing that increased SREBP2 activity can efficiently generate PCs able to reach the tissue (**Fig 5E, F**). These data establish that alteration of SREBP2 activity instruct B cells to assume distinct fates.

SREBP2 is required for the generation of PC in vitro and in vivo
To assess whether SREBP2 is necessary for PC differentiation, we generated *Aicda*^{cre} *Srebf2* $^{flox/flox}$ mice: in these animals GC B cells are unable to transcribe and activate SREBP2. We sorted GC from *Aicda*^{cre} *Srebf2* $^{flox/flox}$, *Aicda*^{cre} *Srebf2* $^{flox/+}$ and *Aicda*^{cre} *Srebf2* $^{+/+}$ and assessed their potential for PC differentiation by co-culturing them with NB21 cells: *Srebf2*-deficient GC B cells were largely unable to differentiate into PCs and maintained their GC B cell identity (**Fig.6A,B,C**). In line with previous reports of SREBP2 transcriptional activity, *Srebf2* gene dosage can affect cell fate, with GC B cells from *Aicda*^{cre} *Srebf2* $^{flox/+}$ showing an intermediate PC output compared to GC B cells from *Aicda*^{cre} *Srebf2* $^{flox/flox}$ and *Aicda*^{cre} *Srebf2* $^{+/+}$ (**Fig. 6A, B,C**). Overall cellular recovery was reduced in cultures containing GC B cells from *Aicda*^{cre} *Srebf2* $^{flox/flox}$, suggesting that SREBP2 deficiency was impairing transition to PCs and impacting GC survival (**Fig.S5B**). IgA secretion was also reduced from *Aicda*^{cre} *Srebf2* $^{flox/flox}$ GC B cell cultures (**Fig.6D**): since IgA class switch was unaffected in GC B cells lacking SREBP2 (**Fig.S5C**), our data show that SREBP2 is required for allowing efficient GC differentiation into PCs. While *Aicda*^{cre} *Srebf2* $^{flox/flox}$ did not show altered overall IgA-ASC in LP (**Fig. S5D**), oral immunization with CT

led to a drastic decrease of Ag-specific-IgA ASC (**Fig. 6F**). Taken together, our data shows an undescribed role for the sterol sensor SREBP2 in controlling B cell fate during intestinal immune response.

Discussion

In summary we find that oxysterol 25-hydroxycholesterol (25-HC) acts on GC B cells in PPs to restrain PC differentiation via inhibition of SREBP2 activation. Our data support a novel, direct effect of 25-HC on adaptive immune cells independent of its role in the generation of 7α,25-dihydroxycholesterol (7α,25-HC), the ligand for the surface receptor EBI2.

We have provided evidence that in absence of 25-HC-producing enzyme CH25H, the Ag-specific IgA PC response is enhanced during oral immunization and enteric infection. CH25H is required in PPs to control PC output during T cell-dependent responses, without any measurable effect on IgA CSR. Our *in vitro* experiments suggest that 25-HC directly affects the ability of GC B cells to undergo PC differentiation. Our data also showed that FDCs are the dominant cell type required to establish the 25-HC niche in the GCs. FDCs have a well-established role in maintaining the anatomical organization of follicular and GC response by enzymatic degradation of metabolites acting on GPCR as described for the EBI2 ligand 7α,25-HC and the P2YR8 ligand S-geranylgeranyl-L-glutathione. Our data show for the first time that a FDC-derived metabolite can be locally maintained and act directly on GC B cells. Other cells have been reported to express *Ch25h*, but anatomy places FDCs at a prime location to directly control GC B cells via 25-HC in an EBI2 independent fashion, without requiring long range tissue diffusion.

Moreover, we demonstrated that increasing cholesterol in the diet is sufficient to modulate 25-HC tissue concentration in PPs and to reduce Ag-specific IgA response. Our data do not exclude that in the presence of increased concentration of tissue cholesterol, cells other than FDCs could produce 25-HC outside of the GC, and therefore contribute to PC inhibition at different stages of B cell activation.

25-HC's ability to block activation, and subsequent transcriptional activity of SREBP2, an ER protein that senses intracellular sterol concentration, is well documented, but its regulation remains largely unexplored in adaptive immune cells. B cells activated *in vitro* by IgM BCR stimulation upregulate SREBP2. Similarly, LZ GC B cells, that routinely test their BCR against Ag displayed by FDCs, are also characterized by increased SREBP2 induction and transcriptional activity. We suggest a model in which SREBP2 activation is modulated by 25-HC in the LZ to reduce PC differentiation. Alterations of SREBP2 levels sharply affect GC B cell fate, suggesting that environmental cues able to control SREBP2 activation might play a role in GC output in addition to the widely studied BCR affinity and T cell help.
Although our over-expression results suggested a striking effect for SREBP2 activation in PC differentiation, the *in vivo* phenotype in mice lacking the SREBP2 inhibitor 25-HC was only evident when Ag-specific response was tracked and was reduced in magnitude. While our data suggest a dominant role for CH25H in mediating dietary cholesterol inhibition of PC response, we cannot exclude the possibility that other oxysterols or cholesterol itself could play a compensatory role in absence of 25-HC. In line with the negative feedback exerted by cholesterol and selected oxysterols on cholesterol metabolism via SREBP2 inhibition, albeit at higher concentration, CH25H-independent PC inhibition could take place when cholesterologenic diet is prolonged or contains an even higher amount of cholesterol.
Moreover, external cues could modulate the strictness of 25-HC/SREBP2 crosstalk in GC B cells, including diet and commensal composition. The original report describing *Ch25h*$^{-/-}$ mice reported increased IgA production: while the authors explained it as an effect on CSR based on *in vitro* experiments, we could not see 25-HC inhibition of IgA CSR in PP or MLN in our mouse colony. However, the initial IgA phenotype reported by Russell could be easily reconciled with our data by invoking a SREBP2 activation and increased PC differentiation mediated by a stronger T cell dependent commensal response in their colony.

We propose a model in which FDCs convert dietary cholesterol into 25-HC in PPs which diminishes Ag-specific PC responses. Given that cholesterol absorption is dependent on bile acid levels, and that selected members of the microbiome can degrade bile acid, we posit that early dysbiosis will result in decreased dietary sterol absorption and consequent 25-HC levels. Since multiple antigens simultaneously drive chronic intestinal GCs, a metabolite-based mechanism to tilt the GC outcome might provide a rapid mode of Ab regulation in response to fluctuations in the luminal commensal composition.
SREBP2 has a well-established role in controlling cholesterol metabolism in non-hematopoietic cells: our findings here have established a critical role for SREBP2 in controlling GC B cell fate. While *in vitro* differentiation into PCs was inhibited in absence of SREBP2, the nature of the genetic model we used prevented us from uncovering the intrinsic role of SREBP2 in PC survival. Our *in vitro* data suggest that BCR stimulation is the dominant signal for SREBP2 transcription

and activation and it is likely to shape GC B cell fate in the LZ; however, we cannot rule out a role for T cells in maintaining SREBP2 activation and promoting differentiation in PCs.

The SREBP2 signature can be clearly observed in LZ GCs of mouse and human, suggesting an evolutionary conserved mechanism to turn on cholesterologenic genes upon selection in the LZ. Therefore, our findings regarding metabolic requirements for controlling PC differentiation may be of broader relevance to understand humoral response in human populations exposed to high cholesterol diet.

Material & Methods:

Mice: C57BL/6J (CD45.2) (Stock No: 00064), Ly5.2 (CD45.1) congenic C57BL/6 (B6) (Stock No 002014), Rosa26DTR (Stock No: 007900), Srebp2 flox/flox (Stock No: 031792), *Ch25h*$^{-/-}$ (Stock No: 016263), *Gpr183*$^{-/-}$ (**Stock No: XXX**), AID cre (Stock No: 007770), CD21^{cre} (Stock No: 006368), TdTomato (Stock No: 007914) were purchased from the Jackson Laboratory. All mice were bred and maintained under standard 12:12 hours light/dark conditions and housed in specific pathogen-free (SPF) conditions at the University of Massachusetts Medical School. All procedures were conformed to ethical principles and guidelines approved by the UMMS Institutional Animal Care and Use Committee.

Mouse Diets: Mice were either fed a standard chow diet (Envigo 2018S) for the duration of the experiment, a purified low fat diet (Envigo TD.00102 customized diet), or purified high saturated milk fat diet (0.15% HCD Envigo TD.180381customized diet).

Immunizations, infections, and treatment: For cholera toxin responses, mice were immunized with 10 ug of cholera toxin (List Biological Laboratories) in PBS by oral gavage. Animals received cholera toxin every 7 days for three consecutive weeks. Mice were analyzed 7 days after last immunization. Metabolically defective (aroA) *S. typhimurium* strain SL1344 was provided by Milena Bogunovic lab (UMMS) and was grown at 37°C in Luria broth supplemented with appropriate antibiotics to preserve mutation and plasmid. Mice were orally gavaged three times on alternate days with 10^9 CFUs of aroA S. typhimurium in 200 ul 5% sodium bicarbonate. Serial dilutions of bacterial preparations were plated onto LB-agar plates to confirm administered dose.
To prevent lymphocyte egress from Peyer's patches, the S1PR1 agonist FTY-720 (Fingolimod-HCl Cat No.S5002 from Selleck Chemicals) was dissolved in saline solution and administered to mice daily at 1mg/kg for 3-7 days as detailed in the figures.

Cell Isolation: Peyer's patches and mesenteric lymph nodes were digested in digestion media (RPMI, 5% heat inactivated FBS, 10 mM hepes, 1% penstrep, 50 ug/mL DNase I (Sigma DN25), and 0.5 mg/mL Collagenase IV (Worthington Biochemical LS004189)) rotating at 37°C for 15 min. The digested tissue was then smashed through 70µm cell strainer. Spleens were harvested and smashed through 70µm cell strainer. Isolated cells were then washed with FACS buffer (1X DPBS, 2% heat inactivated FBS, 2mM EDTA) and counted for further analysis. For isolation of lamina propria (LP) lymphocytes, small intestine was dissected and flushed with cold PBS and Peyer's Patches were removed. The small intestine was divided into 3 equal parts, the proximal (duodenum) and distal (ileum) were opened longitudinally and vortexed in a 50ml conical tube containing HBSS supplemented with 5% heat-inactivated FBS and 10mM Hepes. Epithelial cells were removed by rotating the small intestine tissue in pre-digestion media (RPMI medium, 5% heat-inactivated FBS, 10mM Hepes, 10mM EDTA) for 30 minutes at 37°C. The intestinal pieces were then washed with complete media (10% heat inactivated FBS, 10 mM hepes, 1% penstrep), chopped with scissors, and digested at 37°C for 30 minutes in digestion media. Digested tissue was passed through 70 um cell strainer and isolated cells were resuspended in 40% Percoll-RPMI and layered with 80% Percoll-RPMI and subsequently centrifuged for 20min at 2200 rpm. The isolated LP cells were enumerated on a BD LSRII using AccuCheck Counting Beads (Invitrogen) as per manufacturer recommendations.

Flow cytometry and cell sorting: Mesenteric lymph node, Peyer's patches, and spleen were collected as described above. Cell suspensions were stained with LIVE/DEAD Fixable Aqua Dead Cell Stain (Invitrogen) or Fixable Viability Dye eFluor780 (Invitrogen 65086514) in FACS buffer, and Fc receptors blocked with anti-mouse CD16/32 (2.4G2). BD Cytofix/Cytoperm kit (BD BDB554714) was used for fixation and intracellular staining. Cells were incubated for 20 min on ice with antibodies to B220, IgA, IgD, IgM, IgG1, IgG2b, IgG3, GL7, CD138, CD38, CD45.1, CD90.2, CXCR4 and CD86 (the antibodies used are described in table S1). Data were collected on a BD LSR II and analyzed in FlowJo v10.7 software.

For cell sorting, cells were stained as described above and sorted on a BD FACSAria II with a 70-micron nozzle. Cells were maintained at 4°C until sorting. Sorted plasma cells from PP and mLN were used for either Elispot assay and qPCR analysis. Germinal center, Follicular B cells, Memory and plasma cells were sorted as live IgD- GL7+ CD38- CD138- , live IgD+ GL7- CD138-, live IgD- GL7+ CD38+ CD138-, IgD- GL7- CD38- CD138+ respectively into TRIZOL (Invitrogen) for RNA extraction or into tissue culture plates containing complete media for further analysis. Light zone and dark zone germinal center B cells were sorted based on expression of CXCR4 and CD86.

RNA extraction and Real Time PCR: Total RNA was isolated with 0.5 mL of TRIZOL reagent (Invitrogen) following the manufacturer's protocol. Reverse transcription was performed using Superscript III reagent kit from Invitrogen. All samples were checked for quality (A260/A280 ratio of 1.8-2.0). Real Time PCR was performed by using iQ SYBR Green superMix (BioRad). Gene expression levels were determined using a comparative method (ΔCq) normalizing the target mRNA to b□actin as endogenous internal control. Forward and reverse primers sequences are listed in table S2.

Enzyme-linked ImmunoSorbent Assay (ELISA): Ninety-six–well high binding flat bottom plates (Corning) were coated with 25ul of 2 µg/ml purified anti-IgA (RMA-1, BD), CT (List biological laboratories) diluted in PBS overnight at 4°C. Plates were washed and blocked with PBS–5% BSA before diluted intestinal wash or fecal samples were added and threefold serial dilutions were made. Samples were processed as described previously by *Reboldi et al.* and incubated overnight at 4°C. Bound antibodies were detected by anti-IgA-conjugated horseradish peroxidase (Southern Biotech) and visualized by the addition of Substrate Reagent Pack (Biolegend). Color development was stopped with 3 M H2SO4 stop solution (Biolegend). Purified mouse IgA (Southern Biotech) served as standard. Absorbances at 450 nm were measured on a tunable microplate reader (VersaMax, Molecular Devices). Antibody titers were calculated by extrapolating absorbance values from standard curves where known concentrations were plotted against absorbance using SoftMax Pro 5 software.

ELISpot: ELISpot plates (Millipore) were coated with 100µl of 2 µg/ml purified anti-IgA in PBS overnight at 4°C. Plates were washed three times with PBS then blocked for 2 h at 37°C with 10% FCS-RPMI. Cells were isolated from Peyer's patches, mesenteric lymph nodes and lamina propria and counted as described above then diluted in blocked ELISpot plates and incubated overnight in a 37°C 5% CO_2 tissue culture incubator. The next day, plates were washed three times with PBS-0.1% Tween then PBS. Detection of IgA spots was achieved by using anti-IgA-conjugated horseradish peroxidase and developed with 3-amino-9-ethylcarbazole (Sigma-Aldrich). Color development was stopped by washing several times with water. Once dried, plates were scanned and spots counted using the CTL ELISPOT reader system (Cellular Technology).

NB-21 coculture: Germinal center B cells from Peyer's patches and mesenteric lymph nodes were FACS sorted and cocultured with NB-21 feeder cells as previously described and with adjustments (Kuraoka et al., 2016; Stewart 2018). Briefly, 10^3 NB-21.2D9 cells/well in 100 uL of B cell media (RPMI supplemented with 10% heat-inactivated FBS, 10 mM HEPES, 1mM sodium pyruvate, 100 units/mL penicillin, 100 mg/mL streptomycin, MEM nonessential amino acid and 55 mM 2-Mercaptoethanol) were seeded into 96-well plates 1 day before B cell co-culturing. On the next day, 10^4 B cells in 100 uL of B cell media was added to each well. 100uL of media was replaced on day 2. Cholesterol, 7a,25-HC or 25-HC were added to the B cell/NB-21 co-culture at days 0 and 2. Culture plates were centrifuged at day 3.5 and supernatant was collected for IgA detection.

Nuclei Preparation: Nuclei preparations were carried out as previously described (Gallagher, M. P Immunohorizons, 2018). Briefly, follicular B cells were isolated from spleens of wild type or *Aicda*$^{cre/+}$ *Srebf2*$^{fl/fl}$ mice using EasySep mouse B cell isolation kit and protocol (StemCell). Isolated cells were then cultured *in vitro* in the presence or absence of 5 mg/ml anti-IgM $F(ab)_2$ (Jackson Immunoresearch) and/or 5 mg/ml anti-CD40 (BioXCell) for 24 or 48 hours. Cells were harvested and lysed in Sucrose Buffer A (10 mM HEPES (Gibco), 8 mM $MgCl_2$, 320 mM Sucrose, 0.1 % (v/v) Triton-X 100 (Sigma), and 1X cOmplete, EDTA-free Protease Inhibitor Cocktail (Roche) on ice for 10 minutes. The cells were then centrifuged at 2000g for 10 minutes and washed twice with Sucrose Buffer B (Sucrose Buffer A without Triton-X 100). Nuclei were fixed in 4% Paraformaldehyde (electron microscopy grade, Electron Microscopy Sciences), diluted in Sucrose Buffer B, at room temperature for 25 minutes. Nuclei were stained as described in figure 4 and figureS4 and analyzed by flow cytometry.

Bone marrow chimeras: CD21$^{cre/Rosa26DTR+}$or CD21^{cre} mice were lethally irradiated twice with 550 rads gamma-irradiation, 3 hours apart, then intravenously injected with 1-3 x 10^6 bone marrow cells. Bone marrow was harvested by flushing both tibia and femurs of donor mice. Mice were analyzed 8-12 weeks later.

FDC ablation: For LT-blocking experiments, animals were treated with LTβR-Fc using 100 µg on days −3 and −1 before sacking the mice. FDC ablation with DT was performed as described previously (Wang et al., 2011). In brief, animals expressing CD21-cre and ROSA-DTR alleles or littermate controls were lethally irradiated and reconstituted with WT BM. 6–8 wk after reconstitution, animals were treated with 100 ng DTx (EMD Biosciences) i.p. and analyzed 16–20 h later.

Lipid extraction and 25HC/7a,25-HC quantification: Lipids from 100 mg of Peyer's patches and mesenteric lymph nodes were extracted using the Folch method for lipid extraction (Folch et al., 1957). Lyophilized lipid was dissolved at 100 mg/ml in ethanol. 25-HC was measured by using 293T cells supernatant from cells transfected sequentially with *pENTR-Hsd3b7* and *pENTR-Cyp7b1* vectors. The 7a,25-HC activity was evaluated by transwell chemotaxis assay. Lipid extracts were diluted in 10 volumes of sterile chemotaxis media (RPMI + 0.5% fatty acid free BSA) and tested for EBI2 dependent bioactivity by transwell chemotaxis assays of 50:50 mixed M12 B cell line transduced with an EBI2-IRES-GFP retroviral construct and mock M12 cells. The three-hour migration assay was performed at 37°C and migrated cells were stained for DAPI and analyzed by flow cytometry. The relative migration of EBI2-GFP+ M12 cells over M12 cells was normalized to migration toward lipid free migration media. The purified 7a,25-HC was used as positive control for chemotaxis.

Statistical analysis and quantification: Statistical analyses were performed using GraphPad Prism v8.0 using two-way ANOVA with Bonferroni's multiple comparisons test or unpaired T test Data are presented as means +/- SEM. Differences between group means were considered significant at indicated p value. Sample sizes can be found in the figure legends, indicating the number of animals or cells used for each experiment.

References:

1. van de Pavert, S. A. *et al.* Maternal retinoids control type 3 innate lymphoid cells and set the offspring immunity. *Nature* 1–17 (2014). doi:10.1038/nature13158
2. Mora, J. R. *et al.* Generation of Gut-Homing IgA-Secreting B Cells by Intestinal Dendritic Cells. *Science* **314,** 1157–1160 (2006).
3. Cyster, J. G., Dang, E. V., Reboldi, A. & Yi, T. 25-Hydroxycholesterols in innate and adaptive immunity. *Nat Rev Immunol* **14,** 731–743 (2014).
4. Petta, I., Fraussen, J., Somers, V. & Kleinewietfeld, M. Interrelation of Diet, Gut Microbiome, and Autoantibody Production. *Front. Immun.* **9,** 642–9 (2018).
5. Pereira, J. P., Kelly, L. M., Xu, Y. & Cyster, J. G. EBI2 mediates B cell segregation between the outer and centre follicle. *Nature* **460,** 1122–1126 (2009).
6. Kelly, L. M., Pereira, J. P., Yi, T., Xu, Y. & Cyster, J. G. EBI2 Guides Serial Movements of Activated B Cells and Ligand Activity Is Detectable in Lymphoid and Nonlymphoid Tissues. *J Immunol* **187,** 3026–3032 (2011).
7. Yi, T. *et al.* Oxysterol Gradient Generation by Lymphoid Stromal Cells Guides Activated B Cell Movement during Humoral Responses. *Immunity* **37,** 535–548 (2012).
8. Baptista, A. P. *et al.* The Chemoattractant Receptor Ebi2 Drives Intranodal Naive CD4+ T Cell Peripheralization to Promote Effective Adaptive Immunity. *Immunity* **50,** 1188–1201.e6 (2019).
9. Yi, T. & Cyster, J. G. EBI2-mediated bridging channel positioning supports splenic dendritic cell homeostasis and particulate antigen capture. *eLife* **2,** 309–19 (2013).
10. Chu, C. *et al.* Anti-microbial Functions of Group 3 Innate Lymphoid Cells in Gut-Associated Lymphoid Tissues Are Regulated by G-Protein-Coupled Receptor 183. *CellReports* **23,** 3750–3758 (2018).
11. Emgård, J. *et al.* Oxysterol Sensing through the Receptor GPR183 Promotes the Lymphoid-Tissue-Inducing Function of Innate Lymphoid Cells and Colonic Inflammation. *Immunity* **48,** 120–132.e8 (2018).
12. Melo-Gonzalez, F. *et al.* Antigen-presenting ILC3 regulate T cell–dependent IgA responses to colonic mucosal bacteria. *Journal of Experimental Medicine* **216,** 728–742 (2019).
13. Reboldi, A. *et al.* 25-Hydroxycholesterol suppresses interleukin-1–driven inflammation downstream of type I interferon. *Science* **345,** 679–684 (2014).
14. Dang, E. V., McDonald, J. G., Russell, D. W. & Cyster, J. G. Oxysterol Restraint of Cholesterol Synthesis Prevents AIM2 Inflammasome Activation. *Cell* 1–27 (2017). doi:10.1016/j.cell.2017.09.029

15. Gold, E. S. *et al.* 25-Hydroxycholesterol acts as an amplifier of inflammatory signaling. *Proceedings of the National Academy of Sciences* **111,** 10666–10671 (2014).
16. Liu, S.-Y. *et al.* Interferon-Inducible Cholesterol-25-Hydroxylase Broadly Inhibits Viral Entry by Production of 25-Hydroxycholesterol. *Immunity* 1–14 (2012). doi:10.1016/j.immuni.2012.11.005
17. Blanc, M. *et al.* The Transcription Factor STAT-1 Couples Macrophage Synthesis of 25-Hydroxycholesterol to the Interferon Antiviral Response. *Immunity* 1–13 (2012). doi:10.1016/j.immuni.2012.11.004
18. Goldstein, J. L., DeBose-Boyd, R. A. & Brown, M. S. Protein Sensors for Membrane Sterols. *Cell* **124,** 35–46 (2006).
19. Sakai, J. *et al.* Sterol-regulated release of SREBP-2 from cell membranes requires two sequential cleavages, one within a transmembrane segment. *Cell* **85,** 1037–1046 (1996).
20. Kidani, Y. *et al.* Sterol regulatory element–binding proteins are essential for the metabolic programming of effector T cells and adaptive immunity. *Nat Immunol* **14,** 489–499 (2013).
21. Jung, C., Hugot, J.-P. & Barreau, F. Peyer's Patches: The Immune Sensors of the Intestine. *International Journal of Inflammation* **2010,** 1–12 (2010).
22. Reboldi, A. *et al.* IgA production requires B cell interaction with subepithelial dendritic cells in Peyers patches. *Science* **352,** aaf4822–aaf4822 (2016).
23. Belkaid, Y. & Naik, S. Compartmentalized and systemic control of tissue immunity by commensals. *Nat Immunol* **14,** 646–653 (2013).
24. Hooper, L. V. & Macpherson, A. J. Immune adaptations that maintain homeostasis with the intestinal microbiota. *Nat Rev Immunol* **10,** 159–169 (2010).
25. Martinoli, C., Chiavelli, A. & Rescigno, M. Entry Route of Salmonella typhimurium Directs the Type of Induced Immune Response. *Immunity* **27,** 975–984 (2007).
26. Monack, D. M., Bouley, D. M. & Falkow, S. Salmonella typhimurium Persists within Macrophages in the Mesenteric Lymph Nodes of Chronically Infected Nramp1+/+ Mice and Can Be Reactivated by IFNγ Neutralization. *Journal of Experimental Medicine* **199,** 231–241 (2004).
27. Gohda, M. *et al.* Sphingosine 1-phosphate regulates the egress of IgA plasmablasts from Peyer's patches for intestinal IgA responses. *J Immunol* **180,** 5335–5343 (2008).
28. Bauman, D. R. *et al.* 25-Hydroxycholesterol secreted by macrophages in response to Toll-like receptor activation suppresses immunoglobulin A production. *Proc Natl Acad Sci U S A* **106,** 16764–16769 (2009).
29. Kuraoka, M. *et al.* Complex Antigens Drive Permissive Clonal Selection in Germinal Centers. *Immunity* 1–30 (2016). doi:10.1016/j.immuni.2016.02.010
30. Wang, X. *et al.* Follicular dendritic cells help establish follicle identity and promote B cell retention in germinal centers. *Journal of Experimental Medicine* **208,** 2497–2510 (2011).
31. Iwata, M. *et al.* Retinoic Acid Imprints Gut-Homing Specificity on T Cells. *Immunity* **21,** 527–538 (2004).
32. Rodda, L. B. *et al.* Single-Cell RNA Sequencing of Lymph Node Stromal Cells Reveals Niche-Associated Heterogeneity. *Immunity* **48,** 1014–1028.e6 (2018).
33. Gallagher, M. P., Conley, J. M. & Berg, L. J. Peptide Antigen Concentration Modulates Digital NFAT1 Activation in Primary Mouse Naive CD8 +T Cells as Measured by Flow Cytometry of Isolated Cell Nuclei. *IH* **2,** 208–215 (2018).

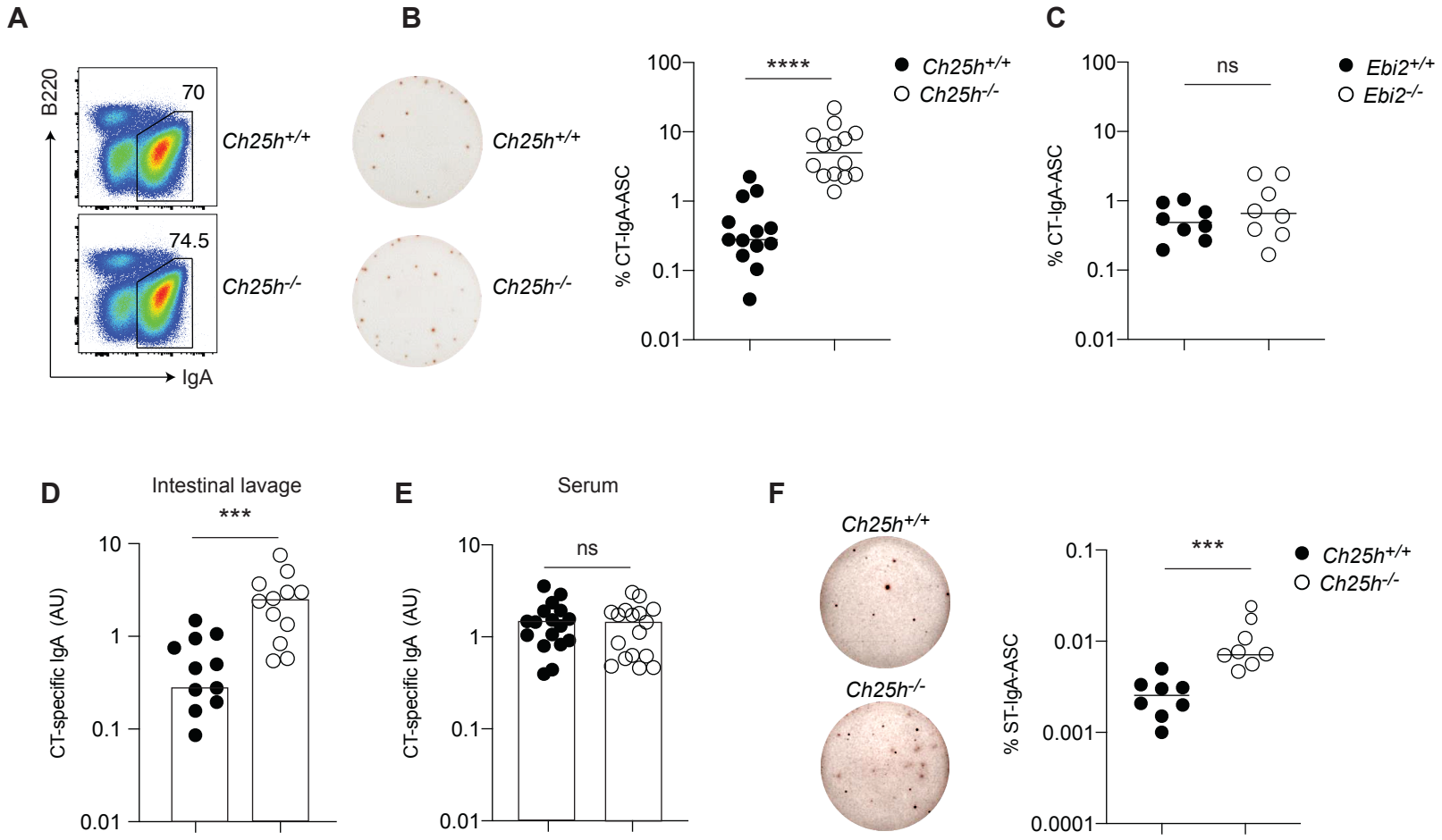
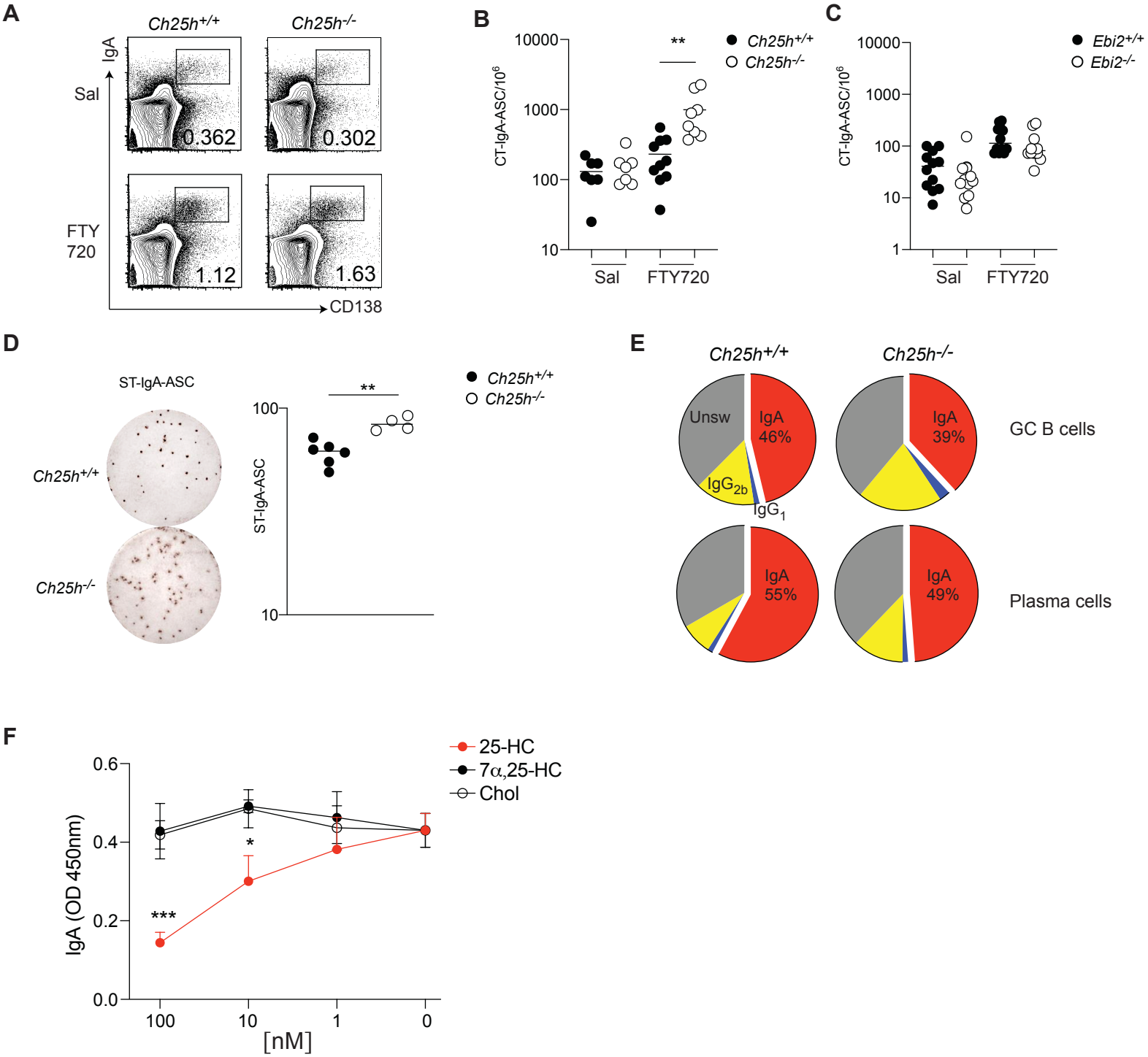

A
B220
70
74.5
Ch25h+/+
Ch25h-/-
IgA
B
Ch25h+/+
Ch25h-/-
****
% CT-IgA-ASC
100
10
1
0.1
0.01
C
ns
Ebi2+/+
Ebi2-/-
% CT-IgA-ASC
D
Intestinal lavage
***
CT-specific IgA (AU)
E
Serum
ns
CT-specific IgA (AU)
F
Ch25h+/+
Ch25h-/-
***
% ST-IgA-ASC
0.1
0.01
0.001
0.0001

Figure 1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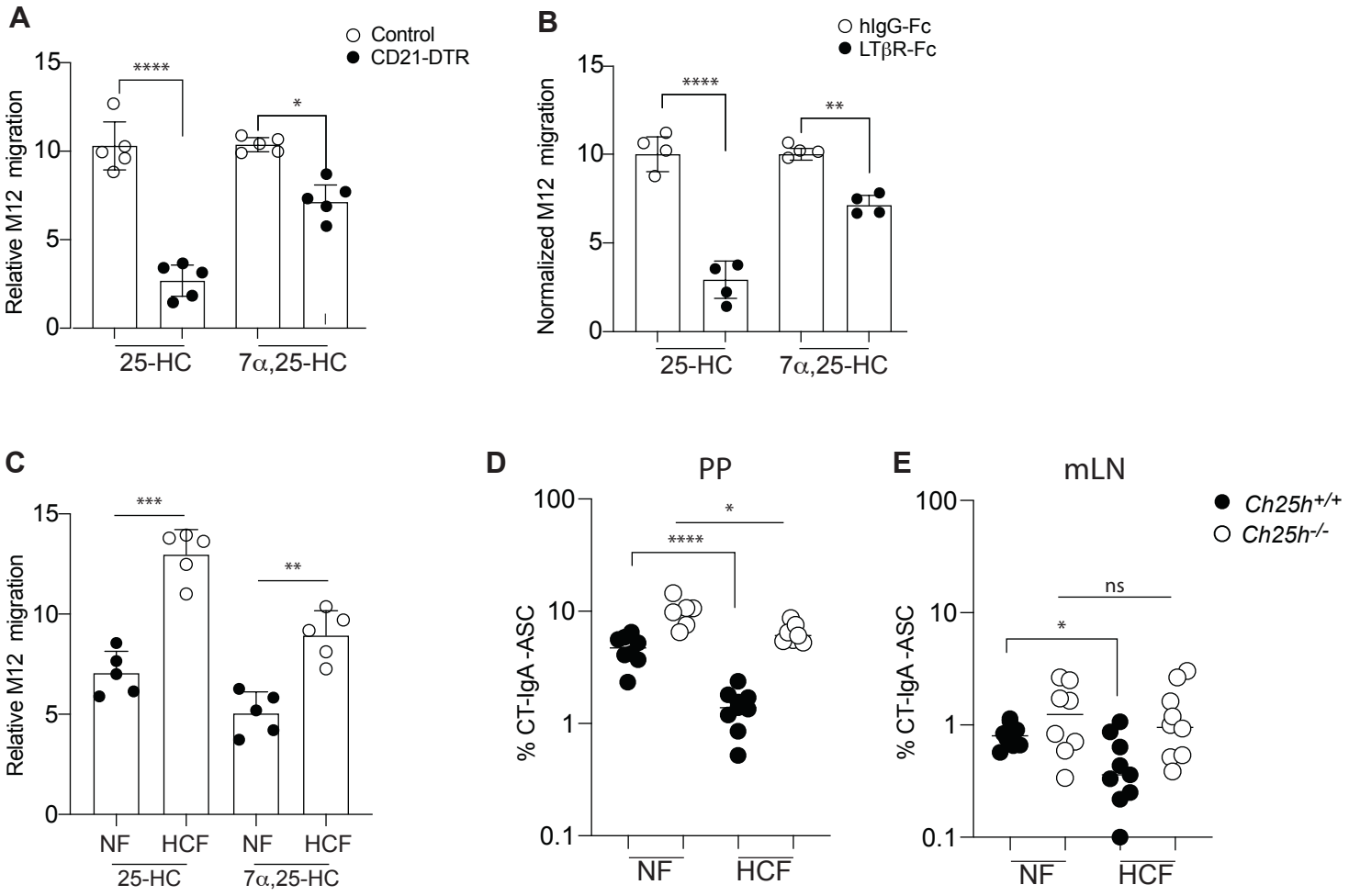
Ch25h+/+
Ch25h-/-
IgA
Sal
FTY 720
0.362
0.302
1.12
1.63
CD138
B
CT-IgA-ASC/10^6
10000
1000
100
10
**
Ch25h+/+
Ch25h-/-
Sal
FTY720
C
Ebi2+/+
Ebi2-/-
1
D
ST-IgA-ASC
Ch25h+/+
Ch25h-/-
100
10
E
Unsw
IgA 46%
IgG2b
IgG1
IgA 39%
GC B cells
IgA 55%
IgA 49%
Plasma cells
F
25-HC
7α,25-HC
Chol
IgA (OD 450nm)
0.6
0.4
0.2
0.0
***
*
100
10
1
0
[nM]

Figure 2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Control
CD21-DTR
Relative M12 migration
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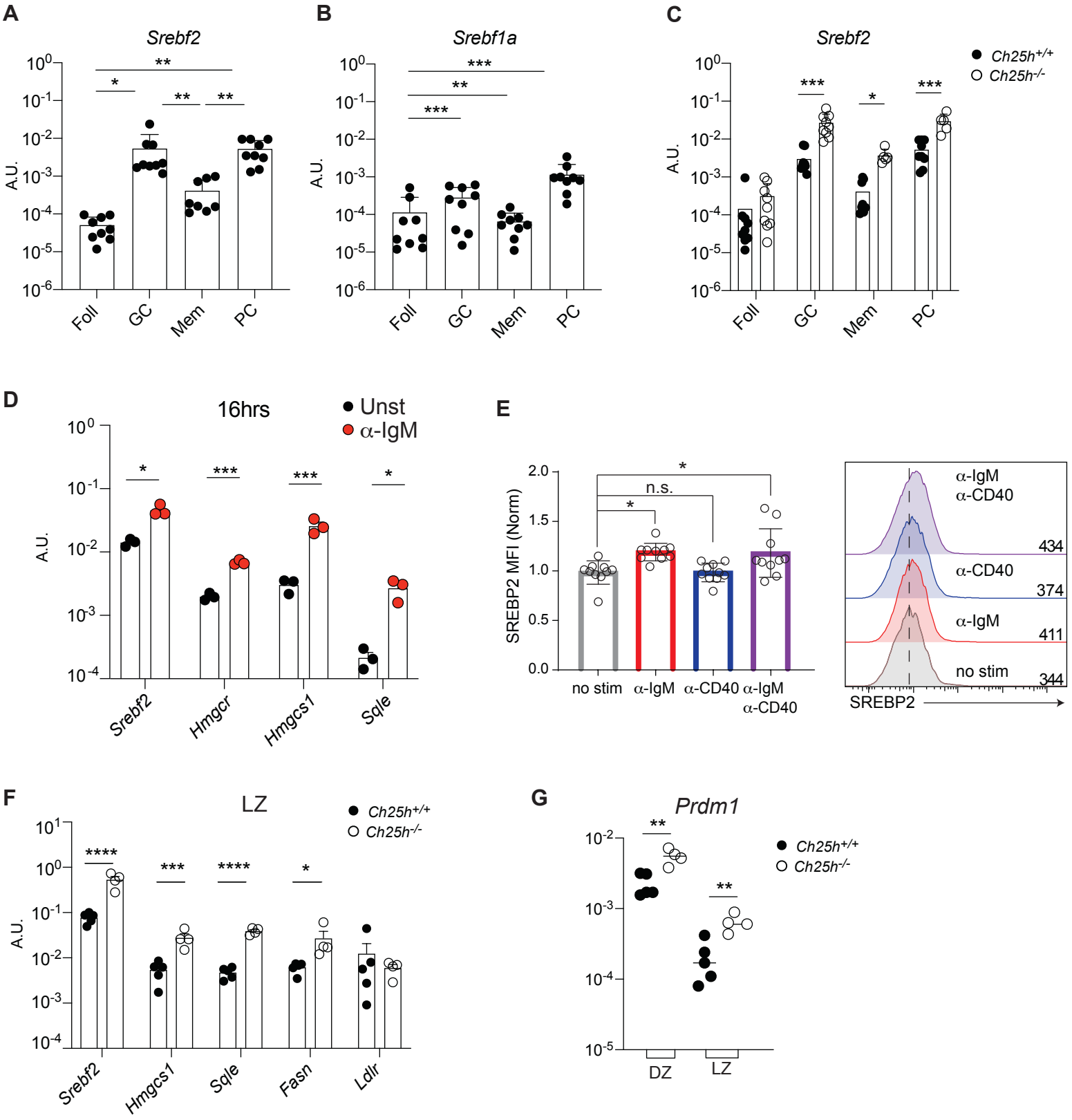
10
5
0
****
*
25-HC
7α,25-HC
B
hIgG-Fc
LTβR-Fc
Normalized M12 migration
15
10
5
0
****
**
25-HC
7α,25-HC
C
Relative M12 migration
15
10
5
0
***
**
NF
HCF
NF
HCF
25-HC
7α,25-HC
D
PP
100
10
1
0.1
% CT-IgA -ASC
*
****
NF
HCF
E
mLN
100
10
1
0.1
% CT-IgA -ASC
ns
*
NF
HCF
Ch25h+/+
Ch25h-/-

Figure 3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Foll
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Mem
PC
B
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C
Srebf2
Ch25h+/+
Ch25h-/-
D
16hrs
Unst
α-IgM
Srebf2
Hmgcr
Hmgcs1
Sqle
E
SREBP2 MFI (Norm)
n.s.
no stim
α-IgM
α-CD40
α-IgM
α-CD40
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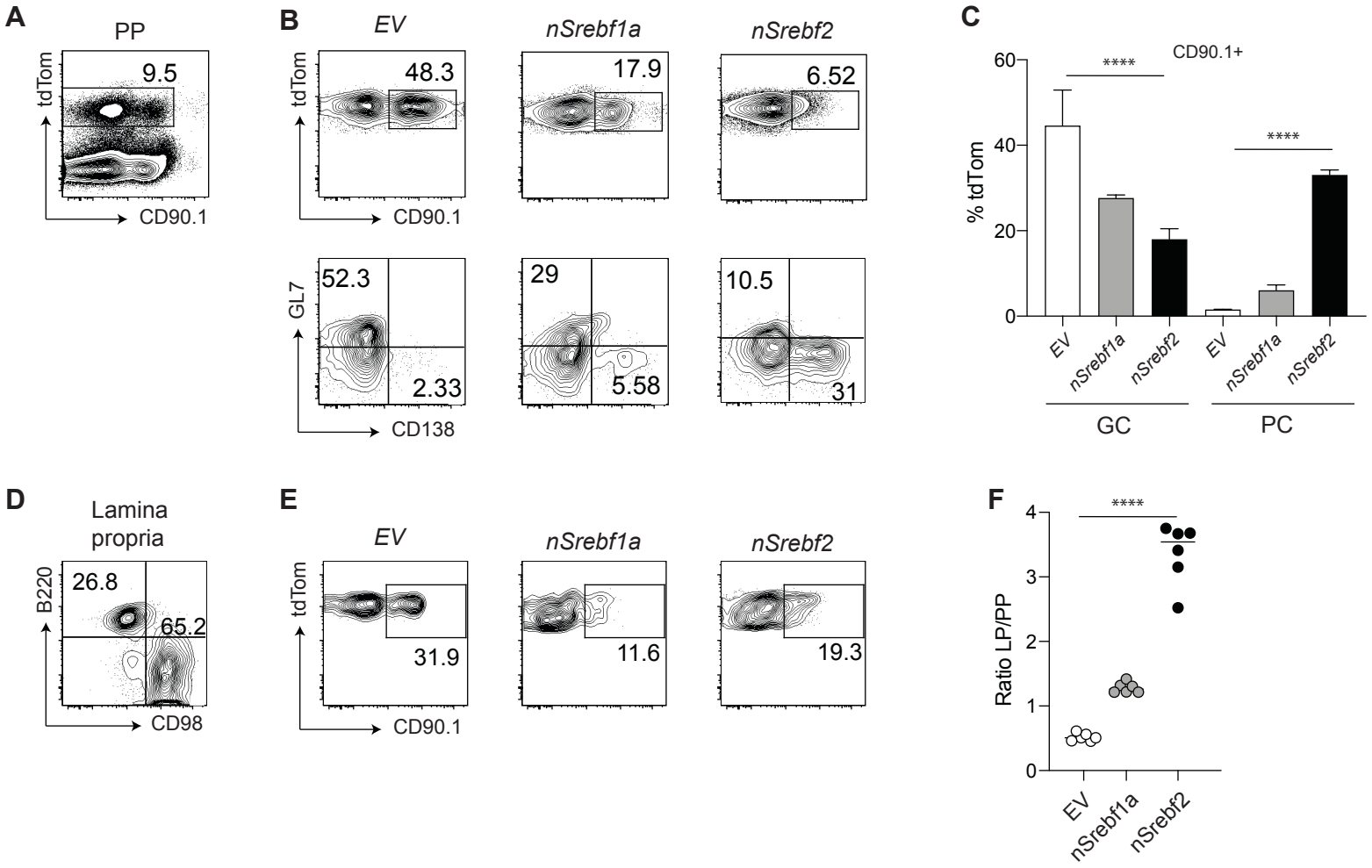
411
344
SREBP2
F
LZ
Fasn
Ldlr
G
Prdm1
DZ
LZ

Figure 4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B
EV
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nSrebf2
tdTom
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17.9
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CD90.1
GL7
52.3
2.33
29
5.58
10.5
31
CD138
C
CD90.1+
% tdTom
60
40
20
0
****
****
EV
nSrebf1a
nSrebf2
EV
nSrebf1a
nSrebf2
GC
PC
D
Lamina propria
B220
26.8
65.2
CD98
E
EV
nSrebf1a
nSrebf2
tdTom
31.9
11.6
19.3
CD90.1
F
****
Ratio LP/PP
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3
2
1
0
EV
nSrebf1a
nSrebf2

Figure 5

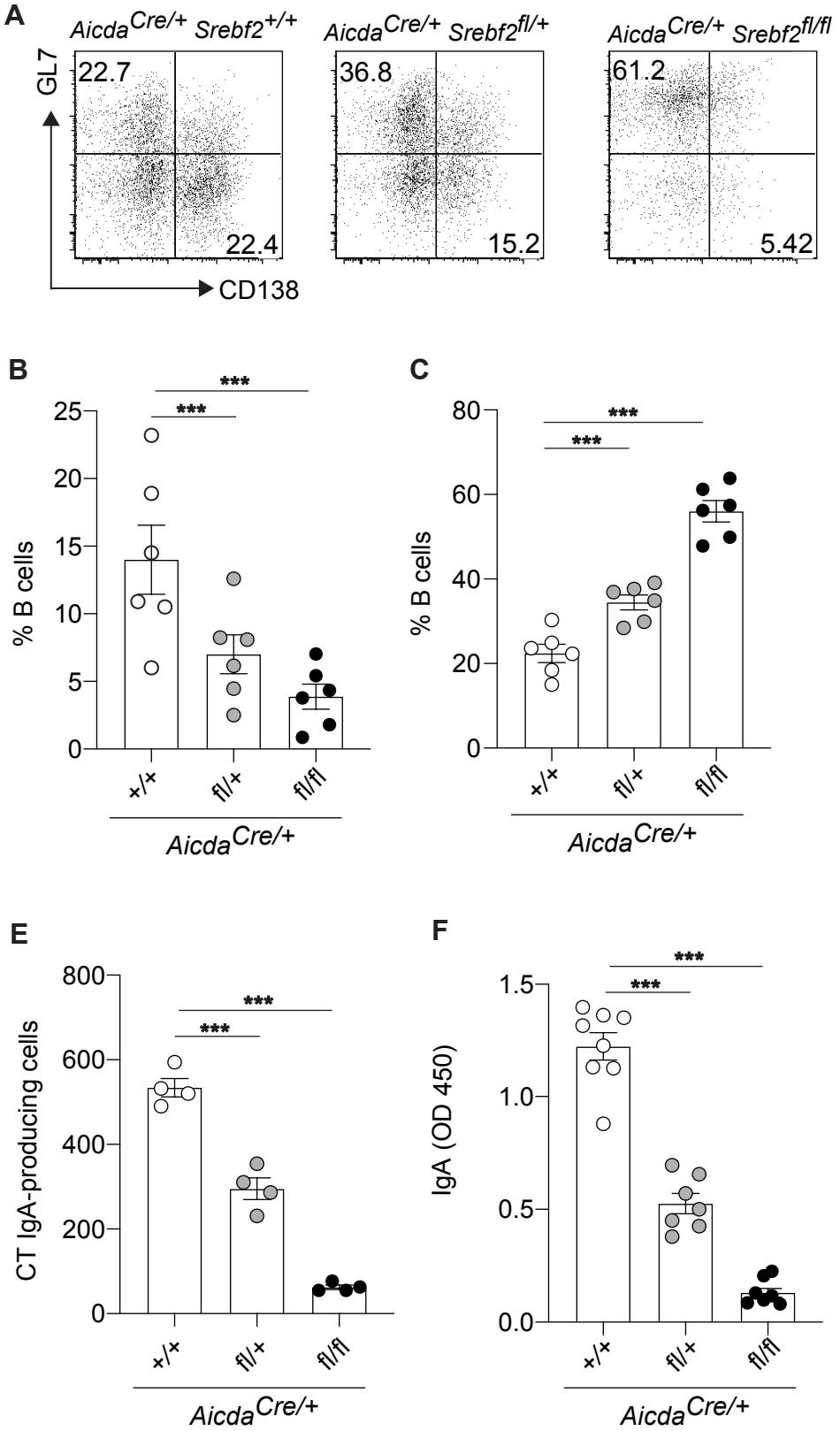

A
Aicda Cre/+ Srebf2 +/+
Aicda Cre/+ Srebf2 fl/+
Aicda Cre/+ Srebf2 fl/fl
GL7
CD138
22.7
22.4
36.8
15.2
61.2
5.42
B
% B cells
C
% B cells
D
IgA (O.D. 450 nm)
E
CT IgA-producing cells
F
IgA (OD 450)
***
+/+
fl/+
fl/fl
Aicda Cre/+

Figure 6

FIGURE LEGENDS

Figure 1. 25-HC, but not EBI2, shapes antigen-specific IgA response in duodenal LP.
(A) Representative Flow cytometric analysis of IgA+ plasma cells in Duodenum (LP) of *Ch25h-/-* and LMC mice. (B) ELISPOT of isolated lamina propria PC from *Ch25h*$^{-/-}$ and (C) *Ebi2*$^{-/-}$ with quantified percentage of CT-specific IgA secreting PC. (D,E) Quantification of secreted CT-IgA by ELISA assay in small intestine lavage and serum of mice immunized for 3 weeks with CT. (F) Percentage of Salmonella T.-specific IgA PC in intestinal lamina propria of *Ch25h-/-* and LMC mice, analyzed by ELISPOT assay two weeks post-infection. Data are representative of three independent experiment, ns=non-significant, *** p<0.005;****p<0.001 (unpaired Student's T test).

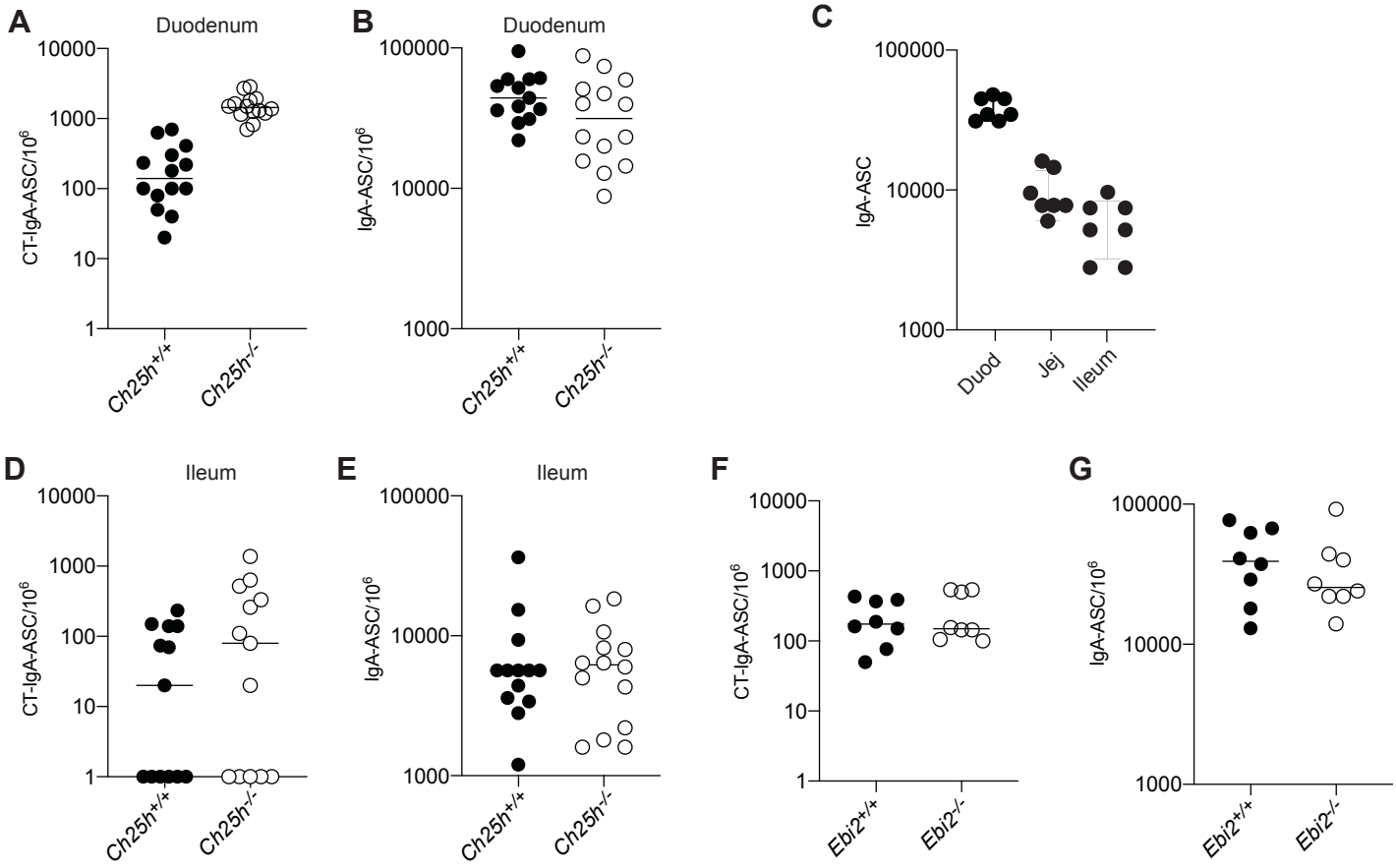
Figure 2. 25-HC modulated Ag-specific plasma cells generation in PPs
(A)Flow cytometric analysis of plasma cells in PP of *Ch25h-/-* and LMC mice treated daily with FTY720 or saline (Sal) for one week. (B) Number of CT-specific IgA secreting cells in *Ch25h*$^{-/-}$. (C) *Ebi2*$^{-/-}$ and LMC mice, measured by ELISPOT. (D) Representative ELISPOT showing Salmonella T. specific IgA secreting cells in PP of *Ch25h*$^{-/-}$ and LMC mice treated with FTY720. Data indicate the frequency of ST-ASC among lamina propria cells. (E) Analysis of Class switch recombination in PP by Flow cytometry. (F) Quantification of secreted IgA by ELISA from GC B cells cultured with NB21 cells and incubated with the indicated sterols for 3.5 days. A,B,C,E from at least three independent experiments, D, F from two independent experiments. Statistical significance was measured by two-way ANOVA using Bonferroni's corrections in B, C, F **p<0.001 (two-way ANOVA using Bonferroni's corrections (B, C, F) unpaired Student's t test (D))

Figure 3. FDCs and diet shape tissue 25-HC levels and control IgA response
Relative migration of EBI2+ M12 in a transwell assay measured by flow cytometry in response to lipid extracted from PP of (A) CD21^{DTR+} and LMC mice treated with DT or PBS and (B) wild type mice treated with recombinant LTβR-FC or isotype control. Relative migration was measured comparing migrated EBI2^{+} M12 cell to M12 cells (EBI2-). (C) Relative migration of EBI2^{+} M12 cells exposed to PP lipid extracts of mice treated for 1 weeks with 0.15 % High cholesterol diet or normal chow. (D,E) Frequency of CT-specific ASC detected by ELISPOT of mLN and PP suspension cells from *Ch25h*$^{-/-}$ and LMC fed with mice normal food or 0.15% of HCF. Results are pooled from 3 or 4 independent experiments. ns=non-significant, *p<0.005, **p<0.001,***p<0.0005, ****p<0.0001 (two-way ANOVA)

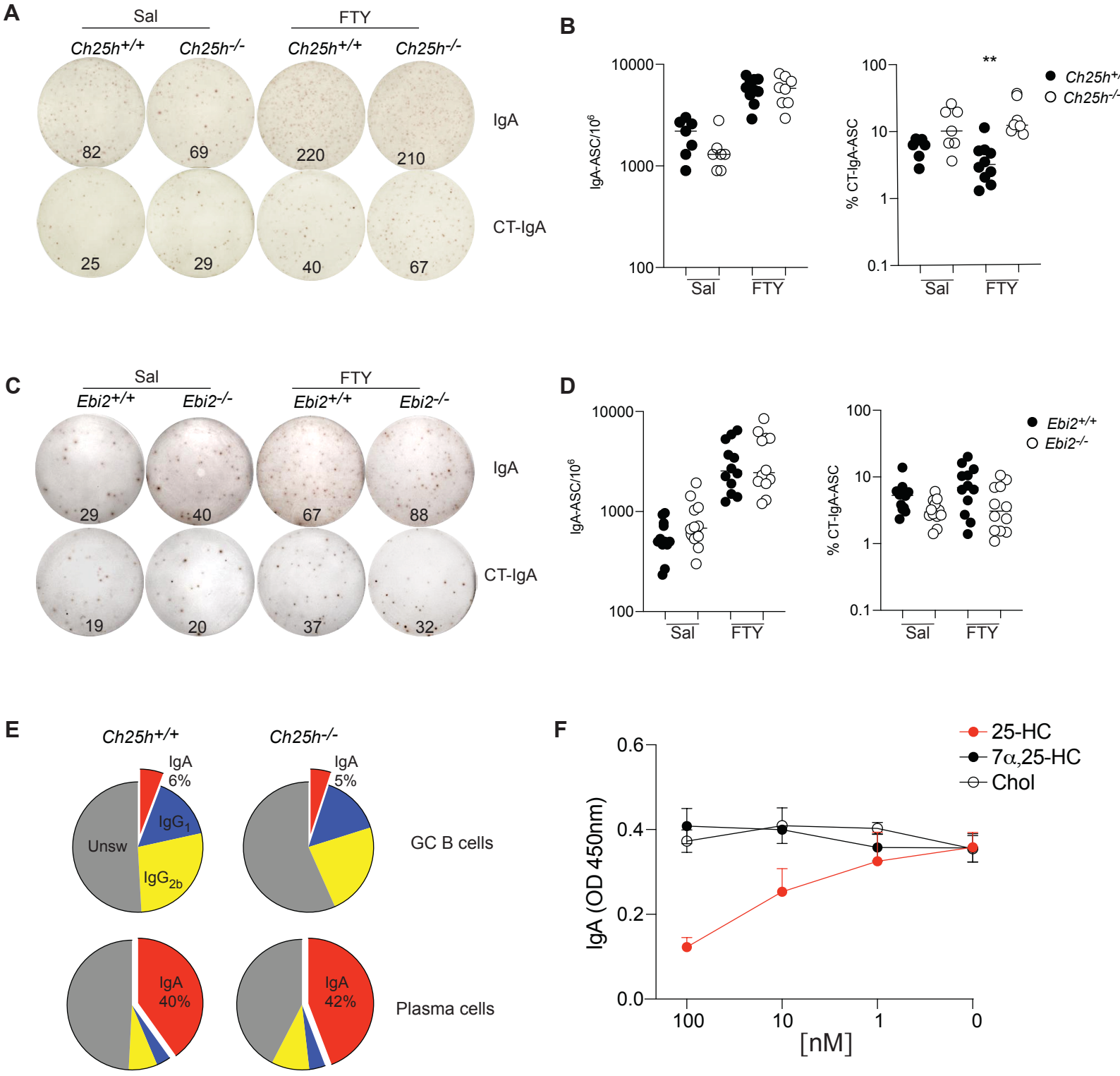
Figure 4. 25-HC regulates SREBP2 transcription and activation in activated B cells
(A,B) mRNA level of *Srebf2* (A) and *Srebf1a* (B) was quantified by Real Time PCR. B cell subsets were isolated from PP of *Ch25h-/-* and LMC mice (C) and further separated by single cell sorting for RNA extraction. (D) Analysis of gene expression of Srebp2 and Srebp2-target in Follicular B cells upon 16 hrs of BCR stimulation. (E) Cumulative and representative flow cytometry of SREBP2 nuclear staining in Follicular B cells stimulated with the indicated stimuli F) qPCR results of *Srebf2* and its target genes in sorted B cells of germinal center LZ. (G) Blimp expression in sorted DZ (CXCR4+CD86-) and LZ (CXCR4-CD86+) GC B cells from PP of *Ch25h-/-* and LMC mice. Each symbol represents one independent mouse from 9-10 (in A,B,C, E) or 4 mice (in F,G). Graphs show mean and SEM. *p<0.005, **p<0.001,***p<0.0005, ****p<0.0001(two-way ANOVA)

Figure 5. SREBP2 overexpression induces PC differentiation.
(A) Representative plot of *Aicda*$^{Cre/+}$ *Rosa26*$^{tdTom/+}$ BM transduced with CD90.1 retroviral vector in PPs.
(B) Representative plot of frequency and fate of tdTom+ B cells transduced with indicated vectors (C) Summary of the data in (B) (D) Representative gating of tdTom+ cells in lamina propria. (E) Representative plot of frequency and fate of tdTom+ B cells transduced with indicated vectors. (F) Ratio of transduced B cells in PP and LP from data in (B) and (E). Data represents 3 independent experiments. Graphs show mean and SEM. ****p<0.0001(two-way ANOVA).

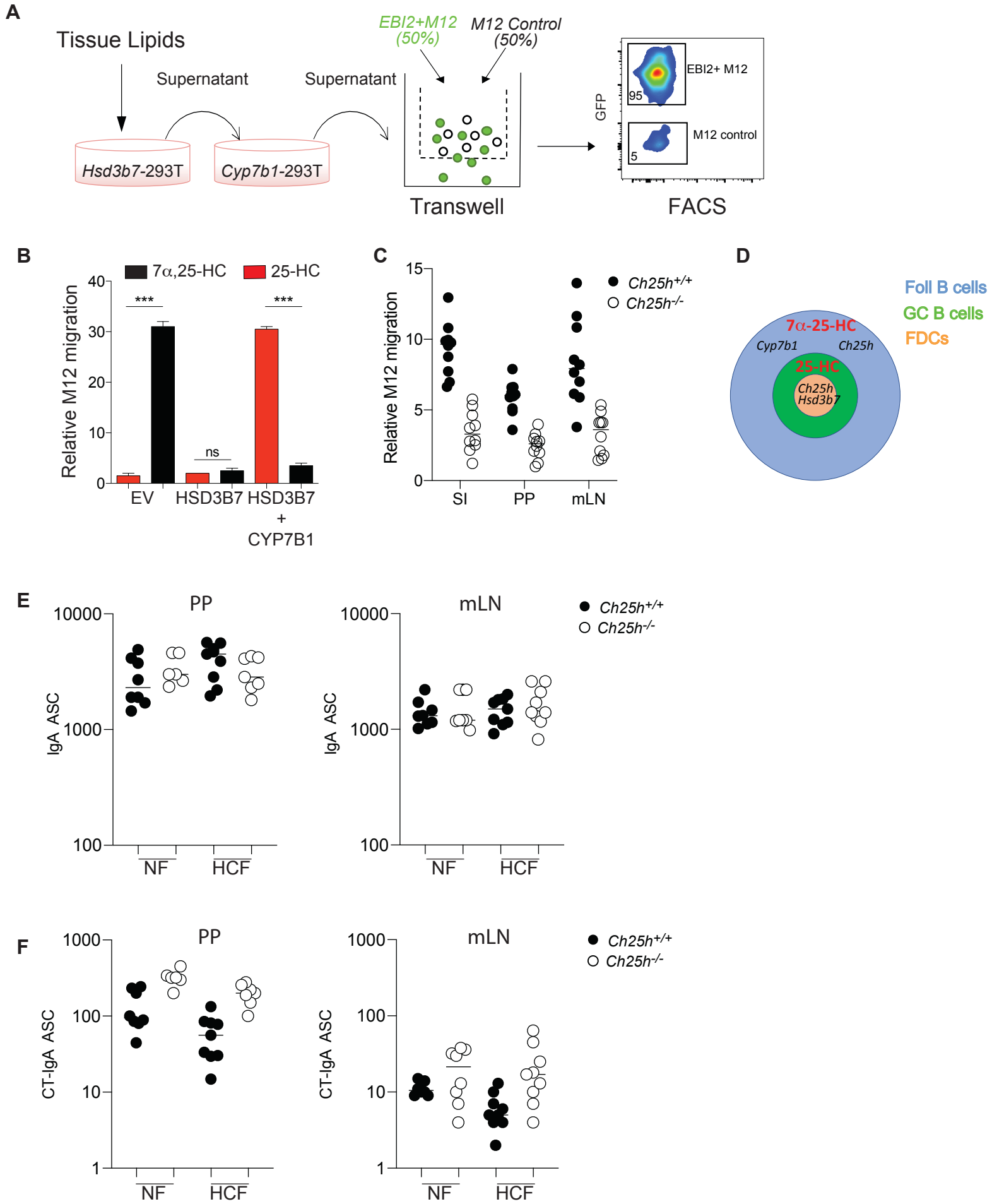
Figure6: SREBP2 control GC differentiation potential *in vitro* and *in vivo.*
(A)Representative flow cytometry of GC B cells from mice of the indicated genotype cultured with NB21 cells for 2.5 days. (B,C) Frequency of GC B cells and PCs from A. (D) Quantification of secreted total IgA in A. (E) Number of CT-IgA ASC cells in LP of AidcreSREBP2F/F and littermate controls, quantified by ELISPOT. (F) Quantification of CT-IgA in small intestine lavage of mice treated 3 weeks with CT. Each symbol represents one mouse from 2-3 independent experiments. ***p<0.0005 (two-way ANOVA).



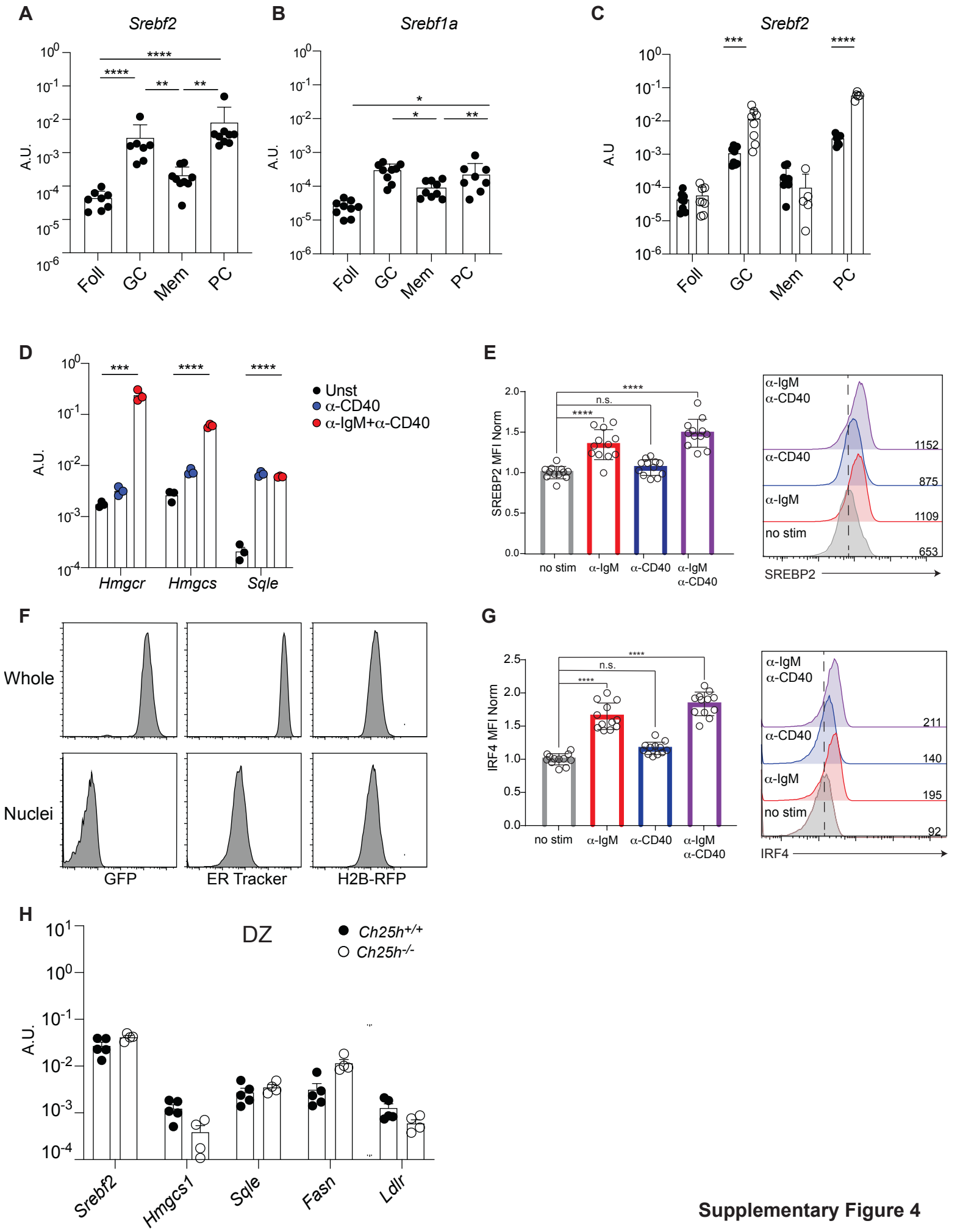
Supplementary Figure 1



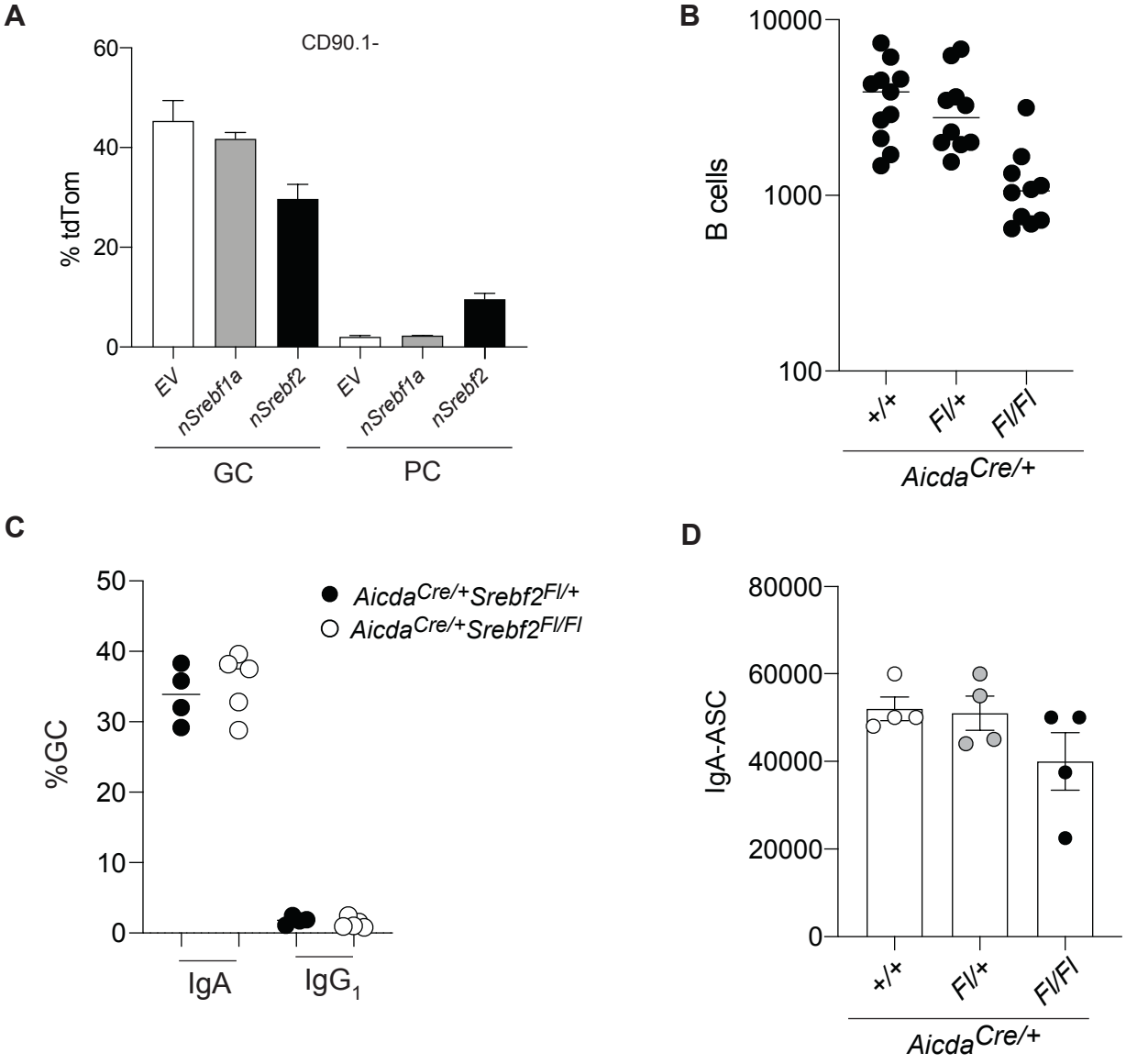
Supplementary Figure 2



Supplementary Figure 3



Supplementary Figure 4



Supplementary Figure 5

SUPPLEMENTARY FIGURE LEGENDS

Figure S1. 25-HC controls antigen-specific IgA secretion in duodenum, but not in the ileum.
(A) ELISPOT of CT-specific IgA and (B) total IgA secretion in Small intestine LP. (C) Number of secreting IgA PC in Duodenum, Jejunum and Ileum, analyzed by ELISPOT. (D) IgA ASC and (E) total IgA in the ileum of *Ch25h-/-* and LMC mice, depicted by ELISPOT. (F,G) ELISPOT data showing IgA ASC and total IgA in EBI2-/- and littermate control mice. Graphs show data from 3≥ experiments. *p<0.005, **p<0.001,***p<0.0005 (ANOVA multiple comparation).

Figure S2. 25-HC modulate CT-specific IgA response in PPs independently of class switch.
(A) Image of representative ELISPOT of PP IgA and CT-specific IgA secreting cells from *Ch25h-/-* and LMC mice orally immunized with CT for 3 weeks and treated for one week with FTY720 or saline by IP.(B) Graphs showing the plotted analysis of the total number of IgA secreting PC and the frequency of the CT-specific IgA PC, detected by ELISPOT assay.(C) Representative ELISPOT of PP IgA and CT-specific IgA secreting cells from $EBI2^{+/+}$ and $EBI2^{-/-}$ mice treated for 3 weeks with CT by oral gavage and one week with FTY720 or saline by IP. (D) Left graph showing the total number of IgA secreting plasma cells in PP of $EBI2^{+/+}$ and $EBI2^{-/-}$ mice. (E) Class switch recombination in GC B cells and PC from mLN of the indicated genotype assessed by FACS. (F) Quantification of secreted IgA by ELISA from GC B cells cultured with NB21 supernatant and anti-CD40 and stimulated with the indicated sterols for 3.5 days.
Graphs in B,D and F depict the mean from 2-3 independent experiments *p<0.005, **p<0.001,***p<0.0005 (two-way ANOVA)

Figure S3. 25HC quantification in PP GC and antigen-specific IgA
(A) Schematic illustration of transwell migration assay with EBI2+ M12 for the quantification of 25HC in intestinal SLO using transfected 293T cell. (B) Validation of the strategy in A using synthetic 25-HC and 7α,25-HC. Graph shows the relative migration of $M12^{EBI2+}$ cells from 3 experiments. (C) 25-HC quantification in small intestine, PP and mLN from *Ch25h-/-* and LMC mice. (D) Model for 25-HC rich niche in GC. (E) ELISPOT of IgA and (F) CT-specific secreting cells on PP and mLN of mice treated for 1 week with regular chow or 0.15% cholesterol supplement diet. ELISPOT data were pooled from at least 3 independent experiment *** p<0.005 (unpaired Student's t test (B) and two-way ANOVA (D, E)).

Figure S4. 25-HC reduces SREBP2 and alters SREBP2-mediated transcription in vitro and in vivo
(A)Quantitative gene expression of *Srebf2* and *(B) Srebf1a* in wild type mLN. (C) Srebf2 mRNA quantification on sorted mLN B cell subsets from Ch25h-/- and LMC. (D) qPCR of mevalonate pathway and SREBP2 targets on B cells stimulated for 48hrs with αCD40 or with αCD40 and αIgM. (E) Mean Fluorescence index (MFI) and representative histograms of total SREBP2 detected by flow cytometry in B cells stimulated for 16h with the indicated stimuli. (F) Whole cell and nuclei isolation stain of transduced WEHI for presence of cytoplasmic, ER and nuclear marker. (G) Mean Fluorescence index (MFI) of intranuclear IRF4 detected by flow cytometry in B cells stimulated for 16h with the indicated stimuli.(H) Gene expression of *Srebf2* and SREBP2- target genes in sorted DZ B cells (gated on CXCR4+ CD86- cells) from mLN of $Ch25h^{-/-}$ and LMC mice. Data were produced from at least experiment. *p<0.005, **p<0.001,***p<0.0005, **** p<0.0001 (two way ANOVA)

Figure S5. SREBP2 deficiency impacts GC B cell differentiation but not IgA class switching.
(A) Frequency of indicated B cells subsets among untransduced tdTom+ B (B). Number of B cells from the indicated genotype recovered after 2.5 days of NB21 culture. (C) IgA and IgG1 class switch in GC B cells from the indicated genotype. (D) ELISPOT of IgA secreting PC on Lamina Propria of the mice of the indicated genotype.