Initiation of Antiviral B Cell Immunity Relies on Innate Signals from Spatially Positioned NKT Cells

Highlights

- NKT cells promote B cell immunity upon viral infection
- NKT cells are primed by lymph-node-resident macrophages
- NKT cells produce early IL-4 wave at the follicular borders
- Early IL-4 wave is required for efficient seeding of germinal centers

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In Brief

NKT cells are required for the initial formation of germinal centers and production of effective neutralizing antibody responses against viruses.
Initiation of Antiviral B Cell Immunity Relies on Innate Signals from Spatially Positioned NKT Cells

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SUMMARY

B cells constitute an essential line of defense from pathogenic infections through the generation of class-switched antibody-secreting cells (ASCs) in germinal centers. Although this process is known to be regulated by follicular helper T (ThF) cells, the mechanism by which B cells initially seed germinal center reactions remains elusive. We found that NKT cells, a population of innate-like T lymphocytes, are critical for the induction of B cell immunity upon viral infection. The positioning of NKT cells at the interfollicular areas of lymph nodes facilitates both their direct priming by resident macrophages and the localized delivery of innate signals to antigen-experienced B cells. Indeed, NKT cells secrete an early wave of IL-4 and constitute up to 70% of the total IL-4-producing cells during the initial stages of infection. Importantly, the requirement of this innate immunity arm appears to be evolutionarily conserved because early NKT and IL-4 gene signatures also positively correlate with the levels of neutralizing antibodies in Zika-virus-infected macaques. In conclusion, our data support a model wherein a pre-ThF wave of IL-4 secreted by interfollicular NKT cells triggers the seeding of germinal center cells and serves as an innate link between viral infection and B cell immunity.

INTRODUCTION

B cells are key elements of the adaptive immune response because they provide protection from pathogenic threats through the production of highly specific antibodies. In the past few years, there has been much attention on the generation of broadly neutralizing antibodies against highly variable or emerging pathogens, such as influenza, HIV, and Zika virus to develop effective vaccines capable of conferring broad and long-lasting immunity. Therefore, it is crucial to understand how viruses elicit B cell immunity during an infection process.

Pathogen recognition occurs in secondary lymphoid organs such as the spleen and lymph nodes (Batista and Harwood, 2009). In the lymph nodes, B cells recognize pathogens on the surface of subcapsular sinus (SCS) macrophages, migratory dendritic cells, and follicular dendritic cells (Carrasco and Batista, 2007; Gaya et al., 2015; Heesters et al., 2013; Junt et al., 2007; Pan et al., 2007; Qi et al., 2006). Following pathogen recognition, B cells process and present pathogen-derived peptides via major histocompatibility complex (MHC) class II molecules (Amigorena et al., 1994). This results in the recruitment of specific CD4+ T cells, which help the B cells to differentiate into either antibody-secreting plasma cells or germinal center (GC) cells. Within germinal centers, B cells undergo class-switch recombination and cycles of proliferation, somatic hypermutation, and affinity-based selection, finally resulting in the production of high-affinity antibodies (Victora and Nussenzweig, 2012). These processes are strictly dependent on co-stimulatory signals, mainly CD40L, ICOS, and cytokines, such as interferon-γ (IFN-γ), interleukin-4 (IL-4), and IL-21, provided by follicular helper T (ThF) cells (Weinstein et al., 2016). Because ThF cells are known to appear later during the immune response, how B cells initially seed germinal center reactions is still unclear.

In addition to pathogen-derived peptides, B cells are able to present glycolipid antigens, such as those present on the surface of Sphingomonas species (spp.), Borrelia burgdorferi, and Streptococcus pneumoniae, on CD1d molecules (Barral et al., 2008; Kinjo et al., 2011; Leadbetter et al., 2008). Populations of...
CD1d-restricted, innate-like T cells, known as natural killer (NK) T cells, recognize these lipids and provide cognate help to B cells. This NK T cell-mediated help enhances the production of antibodies by B cells (Bai et al., 2013; Barral et al., 2008; Leadbetter et al., 2008). Because there are no reports of lipid antigens originating from viruses, it is currently unclear whether NK T cells have a role in inducing antiviral B cell responses.

Here we show that, following respiratory viral infection, NK T cell-deficient mice are impaired in their ability to form germinal centers and immunoglobulin G (IgG) class-switched antibody-secreting cells (ASCs). In a striking contrast to B cell activation by glycolipid antigens, CD1d-mediated interactions between B cells and NK T cells are dispensable during viral infection. Instead, NK T cells promote B cell immunity through the early and localized production of IL-4 at the follicular borders, a process that is likely driven by CXCR3 and requires initial priming by resident macrophages through CD1d and IL-18. Importantly, this seems to be a conserved process because transcriptomic analysis of Zika-virus-infected macaques shows that early NK T and IL-4 gene signatures positively correlate with the levels of Zika-specific neutralizing antibodies. Therefore, our results indicate that the production of IL-4 by NK T cells immediately after viral infection contributes to a pre-TH1 IL-4 wave at the follicular borders that facilitates the initial seeding of germinal center cells.

RESULTS

NKT Cells Promote B Cell Immunity during Viral Infection

The role of NK T cells in inducing the humoral response to glycolipid-bearing antigens has been established (Bai et al., 2013; Barral et al., 2008; Leadbetter et al., 2008); however, because virus-derived lipid antigens are unknown, we sought to investigate the role of NK T cells in the induction of B cell responses during viral infections. Accordingly, we intranasally infected C57BL/6 wild-type and CD1d−/− mice, which lack CD1d and type I and II NK T cells, with 200 plaque-forming units (PFUs) of influenza A virus, Puerto Rico/8/34 (PR8) strain. After 7, 9, and 21 days of infection, lung-draining lymph nodes were harvested, and adaptive immune response development was assessed by flow cytometry. We observed a progressive expansion of Fas+GL7+ germinal center cells and CXCR5+PD-1+ Tfh cells during the course of infection in wild-type animals compared with uninfected ones (Figures 1A and 1B; Figures S1A–S1D). Interestingly, the formation of germinal center cells was significantly reduced in NK T cell-deficient mice at early time points, whereas the expansion Tfh cells remained similar throughout the infection process (Figures 1A and 1B; Figures S1A–S1D). We further compared the spatial organization of germinal centers in wild-type and CD1d−/− mice by immunohistochemistry. In wild-type animals, influenza infection induces both the enlargement of IgG+ B cell follicles as well as the formation of follicular GL7+ structures corresponding to germinal centers (Figure 1C). In contrast, in NK T cell-deficient animals, germinal center structures were reduced by day 9 of infection, a defect that was accompanied by a strong decrease in the generation of influenza-specific IgG1 ASCs (Figures 1C–1E). Interestingly, CD1d−/− mice also displayed a significant reduction in germinal center formation when infected with the vaccinia virus Western Reserve strain, suggesting that the ability of NK T cells to promote B cell responses is a general feature of viral infections (Figures 1F and 1G).

To address whether NK T cells would also induce B cell immunity after vaccination with viral antigens, we intranasally immunized wild-type and CD1d−/− mice with hemagglutinin (HA) trimmer from the PR8 influenza strain in the presence or absence of two different adjuvants: alpha-galactosylceramide (α-GalCer) or poly(I:C). We found that HA-specific germinal centers developed in lung-draining lymph nodes only when HA was administered together with either of these adjuvants (Figures S1E and S1F). Interestingly, the formation of HA-specific germinal centers and ASCs is completely abrogated in NK T-cell-deficient mice when α-GalCer was co-administered, demonstrating that NK T cells are essential to induce B cell immunity when exogenous glycolipids are used as adjuvants (Figures S1G and S1H). In addition, when poly(I:C) was co-administered, we observed a diminution in the number of IgM ASCs and a consistent, although non-significant, reduction in HA-specific germinal centers in CD1d−/− animals. These results indicate that NK T cells may also enhance B cell immunity during intranasal vaccination in the absence of exogenous glycolipids (Figures S1I and S1J).

NKT Cells Deliver Non-cognate Help to B Cells to Induce Antiviral Immunity

Having shown that CD1d−/− mice have diminished B cell responses during influenza and vaccinia virus infection, we sought to determine the mechanism by which NK T cells help B cells to mount an antiviral immune response. In the case of bacterium-derived glycolipids, NK T cells differentiate into CXCR5+PD-1+*Bcl6* follicular helper NK T (NKTfh) cells, which engage in CD1d-mediated cognate interactions with B cells, produce IL-21, and support germinal center responses (Chang et al., 2011; King et al., 2011). To address whether NK T cells differentiate into NKTfh cells after viral infection, wild-type mice were
Figure 2. NKT Cells Deliver Non-cognate Help to B Cells during Viral Infection
(A and B) Flow cytometry analysis of wild-type mice infected with influenza virus.
(A) Gating strategy for NKT and CD4+ T cells. B220+ cells were excluded from the analysis.
(legend continued on next page)
intranasally infected with influenza virus, and mediastinal lymph nodes were harvested after 0–9 days. Flow cytometry analysis revealed that, although conventional CD4+ T cells gradually express CXCR5, PD-1, Bcl6, and SLAM and produce high levels of IL-21 after infection, TCRβ+CD1d-tetramer+ NKT cells do not upregulate Bcl6, CXCR5, or PD-1 and express moderate levels of SLAM and IL-21 (Figures 2A and 2B; Figures S2A–S2D). These results indicate that NKT cells do not acquire NKTh cell phenotype after viral infection.

To further explore the localization of NKT cells within the tissues of infected mice, we stained mediastinal lymph nodes of wild-type mice with R-phycocerythrin-labeled, PBS-57-loaded CD1d-tetramer as reported previously (Lee et al., 2015). Lymph node sections were further stained for GL7 and analyzed by confocal microscopy. We found that tetramer-labeled cells localized in the periphery of GL7+ areas, whereas almost no tetramer-labeled cells were found inside these GL7+ areas (Figure 2C). This observation indicates that NKT cells are excluded from the germinal center after viral infection.

To directly address whether CD1d-mediated cognate interaction between B cells and NKT cells is required for the development of the antiviral B cell response, we generated CD1dfox/fox mice to delete CD1d specifically on B cells. To this end, two loxP sites were inserted, flanking exon 3 of CD1d, and CD1dfox/fox mice were then crossed with mice expressing the Cre recombinase under the promoter of CD19 or Mb1 genes (Figure 2D; Figures S2E–S2H). Deletion of CD1d was observed in approximately 80% of B cells in CD1dfox/foxCD19-Cre+ mice and in 100% of B cells in CD1dfox/foxMb1-Cre+ mice, indicating that the Mb1-Cre system is a more efficient tool to delete CD1d on B cells (Figures 2E–2G). Then, groups of CD1dfox/foxCD19-Cre+ and CD1dfox/foxMb1-Cre+ and their corresponding Cre littermates were intranasally infected with influenza virus, and mediastinal lymph nodes were harvested after 9 days. The formation of germinal centers and Th1 cells was similar in CD1dfox/fox CD19-Cre+ and CD19-Cre+ mice and in CD1dfox/fox Mb1-Cre+ and Mb1-Cre+ animals, indicating that CD1d-mediated interactions between B cells and NKT cells are not required for the development of germinal centers during viral infection (Figures 2H–2K).

**An Early NKT Cell Wave of IL-4 Precedes Late IL-4 Production by Th1 Cells**

One mechanism by which NKT cells might initiate antiviral B cell immunity in a non-cognate manner is likely to be via the production of B cell-modulatory cytokines. To examine the ability of NKT cells to produce these molecules, we first assessed their kinetics of activation after influenza infection by measuring the levels of the surface activation marker CD69. We observed that, although most NKT cells were negative for CD69 in uninfected mice (day 0), the number of CD69+ NKT cells peaked 3 days after infection, indicating that activated NKT cells accumulated early after viral challenge (Figure 3A). Interestingly, influenza infection not only results in an upregulation of activation markers but also in a robust production of IFN-γ and IL-4, with a similar percentage of NKT cells producing either cytokine (Figure 3B; Figure S3A). Notably, even though NKT cells produce IFN-γ after infection, they represent only 1% of the total IFN-γ+ cell population in lymph nodes (Figure 3C). In striking contrast, NKT cells represent up to 50%–70% of the total IL-4+ cells after influenza infection (Figure 3C); however, we did not detect NKT cell production of other type 2 cytokines, such as IL-5 and IL-13 (Figure S3B). These observations demonstrate that, although the population of NKT cells is much smaller than those of conventional T cells, they represent the main source of IL-4 in lymph nodes at the early stages of influenza infection.

Previous studies have shown that TH1 cells constitute essentially all of the IL-4-producing T cells in draining lymph nodes after infection (King and Mohrs, 2009; Reinhardt et al., 2009; Yusuf et al., 2010). However, these cells usually accumulate later during the infection process. To gain insight into the kinetics of IL-4 production by NKT and Th1 cells after influenza infection, we took advantage of IL-4 GFP reporter mice, which concomitantly express endogenous levels of IL-4 in addition to GFP (Mohrs et al., 2001). Flow cytometry analysis revealed an accumulation of NKT cells shortly after influenza infection, reaching a maximum number on day 3, and a gradual but continuous accumulation of Th1 cells, reaching a maximum number on day 9 (Figure 3D). In agreement with previous studies, we observed GFP expression in approximately 20%–40% of NKT cells even on day 0, suggesting that NKT cells already produce IL-4 mRNAs at steady state (Figure 3E; Lee et al., 2013; Mohrs et al., 2005). However, the percentage of GFP+ NKT cells peaks up to 60% to 80% after 3 days of infection, indicating that influenza infection rapidly induces IL-4 production by NKT cells (Figure 3E). Notably, both steady-state IL-4 producers and those that acquire the de novo ability to produce IL-4 can contribute to the early pool of IL-4 secreting (Figures S3C–S3E). In contrast, the percentage of GFP+ TH1 cells increases gradually after infection, with around 10% of GFP+ TH1 cells on day 3 and 30% of GFP-expressing TH1 cells on day 9 of infection (Figure 3E). These results indicate that, although NKT cells accumulate and become IL-4 producers rapidly after infection, TH1 cells differentiate and produce IL-4 later throughout the infection process.

To assess the contribution of NKT and TH1 cells to the pool of IL-4-producing cells at different times of infection, we gated on...
Figure 3. NKT Cells Generate a Pre-TfH Cell Wave of IL-4

(A) Flow cytometry analysis of CD69 levels in the TCRβ⁺ CD1d tetramer⁺ population after 0–6 days of influenza infection. MFI, mean fluorescence intensity.

(B) Representative plots showing the percentage of NKT cells producing IFN-γ (blue) and IL-4 (red) after 3 days of influenza infection.

(C) Flow cytometry analysis of IFN-γ (blue) and IL-4 (red) production on day 3 of influenza infection. Gates were drawn on IFN-γ⁺ and IL-4⁺ lymph node cells to analyze the percentage of NKT cells inside each cytokine⁺ population.

(D–F) Flow cytometry analysis of NKT (blue) and TfH (red) cells from IL-4 GFP mice infected with influenza virus.

(D) Numbers of NKT and TfH cells at different stages of influenza infection.

(E) GFP expression in NKT and TfH cells at different times of infection.

(F) Percentage of CD1d-tetramer⁺ NKT cells and CXCR5⁺ TfH cells inside the IL-4 GFP⁺ lymph node cell population at different stages of influenza infection.

In all charts, each dot represents one mouse. In graphs showing statistical analysis, data are representative of three independent experiments. Horizontal bars, mean; error bars, SEM. Student’s t test, ****p < 0.0001.
An Early NKT Cell Wave of IL-4 Occurs at the Follicular Borders

So far, we have defined the temporal frame of IL-4 production by NKT cells during the early stages of influenza infection. To gain insight into the spatial distribution of these IL-4-producing NKT cells, we infected wild-type, CD1d−/−, and IL-4 GFP reporter mice with influenza virus and harvested mediastinal lymph nodes after infection. The lymph nodes were incubated with labeled PBS-57-loaded CD1d-tetramer, and sections were further stained against B220 and CD169, a macrophage marker, and analyzed by confocal microscopy. Although CD1d-tetramer+ cells were nearly undetectable in uninfected animals, they were observed inside B cell follicles and in direct contact with CD169+ macrophages at the subcapsular sinus and interfollicular areas by day 3 of infection (Figures 4A and 4B). In contrast, CD1d-tetramer− cells were almost absent in lymph nodes from CD1d−/− animals, indicating that CD1d-tetramer+ cells are most likely NKT cells (Figures 4A and 4B). Interestingly, although the great majority of CD1d-tetramer+ cells located inside the B cell follicles do not express GFP, most of the GFP+ cells appear to be located in the areas surrounding B cell follicles (Figure 4C). These results indicate that early IL-4 production is restricted to the periphery of the B cell follicles, where antigen-specific B cells relocate to recruit T cell help after activation.

To get insight into the extent of cellular heterogeneity within the immune cells contributing to the early IL-4 wave, we characterized the IL-4 GFP+ population by flow cytometry 3 days post-infection with influenza virus. This experiment shows that approximately 70% of this population corresponds to NKT cells (CD1d-tet+), whereas the remaining 30% (CD1d-tet−) are mostly T cells (TCRb+), excluding a significant contribution of ILC2s to the early wave of IL-4 (Figure 4D; Figure S4B). To further study these two cell populations, IL-4 GFP+ CD1d-tet+ and IL-4 GFP+ CD1d-tet− cells were single cell-sorted and analyzed by RNA sequencing (RNA-seq). After processing (STAR Methods), we retained 222 cells from the CD1d-tet+ (n = 113) and CD1d-tet− (n = 109) populations in our single-cell RNA-seq analysis. T-distributed stochastic neighbor embedding (t-SNE) and shared nearest neighbor (SNN) clustering revealed that single cells from the two subsets (CD1d-tet+ and CD1d-tet−) each bifurcate into two separate clusters, reflecting divergent transcriptional states within these populations (Figures 4E; Figure S4C; Table S1). Independent of the cluster, most of these cells express CD3e, confirming their T cell origin (Figure 4F). A feature of the CD1d-tet− subset clusters is the expression of CCR7, L-selectin, and S1PR1, mRNAs encoding proteins that control T cell migration (Figure S4D). Furthermore, one of the CD1d-tet− clusters is characterized by cells expressing high levels of CD4 and the other one by cells expressing CD8a. Importantly, only a small proportion of the cells in these two clusters express Bcl-6, ruling out a significant contribution of TH cells to this early IL-4 wave (Figures 3F; Figures S4D and S4E). On the other hand, significant numbers of cells from the CD1d-tet+ subset express PLZF, a transcription factor that directs the NKT cell effector program (Figure 4F). Interestingly, a large proportion of cells from both CD1d-tet+ clusters express the chemokine receptor CXCR6, whereas only one of them expresses CXCR3 (Figure 4F). Both CXCR3+ and CXCR3− clusters express GATA-3, whereas the CXCR3+ group specifically expresses T-bet, and the CXCR3− one expresses RORγt, an observation that was further confirmed by flow cytometry (Figures 4G; Figures S4F and S4G).

CXCR3 has been reported to recruit T cells to the interfollicular zones, the areas where we find most of the IL-4 GFP+ NKT cells after infection (Groom et al., 2012; Sung et al., 2012). Interestingly, flow cytometry analysis of lymph nodes from influenza-infected mice revealed that 80% of IL-4 GFP+ NKT cells express CXCR3, whereas only 30% of IL-4 GFP− NKT cells express this marker (Figure 4H). Therefore, we wondered whether

Figure 4. The Early IL-4 Wave Is Localized at the Periphery of B Cell Follicles
(A–C) Confocal microscopy analysis of (A and B) wild-type and CD1d−/− and (C) IL4-GFP mice on day 3 of influenza infection. Lymph nodes were labeled with CD1d tetramer (magenta) and anti-B220 antibody (green in A and B and white in C). The arrows in (C) indicate CD1d tetramer+ cells expressing IL-4 (IL-4 GFP, green). Scale bars, 300 μm (lymph node) and 60 μm (section).
(D) Flow cytometry analysis of IL-4 GFP+ cells in mediastinal lymph nodes on day 3 of influenza infection, showing CD1d-tet− and CD1d-tet+ cells. The CD1d-tet+ population separates into two clusters (1, dark purple; 2, light purple), as does the CD1d-tet− population (3, light green; 4, dark green).
(E) t-SNE plots of CD1d-tet+ and CD1d-tet− subsets. The CD1d-tet+ population separates into two clusters (1, dark purple; 2, light purple), as does the CD1d-tet− population (3, light green; 4, dark green).
(F and G) Expression distribution (violin plots) in each population (horizontal axes) for (F) CD3e, Zbtb16, CXCR6, and CXCR3 and (G) Gata3, Tbx21, and Rorc. Expression level values are log2(tpm+1).
(H) Flow cytometry analysis of CXCR3 levels in IL-4-GFP− (red) and GFP+ (blue) NKT cells on day 3 of influenza infection. Lines connect cells from the same mouse.
(I) Flow cytometry analysis of IL-4 production by CXCR3− (red) and CXCR3+ (blue) NKT cells on day 3 of influenza infection. Lines connect cells from the same mouse.
(J) ELISPOT analysis of IL-4-producing CXCR3− (red) and CXCR3+ (blue) NKT cells sorted on day 3 of influenza infection. Each dot represents pooled NKT cells from 6 mice.
In graphs showing statistical analysis, data are representative of at least two independent experiments. Horizontal bars, mean; error bars, SEM. Student’s t test, *p < 0.05, **p < 0.01.
CXCR3+ and CXCR3+ NKT cells would have a differential ability to produce IL-4 after infection. To this end, we analyzed IL-4 production by both CXCR3+ and CXCR3+ NKT cell groups on day 3 of influenza infection by flow cytometry and enzyme-linked immunosorbent assay (ELISPOT). Strikingly, the number of cells secreting IL-4 in the CXCR3+ NKT cell sub-population is five times higher than in the CXCR3+ group (Figures 4I and 4J). These results suggest that CXCR3 preferentially drives the recruitment of IL-4-producing NKT cells to interfollicular areas.

Taken together, our results indicate that influenza infection triggers two spatiotemporally divergent waves of IL-4: an early wave, mainly produced by NKT cells and restricted to the periphery of B cell follicles, and a late one, produced by germinal center-resident TFH cells.

**Resident Macrophages Trigger an Early NKT Cell Wave of IL-4 and B Cell Immunity**

To define the immune machinery involved in the induction of this early NKT cell-derived IL-4, we specifically abrogated CD1d expression on different immune cell populations by crossing our CD1dflox/flox mice with animals expressing the Cre recombinase under the control of the Mb1, Lyz2, or CD11c gene promoters. Flow cytometry analysis showed an effective deletion of CD1d on Gr-1+CD11b+ neutrophils/myeloid-derived suppressor cells (MDSCs) in CD1dflox/flox-Lyz2-Cre+ mice and on MHCIId11c+ dendritic cells in CD1dflox/exCD11c-Cre+ mice (Figures S5A–S5C). However, no deletion of CD1d could be obtained in lymph node CD169+ macrophages in these transgenic lines (Figure S5D). Groups of CD1dflox/flox/Mb1-Cre+, CD1dflox/flox/Lyz2-Cre+, and CD1dflox/exCD11c-Cre+ and their corresponding Cre littermates were intranasally infected with influenza virus, and mediastinal lymph nodes were harvested after 3 or 9 days. Interestingly, similar percentages of IL-4+ NKT cells (day 3) and germinal centers and TFH cells (day 9) were observed among these transgenic lines (Figures 2H–2K and 5A–5C; Figures S5D–S5F). These results indicate that CD1d-mediated cognate interactions between NKT cells and B cells, neutrophils/MDSCs, or dendritic cells are not required for NKT cell-mediated induction of IL-4 production or germinal center formation.

Because we were unable to delete CD1d on CD169+ macrophages by using CD11c or Lyz2 Cre lines, we decided to take a macrophage depletion approach to assess the importance of these macrophages in the induction of IL-4 production by NKT cells. Accordingly, mice expressing the diphtheria toxin receptor (DTR) under the CD169 promoter, here named CD169-diphtheria toxin (DTR) cells, were treated with diphtheria toxin (DT) to deplete both subcapsular sinus and medullar macrophages from lymph nodes (Figure S6A). Although we observed a robust production of IL-4 in CD169-DTR/wild-type chimeras by day 3 of infection, NKT cells from CD169-DTR/CD1d−/− chimeras were impaired regarding this cytokine (Figure 6A). This result demonstrates that close interactions between CD169+ macrophages and NKT cells through CD1d are necessary to induce IL-4 production by NKT cells after viral infection.

The depletion of CD169+ macrophages has a more profound effect on the induction of IL-4 production by NKT cells than solely the deletion of CD1d on these cells. Therefore, we wondered whether CD169+ macrophages could enhance IL-4 synthesis through the secretion of co-stimulatory cytokines. To address this question, we stained lymph nodes on day 2 of influenza infection with antibodies against CD169, IL-18, and IL-33. These cytokines are secreted early after infection and have been shown to modulate the differentiation and function of innate and adaptive lymphoid cells (Kastenmüller et al., 2012; Sagoo et al., 2016). We detected the accumulation of IL-18, IL-19, IL-18, and IL-33 in both subcapsular sinus and medullar macrophages, suggesting that these resident macrophages are a source of inflammasome cytokines (Figures S6C and 6D; Figure S6A). Interestingly, their production is not the consequence of direct infection of macrophages because no viral replication was detected on them when a recombinant influenza virus carrying a GFP reporter gene was used (Manicassamy et al., 2010; Figure S6B).

Next we tested whether macrophage-derived IL-1β, IL-18, or IL-33 are required to induce NKT cell production of IL-4. These three cytokines are known to signal through the MyD88 adaptor protein (Adachi et al., 1998; Schmitz et al., 2005); hence, as a first approach, we compared the ability of wild-type versus MyD88−/− NKT cells to produce IL-4 after influenza infection. Our flow cytometry results show that, although a substantial proportion of lymph node NKT cells from wild-type mice produce IL-4, NKT cells from MyD88−/− animals are severely impaired in the production of this cytokine (Figure 6E). Importantly, their impairment is not due to lack of Toll-like receptor (TLR) signaling because NKT cells from TLR7−/− mice secrete IL-4 to the same extent as NKT cells from wild-type animals (Figure 6F). We then
Figure 5. Resident Macrophages Promote Early IL-4 Production by NKT Cells and Antiviral B Cell Immunity

(A–C) Flow cytometry analysis of IL-4 production by NKT cells in (A) CD1d<sup>flox/flox</sup>Mb1-Cre, (B) CD1d<sup>flox/flox</sup>Lyz2-Cre, and (C) CD1d<sup>flox/ex</sup>CD11c-Cre mice on day 3 of influenza infection: Cre<sup>−</sup> (blue dots) and Cre<sup>+</sup> (red dots).

(D and E) Confocal microscopy analysis of CD169<sup>DTR/+</sup> mice after 4 days of (D) PBS or (E) diphtheria toxin (DT) administration. Sections were stained with antibodies to B220 (white) and CD169 (green). Scale bars, 300 μm (left) and 60 μm (center and right).

(F) Flow cytometry analysis of IL-4 production by NKT cells from CD169<sup>DTR/+</sup>/PBS-treated or CD169<sup>DTR/+</sup>/DT-treated mice on day 3 of influenza infection: PBS (blue dots) and DT (red dots).

(G and H) Flow cytometry analysis of germinal center (G) and ThF (H) cells in CD169<sup>DTR/+</sup>/PBS-treated and CD169<sup>DTR/+</sup>/DT-treated mice on day 9 of influenza infection: PBS (blue dots) and DT (red dots).

In all panels, each dot represents one mouse. Data show one representative result from three experiments. Data represent mean ± SEM; two-tailed paired Student’s t test, **p < 0.01, ****p < 0.0001.

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Figure 6. Resident Macrophages Promote Early NKT Cell IL-4 Production through CD1d and IL-18

(A) Experimental scheme showing the strategy for the generation of chimeras lacking CD1d specifically on CD169+ macrophages. (B) Flow cytometry analysis of IL-4 production by NKT cells from mice subjected to the experimental scheme show in (A): wild-type chimeras (blue dots) and CD1d-/- chimeras (red dots).

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An Early IL-4 Wave Initiates B Cell Immunity to Viral Infection

Because of the central role of NKT cells in producing IL-4 early after infection, we wondered whether CD1d−/− mice would have reduced numbers of IL-4-secreting cells. To address this question, wild-type and CD1d−/− mice were infected with influenza virus, and the amount of IL-4+ cells was measured on day 3. The number of secreting IL-4− cells was significantly reduced in CD1d−/− compared with wild-type animals, indicating that other immune cells cannot compensate for the absence of NKT cells (Figure 7A). Next, to test whether the impaired B cell responses observed in CD1d−/− animals could be due to reduced IL-4 production, FVB wild-type and IL-4−/− mice were infected with influenza virus, and, 8 days later, mediastinal lymph nodes were harvested. A substantial expansion of germinal center and TFH cells was observed in wild-type mice compared with uninfected animals (Figures 7B and 7C). Interestingly, in IL-4−/− mice, there was a significant reduction in the proportion of germinal centers, whereas the differentiation of TFH cells remained similar to that of wild-type animals (Figures 7B and 7C). Furthermore, germinal center cells from IL-4−/− mice have a reduced capacity to class-switch to IgG1 (Figure 7D). These results show that IL-4 is necessary for the induction of the B cell response after influenza infection.

Then we asked whether the early wave of IL-4 was necessary for the induction of B cell responses after influenza infection. To address this question, wild-type mice were infected with influenza virus and treated them with either PBS or anti-IL4 blocking antibody (anti-IgM) in the presence or absence of IL-4. Interestingly, we observed that the addition of extracellular IL-4 to our ex vivo B cell cultures increases the oxygen consumption rate (OCR) in both naive and antigen-stimulated lymphocytes (Figure S7G). To address whether early IL-4 secretion could induce metabolic changes on B cells in vivo, we infected groups of mice with influenza virus and treated them with either PBS or anti-IL4 blocking antibody. We isolated lymph node cells on day 3 of infection, cultured them with anti-IgM and anti-CD3ε for 16 hr, and measured OCR levels in purified B cells after the incubation. We found that B cells from lung-draining lymph nodes have a higher OCR than B cells from non-draining lymph nodes, indicating that there is a metabolic reprogramming of B cells early after influenza infection (Figure S7H). Interestingly, the ability of B cells to increase their respiratory capacity after viral infection requires early IL-4 production because the upregulation of the OCR is diminished when the early IL-4 wave is inhibited (Figure S7H). These results suggest that the early IL-4 wave triggered after
infection could be regulating seeding of germinal center cells through the induction of metabolic changes in antigen-specific B cells.

**Early NKT Cell and IL-4 Signatures in Lymph Nodes from Zika-Infected Macaques**

We were interested to find out whether the early role for NKT cells and IL-4 in the initiation of antiviral B cell immunity might extend to other species. To address this question, we analyzed transcriptomic data from lymph nodes of rhesus macaques that were subcutaneously infected with Zika virus (Aid et al., 2017). We assessed the transcriptomic profiles of lymph nodes obtained on day 3 or 14 post-infection to identify pathways that correlated with Zika-specific neutralizing antibody titers on day 7 and day 14, respectively (Figure 7I; Figure S7I). Pathway enrichment analysis revealed a significant and positive correlation of NKT cells (normalized enrichment score [NES] = 1.48, false discovery rate [FDR] = 0.06), IL-4 (NES = 1.43, FDR = 0.02), IL-18 (NES = 1.42, FDR = 0.05), and STAT6 signaling (NES = 1.83, FDR = 0.02) signatures as early as day 3 of infection with neutralizing antibody titers at day 7 (Figure 7J). Furthermore, gene-interacting network inference showed the high interconnectivity of all these pathways, highlighted by co-expression of their leading genes with a central role of IL-4 in connecting all of these genes (Figure 7K). Importantly, no significant correlation between Th1 cell signatures (FDR > 0.20) and neutralizing antibodies was found at this early time point (Figure 7J). In contrast, a positive correlation between Th1 cell signatures and neutralizing antibodies was observed on day 14 of infection (NES = 1.23, FDR = 0.13) (Figures S7J and S7K). These correlates are in line with our previous findings, pointing out to a possible role for NKT cells at the early stages of B cell immunity, with Th1 cells only appearing later on the infection process. Furthermore, these analyses indicate that an early pre-Th1 wave of NKT cells and IL-4 might be conserved upon influenza and Zika virus infection in mice and primates, respectively.

**DISCUSSION**

Several studies have focused on the use of NKT cell agonists as vaccine adjuvants to boost NKT cell activity and immune responses. In particular, immunization with inactivated influenza virus or virus-derived proteins in combination with α-GalCer, a synthetic glycosphingolipid, results in strong protection against subsequent challenges with various strains of influenza virus (Artiaga et al., 2016; Galli et al., 2007). However, whether NKT cells could play a role in the induction of adaptive immune responses in the absence of exogenous glycolipids was not clear. Here we showed that NKT cells are required for the early seeding of germinal center B cells and the production of class-switched antibody-secreting cells in response to influenza infection. Furthermore, we found that NKT cells can enhance B cell immunity to influenza-derived proteins when administered together with TLR adjuvants. Our findings are in line with previous studies showing that NKT cells enhance the activity of CD8+ T cells after influenza infection even in the absence of exogenous glycolipoids (De Santo et al., 2008; Ishikawa et al., 2010). Therefore, the importance of NKT cells in the induction of adaptive immunity goes beyond the presence of glycolipid antigens.

How do NKT cells promote B cell responses during infection? Previous work showed that, in response to bacterium-derived glycolipids, NKT cells differentiate into NKTfH cells that engage in prolonged cognate interactions with B cells. These CD1d-mediated interactions rapidly induce germinal center and extrafollicular plasma cell formation, leading to the production of high titers of IgM and early class-switched antibodies (Bai et al., 2013; Barral et al., 2008; Chang et al., 2011; King et al., 2011; Leadbetter et al., 2008). However, we found that, upon influenza virus infection, NKT cells do not differentiate into NKTfH cells, nor they localize in germinal centers. Furthermore, CD1d-mediated cognate interactions between B cells and NKT cells are dispensable for the induction of antiviral B cell responses. Therefore, even though NKT cells promote B cell immunity in response to both bacterium-derived glycolipids and viral infection, the mechanism behind this regulation varies depending on the absence or presence of pathogen-derived glycolipids on the surface of the infectious agent.

Our results show that, during the initial stages of a viral infection, NKT cells become the main source of IL-4 in draining lymph nodes. Flow cytometry and single-cell RNA-seq analysis revealed that around 70% of the IL-4-producing cells are NKT cells, whereas the remaining ones are mainly T cells, excluding a significant contribution by ILC2s and Tfh cells to the IL-4 production at early stages of infection. This finding was unexpected because previous studies reported that Tfh cells constitute essentially all of the IL-4 producing cells in lymph nodes in

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**Figure 7. A Conserved Early IL-4 Wave Is Critical for Induction of B Cell Immunity**

(A) Flow cytometry analysis on day 3 of influenza infection. Graphs show the number of IL-4+ lymph node cells in wild-type (blue dots) and CD1d<sup>−/−</sup> (red dots) animals.

(B–D) Flow cytometry analysis on day 8 of influenza infection. Graphs show the percentage of (B) germinal center, (C) Thfh, and (D) IgG1+ cells in wild-type (blue dots) and IL-4<sup>−/−</sup> mice (red dots).

(E and F) Flow cytometry analysis on day 8 of influenza infection of wild-type mice treated with PBS or blocking α-IL-4 antibody during the initial 3 days of infection. Graphs show (E) germinal center and (F) IgG1+ cells in wild-type (blue dots) and α-IL-4 treated mice (red dots).

(G and H) Flow cytometry analysis on day 8 of influenza infection of CD1d<sup>−/−</sup> mice treated with PBS or IL-4 complex (IL-4c) during the initial 3 days of infection. Graphs show (G) germinal center cells and (H) IgG1+ cells in wild-type (blue dots) and IL-4c-treated mice (red dots).

(I) Experimental scheme showing the Zika virus infection strategy.

(J) Linear regression model of gene signatures for NKT cells, Thfh cells, IL-4, STAT6, and IL-18 from harvested lymph nodes on day 3 with neutralizing antibodies measured in plasma on day 7. Red squares, positive correlation; blue squares, negative correlation; white squares, no significant correlation.

(K) Gene-interacting network inference representing interconnectivity of NKT cells, IL-4, STAT6, and IL-18 pathways.

In (A)–(H), each dot represents a single mouse. Data are representative of three independent experiments. Statistical analysis: two-tailed Student’s t test, *p < 0.05, **p < 0.01, ***p < 0.001.
response to infection with different pathogens (King and Mohrs, 2009; Reinhardt et al., 2009; Yusuf et al., 2010). However, this previously unreported NKT cell-mediated wave of IL-4 production occurs at the early stages of infection, even before the appearance of ThF cells or germinal centers. Furthermore, this phenomenon is restricted to the follicular borders, where B cells migrate immediately after pathogen encounter. The spatiotemporally restricted production of IL-4 by NKT cells appears to instruct antigen-activated B cells at the periphery of B cell follicles early after infection for the subsequent seeding of germinal centers. Interestingly, a recent report showed that ThF cells only switch to IL-4 production at the late stages of the infection process (Weinstein et al., 2016). Therefore, it will be important to discern the contribution of the early NKT cell IL-4 wave at the follicular borders versus the late ThF cell IL-4 wave at germinal centers to the different processes involved in the development of B cell immunity.

The positioning of NKT cells at the follicular borders during the early stages of infection appears to be at the core of the NKT cell ability to initiate B cell immunity. Single-cell RNA-seq and flow cytometry analysis revealed that IL-4-producing NKT cells preferentially express CXCR3, a chemokine receptor used by T cell subsets to relocate to the interfollicular areas of the lymph node (Groom et al., 2012; Sung et al., 2012). However, whether NKT cells acquire CXCR3 after activation or whether a population of NKT cells already expressing CXCR3 expands after infection remains to be elucidated. Interestingly, Th2 cells have also been shown to locate and exert their functions at the periphery of B cell follicles (Turqueti-Neves et al., 2014). Therefore, we believe our study reveals a new layer of regulation of B cell immunity at the initial stages of the infection process, when B cells relocate to follicular borders and recruit T cell help.

The early production of IL-4 by NKT cells is mainly triggered by CD169+ macrophages and is mediated by CD1d interactions and IL-18 secretion. Interestingly, a great number of NKT-macrocephage contacts can be observed at the subcapsular sinus and interfollicular areas as early as 2 days post-influenza infection. Our findings are in line with the notion that NKT cell activation after infection is the result of a combination of both CD1d and cytokine signaling, with CD1d helping to position the cells and cytokines inducing the effector functions of NKT cells (Brennan et al., 2013). Furthermore, previous studies by others and us have shown that these macrophages can present bacterium-derived glycolipids on CD1d to mediate early NKT cell expansion and that IL-18 can enhance NKT cell activity in response to lipopolysaccharide (LPS), when co-administered with z-GalCer or in autoimmune models (Barral et al., 2010; Hägglof et al., 2016; Leite-De-Moraes et al., 2001; Nagarajan and Kronenberg, 2007). Therefore, it seems that the induction of diverse NKT cell effector functions by CD169+ macrophages and IL-18 is a conserved mechanism observed upon infection with various pathogens and in autoimmune disorders.

In this study, we have unveiled a new layer of regulation of B cell immunity at the early stages of viral infection that involves a tightly regulated NKT cell-IL-4-B cell axis. Interestingly, this phenomenon seems to be conserved upon viral infection in other species because our transcriptomic analysis of Zika-virus-infected macaques revealed a positive correlation between early NKT cell-IL-4 gene signatures and the serum levels of Zika virus-neutralizing antibodies. We believe our findings raise the prospect of harnessing the interplay between macrophages and NKT cells and the early production of IL-4 at follicular borders in future approaches for the design of new vaccination strategies.

STAR+METHODS
Detailed methods are provided in the online version of this paper and include the following:

- KEY RESOURCES TABLE
- CONTACT FOR REAGENT AND RESOURCE SHARING
- EXPERIMENTAL MODEL AND SUBJECT DETAILS
  - Mice
  - Infections and injections
  - Flow Cytometry
  - Immunohistochemistry
  - qRT-PCR
  - ELISPOT
  - Extracellular flux assay
  - scRNA-Seq Library Preparation
  - Single-cell RNA-seq Data Analysis
  - Transcriptomic analysis of samples from Zika infected macaques
- QUANTIFICATION AND STATISTICAL ANALYSIS
- DATA AND SOFTWARE AVAILABILITY

SUPPLEMENTAL INFORMATION
Supplemental Information includes seven figures and one table and can be found with this article online at https://doi.org/10.1016/j.cell.2017.11.036.

AUTHOR CONTRIBUTIONS
M.G. designed and performed experiments, generated CD1d-floxed mice, analyzed the data, and wrote the manuscript. P.B. performed some initial experiments, assist with the setup of in vivo infections, and contributed to scientific discussions. A.W.N. and A.K.S. performed and analyzed single-cell RNA-seq data. M.B., C.T., S.A., and P.M. performed experiments and contributed to scientific discussions. F.D.B. conceived the project, supervised the work, and wrote the manuscript. P.G.F. provided IL-33R/C0 Cre, and Lyz2-Cre mice and for scientific discussions. We thank Michael Kronenberg for the Mb1-Cre strain. We thank Mark Wilson for the IL-4-GFP reporter mice. We thank Andrew McKenzie for the IL-33R/C0 bone marrow. We thank Samuel W. Kazer and Sanjay M. Prakadan for assistance with RNA-seq.
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DECLARATION OF INTERESTS

The authors declare no competing interests.

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# STAR METHODS

## KEY RESOURCES TABLE

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**Bacterial and Virus Strains**

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**Chemicals, Peptides, and Recombinant Proteins**

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## Critical Commercial Assays

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## Deposited Data

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## Experimental Models: Cell Lines

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## Experimental Models: Organisms/Strains

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**Oligonucleotides**

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**Software and Algorithms**

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EXPERIMENTAL MODEL AND SUBJECT DETAILS

**Mice**

8-10 week old wild-type C57BL/6 and FVB/N mice were obtained from Jackson and Charles River Laboratories. Congenic CD45.1 mice were from the Francis Crick Institute, UK. Siglec1DTR/+ (CD169-DTR) mice were obtained from the Riken BioResource Center, Japan. CD19-Cre, CD1d−/−, IL-1R−/− and IL-18R−/− mice were obtained from Jackson Laboratories, USA. Lyz2-Cre, Cd11c-Cre, TLR7−/−, OT-II and Myd88−/− mice were obtained from Dr. Caetano Reis e Sousa, Francis Crick Institute, UK. IL-4/GFP-enhanced transcript (4Get) mice were obtained from Dr. Mark Wilson, Francis Crick Institute, UK. FVB/N IL-4−/− mice were obtained from Dr. Jessica Strid, Imperial College London, UK. Mb1-Cre mice were obtained from Dr. Michael Reth, Max Planck Institute, Germany. IL-33R−/− bone marrows (Townsend et al., 2000) were obtained from Andrew McKenzie, MRC laboratory of Molecular Biology, UK and Padraic Fallon, Trinity College Dublin, Ireland.

For the generation of CD1d conditional mice, we used homologous recombination to insert loxP sites flanking exon 3 of cd1d1 gene in B6 embryonic stem (ES) cells. Identification of ES cell clones with a single correct insertion of the targeted locus in the genome was performed through long-range PCR and Southern blot. Transgenic mice derived from these ES cells were crossed with B6 mice expressing the FLP recombinase to remove the frt-flanked PGK-neomycin sequence used for selection of ES cells that have inserted the targeted vector into their genome. Mice carrying the fox allele were crossed with mice expressing the Cre recombinase under the promoters of CD19, Mb1, Lyz2 and CD11c genes. Finally, mice were crossed again with the same floxed mice to make them homozygous for the floxed allele.

For the generation of mixed bone marrow chimeras, CD45.1 C57BL/6 or CD169-DTR of 6-8 weeks of age were irradiated with 2 doses of 500 rad, 4 hours apart. One day later, bone marrow cells were injected i.v. in recipient animals (2x10⁶ cells in total). Experimental animals were kept on acidified water for one week prior and 3 weeks post irradiation treatment. Chimeras were used after 8 weeks of reconstitution.

Mice were bred and maintained at the animal facilities of the Francis Crick and Ragon Institutes. All mice were maintained in individually ventilated cages and were used at the age of 8 to 12 weeks. Up to five mice per cage were housed in individually ventilated cages in a 12 hr light/dark cycle, with room temperature at 22 C (19 C-23 C change). They were fed with autoclaved standard pellet chow and reverse osmosis water. All cages contained 5 mm of aspen chip and tissue wiles for bedding and a mouse house for environmental enrichment. Littermates (males or females) were randomly assigned to experimental groups. Generally between 4 to 8 mice were used per experimental group. All experiments were approved by the Animal Ethics Committee of Cancer Research UK and the United Kingdom Home Office, and by the Institutional Animal Care and Use Committee of the United States.

**Infections and injections**

Mice were anesthetized i.p. with Ketamine/Xylazine (100 mg/kg body) and intranasally infected with 200 PFU of Influenza virus A/Puerto Rico/8/1934 (PR8) H1N1 strain or 100 PFU of PR8-OTII virus (Thomas et al., 2006) or 10³ PFU of Influenza NS1-GFP virus (Manicassamy et al., 2010) or 200 PFU of Vaccinia virus Western Reserve strain in 20 μl of PBS. Influenza virus was amplified on MDCK cells and VACV on DF-1 cells. Purification of viral particles was performed in a sucrose 30% cushion at 25,000 RPM for 2 hours in an SW32Ti rotor. For protein immunizations, 5 μg of PR8 Hemagglutinin (HA) trimmer, kindly provided by Daniel Lingwood, was intranasally administered together with 10 μg of Poly I:C (Inovigen) or 2 μg of α-GalCer (Enzo) in 20 μl of PBS. For depletion of CD169+ macrophages, mice were i.p. injected with 500 ng of Diphtheria toxin (Sigma) in 200 μl of PBS. For in vivo IL-4 blocking, mice were i.p. injected with 1.5 mg of anti-IL4 (11B11, BioXCell) in 200 μl of PBS at different days of infection. For the administration of IL-4 complex (IL-4c), 5 μg of recombinant mouse IL-4 (PrepoTech) was complexed to 25 μg of anti-mouse IL-4 (BioXCell, 11B11), 5 μg of recombinant mouse IL-12 (PrepoTech) was complexed to 25 μg of anti-mouse IL-12 (BioXCell, 7B23), 5 μg of recombinant mouse IL-15 (PrepoTech) was complexed to 25 μg of anti-mouse IL-15 (BioXCell, 14D11), 5 μg of recombinant mouse IL-21 (PrepoTech) was complexed to 25 μg of anti-mouse IL-21 (BioXCell, 11B11).
Splenic naive B cells were purified using Pan B cell Isolation Kit (Miltenyi) and stimulated with 10 ng/ml IL-4 (PrepoTech) and/or Extracellular flux assay instructions. Cells/well. IL-4 secreting cells were detected using the mouse IL-4 ELISPOT kit from BD Biosciences following manufacturer’s instructions.

To measure influenza-specific ASCs, Enzyme-linked immunosorbent spot (ELISPOT) multiscreen filtration plates (Millipore) were coated overnight at 4°C with serial dilutions of lymph node single cell suspensions. Plates were washed with PBS 0.01% Tween-20 and incubated for 1 hour with 1μg/ml Streptavidin-Alkaline Phosphatase (Sigma). Finally plates were washed and developed with 3-amino-9-ethyl-carbazole (Sigma).

To measure IL-4 production after influenza infection, NKT cells were sorted from mediastinal lymph nodes (TCRβ4+ CD1d-tetramer+) and incubated in RPMI complete medium supplemented with 1X Cell stimulation cocktail (eBioscience) for 32 hours at a density of 2.10^4 cells/well. To measure IL-4 production in the presence of IL-18, NKT cells were sorted from spleen (TCRβ4+ CD1d-tetramer+) and incubated with increasing concentrations of mouse recombinant IL-18 (R&D systems) for 32 hours at a density of 2.10^4 cells/well. IL-4 secreting cells were detected using the mouse IL-4 ELISPOT kit from BD Biosciences following manufacturer’s instructions.

Flow Cytometry
For labeling of surface markers, single cell suspensions from lymph nodes or lungs were incubated with anti-mouse CD16/32 (BD Biosciences) and LIVE/DEAD Fixable blue (Life Technologies) in PBS 2% FCS for 20 minutes on ice to block nonspecific antibody binding and exclude dead cells. Cells were then washed and stained for 20 minutes on ice with the indicated anti-mouse antibodies, PBS57-loaded CD1d tetramer (kindly provided by NIH tetramer core facility) and/or biotinylated HA (Sino biological). When using biotinylated antibodies or proteins, cells suspensions were washed following antibody labeling and incubated for 20 minutes on ice with labeled streptavidin. If labeling of transcription factors was performed, cells were fixed and permeabilized with Foxp3 transcription factor staining buffer set (eBioscience) according to manufacturer instructions and incubated for 30 min with anti-mouse antibodies. Finally, cells were either resuspended in 400μl of FACS buffer and analyzed on a Fortessa cytometer (BD Biosciences) or in RPMI media and used for cell sorting on a FACSaria II (BD, Bioscience).

For analysis of cytokine production, lymph node single cell suspensions were incubated 4 hours at 37°C in RPMI complete medium supplemented with 1X Cell stimulation cocktail plus protein transport inhibitor (eBioscience). After surface staining, cells were fixed and permeabilized with Cytofix/Cytoperm (BD Biosciences) according to manufacturer instructions and incubated for 30 min with anti-mouse antibodies. Finally, cells were resuspended in 400μl of FACS buffer and data were collected on a Fortessa cytometer (BD Biosciences) and analyzed with FlowJo (TreeStar).

Elispot
To measure influenza-specific ASCs, Enzyme-linked immunosorbent spot (ELISPOT) multiscreen filtration plates (Millipore) were activated with absolute ethanol, washed with PBS and coated overnight at 4°C with 1 μg/ml of PR8 virus or 2 μg/ml of PR8 HA (Sino Biological) diluted in PBS. Plates were subsequently blocked for 1 hour with complete medium and incubated for 24 hours at 37°C with serial dilutions of lymph node single cell suspensions. Plates were washed with PBS 0.01% Tween-20 and incubated for 1 hour with 1μg/ml of biotinylated anti-IgM or IgG1 (Southern Biotech) diluted in PBS 1% BSA. Then, plates were washed and incubated for 1 hour with 1μg/ml Streptavidin-Alkaline Phosphatase (Sigma). Finally plates were washed and developed with 3-amino-9-ethyl-carbazole (Sigma).

Flow Cytometry
For labeling of surface markers, single cell suspensions from lung or spleen were incubated with anti-mouse CD16/32 (BD Biosciences) and LIVE/DEAD Fixable blue (Life Technologies) in PBS 2% FCS for 20 minutes on ice to block nonspecific antibody binding and exclude dead cells. Cells were then washed and stained for 20 minutes on ice with the indicated anti-mouse antibodies, PBS57-loaded CD1d tetramer (kindly provided by NIH tetramer core facility) and/or biotinylated HA (Sino biological). When using biotinylated antibodies or proteins, cells suspensions were washed following antibody labeling and incubated for 20 minutes on ice with labeled streptavidin. If labeling of transcription factors was performed, cells were fixed and permeabilized with Foxp3 transcription factor staining buffer set (eBioscience) according to manufacturer instructions and incubated for 30 min with anti-mouse antibodies. Finally, cells were either resuspended in 400μl of FACS buffer and analyzed on a Fortessa cytometer (BD Biosciences) or in RPMI media and used for cell sorting on a FACSaria II (BD, Bioscience).

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Immunohistochemistry
For surface staining, cryostat sections (10μm thick) of lymph nodes were dried, fixed in 4% PFA for 10 minutes, blocked with PBS 5% BSA and then incubated with the following anti-mouse antibodies in 1% BSA for 1 hour: B220 Pacific Blue (RA3-6 B2, Biolegend), CD169 AF488 (Ser-4, ATCC), IgD FITC (11-26c.2a, BD Biosciences), Langerin AF647 (929F3.01, Dendritics) and GL-7 AF647 (GL7, Biolegend). For intracellular staining, sections were permeabilized with PBS Triton 0.3% for 5 minutes before blocking and then incubated with anti-mouse IL-10 (polyclonal, R&D), IL-18 (polyclonal, Abcam) or IL-33 (polyclonal, R&D), followed by incubation with AF555 anti-goat or anti-rabbit IgG (Invitrogen). CD1d tetramer staining was performed as previously described (Lee et al., 2015). Briefly, lymph nodes were incubated overnight with PE-labeled PBS-57 loaded CD1d tetramer in 2% FCS at 4°C. Next day, lymph nodes were washed and fixed with 4% PFA for 1 hour and snap frozen. 10μm sections were blocked with 5% BSA and stained with anti-PE antibody (polyclonal, Novus Biologicals) followed by goat anti-rabbit AF555 (Life Technology). Imaging was carried out on a LSM 780 (Zeiss) inverted confocal microscope using a Plan-Apochromat 40x NA 1.3 oil immersion objective.

qRT-PCR
NKT, Tfh cells and conventional CD4+ T cells from mediastinal lymph nodes were sorted at day 3 and 9 of infection using the gating strategy shown in Figure S2A. RNA was extracted from sorted cells using the RNeasy micro Kit (QIAGEN). Between 10^3 and 10^5 cells were used in each condition. mRNA was reversely transcribed into cDNA using random hexamers from the SuperScript III first-strand system (Thermofisher). cDNA was amplified using custom primer sets (Sigma) and SYBR green PCR master mix (Thermofisher) on a 7500 real-time PCR system (Applied Biosystems). The relative mRNA abundance of target genes was determined by means of the ΔΔCT method, using HPRT as housekeeping gene.

ELISPOT
To measure influenza-specific ASCs, Enzyme-linked immunosorbent spot (ELISPOT) multiscreen filtration plates (Millipore) were activated with absolute ethanol, washed with PBS and coated overnight at 4°C with 1 μg/ml of PR8 virus or 2 μg/ml of PR8 HA (Sino Biological) diluted in PBS. Plates were subsequently blocked for 1 hour with complete medium and incubated for 24 hours at 37°C with serial dilutions of lymph node single cell suspensions. Plates were washed with PBS 0.01% Tween-20 and incubated for 1 hour with 1μg/ml of biotinylated anti-IgM or IgG1 (Southern Biotech) diluted in PBS 1% BSA. Then, plates were washed and incubated for 1 hour with 1μg/ml Streptavidin-Alkaline Phosphatase (Sigma). Finally plates were washed and developed with 3-amino-9-ethyl-carbazole (Sigma).

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Extracellular flux assay
Splenic naive B cells were purified using Pan B cell Isolation Kit (Miltenyi) and stimulated with 10 ng/ml IL-4 (PrepoTech) and/or 5 μg/ml F(ab')2 anti-IgM (Jackson ImmunoResearch) for 16 hours. Lymph node cells were obtained after 3 days of influenza infection, cultured with 5 μg/ml F(ab')2 anti-IgM and 3 μg/ml anti-CD3ε (Biolegend, 145-2C11) and B cells were purified after 16 hours. B cells were diluted in 200 μl of PBS and administered i.p. on days 0, 1 and 2 of infection. For adoptive transfers, sorted NKT cells (8.10^5), OT-II CD4+ isolated CD4+ T cells (2.10^6) or bone marrow cells (2.10^6) were resuspended in 200 μl of PBS and injected in the tail vein.
were resuspended in Seahorse medium supplemented with 11mM glucose (Sigma Aldrich), 2 mM glutamax (Life Technologies) and 1 mM Sodium pyruvate (Sigma Aldrich) for 30 minutes. Data was recorded with the XF96 Extracellular Flux analyzer. Oligomycin A (3 μM), FCCP (5 μM) and Rotenone (5 μM) (all from Agilent) were added sequentially at indicated time points to generate the OCR profile.

**scRNA-Seq Library Preparation**
Plate-based scRNA-Seq libraries were prepared as previously described (Trombetta et al., 2014). Briefly, cells were FACS sorted into 96-well plates containing 10 μL of RLT lysis buffer (QIAGEN) supplemented with 1% β-mercaptoethanol (Aldrich) in each well. Two plates of GFP+CD1d-tetramer+ and two plates of the GFP+CD1d-tetramer− cells were sorted. Smart-Seq2 whole transcriptome amplification (WTA) was performed as per (Trombetta et al., 2014) with Maxima Reverse Transcriptase used in place of SuperScript II (Life Technologies). Libraries were sequenced on an Illumina NextSeq500, with 30 base-pair paired-end reads, to an average depth of 670k ± 30k reads per cell.

**Single-cell RNA-seq Data Analysis**
Sequenced reads were aligned to the UCSC mm10 mouse transcriptome and gene expression was calculated using RSEM as previously described (Tirosn et al., 2016). Cells with greater than 1,000 genes mapped and with more than 10,000 transcripts detected were used for downstream phenotypic analysis. Genes expressed in fewer than 15 cells were filtered out from further analysis, leaving 222 cells and 9,271 genes. Expression data was log transformed (Log2(tpm+1)) before further analysis. Clustering and differential expression were performed using the Seurat package (version 1.4.0.16) in R (version 3.3.2) (Satija et al., 2015). Briefly, we identified 708 variable genes with log-mean expression values greater than 0.05 and dispersion (variance/mean) greater than 0.8. A principal components analysis was then run over variable genes, and the first five principal components were selected for downstream analyses based on a combination of Q-Q plots and the standard deviation of PCs as determined by an elbow plot (calculated using the Jackstraw algorithm in Seurat). tSNE and SNN clustering (FindClusters function in Seurat with k.param = 20 & resolution = 0.6) were used to depict and cluster the cells in each analysis, respectively. Genes differentially expressed between identified clusters were calculated using the bimodal distribution option. Differentially expressed markers were removed if identified in less than 20 percent of cells in a given cluster or if the average log fold change between clusters was less than 0.5. Heatmaps depict the 25 most significantly differentially expressed genes (p values < 0.002) in each cluster (full data available in Table S1). Violin plots were generated over curated NKT genes.

**Transcriptomic analysis of samples from Zika infected macaques**
Genes signatures for NKT, NKT subpopulations, IL4, IL18, TfH, STAT6 and B cell signaling were compiled from different public repository (http://software.broadinstitute.org/gsea/mSigdb/, http://pathcards.genecards.org/, PMID:16698943, PMCID:PMC3528072). We correlated the expression of these signatures in lymph nodes from ZIKA virus infected Rhesus monkeys to neutralizing antibodies measured in plasma from the same animals on day 7 and day 14 (Aid et al., 2017). We used a linear regression analysis (R language) to identify genes that correlated significantly to neutralizing antibodies or viral loads using a p value cut-off of 0.05. Significant genes were then used to perform a genesets enrichment analysis using the above signatures (http://software.broadinstitute.org/gsea/index.jsp). Significant genes were used to infer gene interacting networks using GeneMania (http://genemania.org/) and Cytoscape (http://cytoscape.org/).

**QUANTIFICATION AND STATISTICAL ANALYSIS**
Statistical parameters including the exact value of n with the description of what n represents, the mean, the SEM and the p value are reported in the Figures and the Figure Legends. Statistical analyses were performed using Prism (GraphPad Software), two-tailed t test, p values < 0.05 were considered significant. In figures, asterisks stand for: *, p < 0.05; **, p < 0.01; ***, p < 0.001; ****, p < 0.0001).

**DATA AND SOFTWARE AVAILABILITY**
The single-cell RNA-seq data reported in this paper have been deposited in the NCBI Gene Expression Omnibus (GEO) database under accession number GEO: GSE103753.
Figure S1. NKT Cell-Deficient Mice Display Early Impairment in B Cell Immunity, Related to Figure 1

(A–D) Flow cytometry analysis of mediastinal lymph node cells from wild type or CD1d−/− mice that were intranasally infected with 200 PFU of influenza virus for (A–B) 7 or (C–D) 21 days. Representative contour plots display the percentage of (A–C) B220+ cells bearing biomarkers of germinal cell activity, Fas+GL7+, and (B–D) CD4+ T cells bearing biomarkers of TFH cells, CXCR5+PD-1+. Dot plots show quantification of (A–C) germinal center and (B–D) TFH cells.

(E and F) Flow cytometry analysis of mediastinal lymph nodes harvested at day 10 from wild type mice that were intranasally challenged with (E) 10 μg of HA trimmer or (F) PBS in the presence or absence of poly I:C or α-GalCer. Representative contour plots display the percentage of HA-specific B220+ cells differentiated into Fas+GL7+ germinal center cells.

(G and I) ELISPOT analysis of IgG1 HA-specific ASCs in mediastinal lymph nodes from wild type and CD1d−/− mice treated as in figure S1E.

(H and J) Flow cytometry analysis of HA-specific germinal center cells in mediastinal lymph nodes from wild type and CD1d−/− mice treated as in figure S1E.

In all quantification charts, each dot represents one mouse. Data are representative from two experiments. Statistical analysis two-tailed Student’s t test; *p < 0.05.
Figure S2. Expression Analysis of TfH Markers by NKT Cells and Strategy for the Generation of CD1d-Floxed ESCs, Related to Figure 2

(A) Flow cytometry gating strategy for the analysis of NKT cells (TCRβ+ CD1d tetramer+), CD4+ T cells (TCRβ+ CD1d tetramer− CD4+) and TfH cells (TCRβ+ CD1d tetramer− CD4+ CXCR5+ PD-1+) in mediastinal lymph nodes at day 9 of influenza infection.

(B) RT-qPCR analysis of IL-21 expression at day 3 and 9 of influenza infection by sorted NKT and TfH cells compared to CD4+ cells.

(C and D) Representative contour plots show the expression of (C) Bcl-6 and (D) SLAM by NKT, TfH and CD4+ T cells at day 3 and 9 of infection. Quantification on the right charts shows the mean fluorescence intensity for (C) Bcl-6 and (D) SLAM.

(E) Scheme showing the wild type and targeted CD1d locus containing two loxP sites flanking the exon 3 of CD1d as well as two FRT sites for the removal of the Neomycin resistance cassette.

(F) Specific fragments amplified by long-range PCR for the 5’ end (7.1kb) and 3’ end (6.6kb) of the clones that correctly performed homologous recombination.

(G) DNA from three positive clones was digested with SacI restriction enzyme and Southern blot performed using a labeled Southern probe. The upper band corresponds to the wild type locus (10.6kb) and the lower band to the targeted locus (7.3kb).

(H) Southern blot of the three positive clones performed with a labeled Southern probe against the Neomycin cassette. Orange arrow indicates the single 5.2kb band recognized by this probe, indicating single insertion of the construct. The clone 2, in red, was chosen for injections.

In all quantification charts, each dot represents one mouse. Data are representative from three experiments. Statistical analysis two-tailed Student’s t test; **p < 0.01, ***p < 0.001 and ****p < 0.0001.
A

B

C

Groups

Day -1: Adoptive transfer

1
IL4-GFP+ CD45.2
No adoptive transfer

2
IL4-GFP+ CD45.1
No adoptive transfer

3
IL4-GFP+ CD45.1 + IL4-GFP+ CD45.2 NKTs

4
IL4-GFP+ CD45.1 + IL4-GFP+ CD45.2 NKTs

D

E

(legend on next page)
Figure S3. Analysis of the NKT Cell Source of IL-4, Related to Figure 3

(A) Quantification of IFN-γ⁺ (blue) and IL-4⁺ (red) producing lymph node (left chart) or NKT cells (right chart) in mediastinal lymph nodes after 3 days of influenza infection.

(B) RT-qPCR analysis of IL-4, IL-5 and IL-13 expression at day 3 of influenza infection by sorted NKT cells compared to CD4⁺ T cells. Each dot represents one mouse.

(C) Adoptive transfer strategy for the analysis of the source of IL-4⁺ NKT cells after influenza infection.

(D and E) Flow cytometry analysis at day 3 of influenza infection of (D) mediastinal lymph node or (E) lung NKT cells from mice that were treated as in (C) to detect IL4-GFP expression.

Data are representative from two experiments.
Figure S4. Single-Cell RNA-Seq Analysis of IL-4⁺ Lymph Node Cells, Related to Figure 4

(A) Confocal microscopy of mediastinal lymph nodes sections at day 3 of influenza infection. Lymph nodes were labeled with PBS-57 loaded CD1d tetramer (magenta) and antibodies to B220 (white) and CD169 (green). IF, interfollicular. Scale bars, 300 μm (left); 60 μm (right). Chart on the right shows the quantification of NKT cells-macrophages contacts observed in the different areas of the lymph node at day 0, 1, 2 and 3 of infection in mediastinal lymph nodes. Data are representative from three experiments.

(B) Flow cytometry analysis of IL4-GFP⁺ CD1d-tet⁻ cells in mediastinal lymph nodes at day 3 of influenza infection showing TCRβ⁻ and TCRβ⁺ cells.

(C) Single-cell RNA-seq heatmap showing the expression of 25 significant marker genes (Table S1) for the CD1d-tet⁺ clusters (columns 1 and 2) and CD1d-tet⁻ clusters (columns 3 and 4). Single cells and genes were determined on the basis of unsupervised hierarchical clustering (Table S1).

(D) Expression distribution (violin plots) in each population (horizontal axes) for Ccr7, Seli, Stpr1, CD4, CD8a and Bcl6. Expression levels are Log₂(tpm+1).

(E) Flow cytometry analysis of IL4-GFP⁺ TCRβ⁻ CD1d-tet⁻ cells in mediastinal lymph nodes at day 3 of influenza infection showing CD4⁺ and CD8a⁻ cells.

(F) Representative contour plot show the expression of GATA-3 by NKT cells, ILC2s (Live/Dead⁻ B220⁻ CD3⁻ CD5⁻ CD11b⁻ CD11c⁻ CD127⁺ CD45⁺ GATA-3⁺) and CD4⁺ T cells at day 3 of infection. Quantification on the right chart shows the mean fluorescence intensity for GATA-3.

(G) Flow cytometry analysis of RORγt⁺ and T-bet⁺ NKT cells in mediastinal lymph nodes at day 3 of influenza infection. Histogram shows the levels of CXCR3 in RORγt⁺ and T-bet⁺ populations. Dot graphs show the quantification of CXCR3⁺ NKT cells in the RORγt⁺ (red dots) and T-bet⁺ populations (blue dots), with lines connecting NKT cells from the same mouse.

In graphs showing statistical analysis, data are representative of at least two independent experiments. Statistical analysis two-tailed Student’s t test; **p < 0.01, ***p < 0.001 and ****p < 0.0001.
**Dendritic cells**

Neutrophils/MDSCs

**CD11b**

Gr-1

**CD11c**

MHCII

**CD1d**

Events (% of max)

**CD1d+ DCs (%)**

**CD1d+ NFs/MDSCs (%)**

**Cre**

**Cre**

**A**

**B**

**C**

**D**

**E**

**F**

**G**

**H**

(legend on next page)
Figure S5. Conditional Targeting of CD1d in the Myeloid Lineage, Related to Figure 5

(A–C) Flow cytometry plots showing the gating strategy for (A) Gr-1+CD11b+ neutrophils (NFs) and myeloid-derived suppressor cells (MDSCs) and (C) MHCII+CD11c+ dendritic cells (DCs). Red and blue histograms depict the levels of CD1d in the gated populations in (A) CD1d^{floxed/Lyz2-Cre-} versus Cre+ and (B) CD1d^{−/−} mice or (C) CD1d^{floxed/CD11c-Cre-} versus Cre+ mice. These quantitative differences are depicted via dot plots; blue and red dots show CD1d+ cells in the Cre− and Cre+ animals, respectively.

(D) Confocal microscopy of mediastinal lymph nodes sections from wild type, CD1d^{−/−}, CD1d^{floxed/Lyz2-Cre-} and CD1d^{floxed/CD11c-Cre-} mice. Sections were stained with antibodies to CD169 (green) and CD1d (magenta). Scale bars, 60 μm.

(E–H) Flow cytometry analysis of mediastinal lymph node cells at day 9 of influenza infection. Representative contour plots show the percentage of B cells and T cells from (E–F) CD1d^{floxed/Lyz2-Cre-} and Cre+ or (G–H) CD1d^{floxed/CD11c-Cre-} and Cre+ mice that acquire (E and G) germinal center markers, Fas and GL7, or (F and H) Tfh cell markers, CXCR5 and PD-1. Quantifications are shown in dot plots, Cre− mice (blue dots) and Cre+ mice (red dots).

In all panels, each dot represents a single mouse. Horizontal bars – mean; error bars – SEM. Data are representative of three independent experiments. Statistical analysis, two tailed Student’s t test, *p < 0.05, ****p < 0.0001.
Figure S6. Intrinsic Role of MyD88 and IL-18R in NKT Cell Production of IL-4, Related to Figure 6
(A) Quantification of IL-1β+, IL-18+ and IL-33+ cells in the SCS and medullar areas from the images shown in Figures 6C and 6D.

(B) Confocal microscopy of mediastinal lymph nodes from wild type mice that were infected with 200 PFU of PR8 NS1-GFP and sacrificed after 3 days. Sections were stained with antibodies to CD169 (red), B220 (gray) and Langerin (magenta). Scale bars, 60 μm. Quantification of GFP+ cells is shown in the right chart.

(C–E) Flow cytometry analysis of chimeric mice that were generated by transferring a 1:1 bone marrow mixture of wild type (CD45.1+) and (C) wild type (CD45.2+), (D) MyD88−/− (CD45.2+) or (E) IL-18R−/− (CD45.2+) to sub-lethally γ-irradiated CD45.1 recipient mice. Eight weeks after bone marrow transplant, mice were infected with 200 PFU of influenza virus and mediastinal lymph nodes were harvested after further 3 days. Flow cytometry panels show the production of IL-4 by CD45.1+ and CD45.2+ NKT cells in the different sets of chimeras. Dot graphs show the quantification of IL-4+ NKT cells in the CD45.1+ population (blue dots) and the CD45.2+ population (red dots), with lines connecting the two NKT cell subpopulation coming from the same chimera.

In all panels, each dot represents a single mouse. Horizontal bars – mean; error bars – SEM. Data are representative of two independent experiments. Statistical analysis, two tailed Student’s t test, *p < 0.05, **p < 0.01.
A. Uninfected, Control/influenza, α-IL-4 (d0-d3)/influenza

B. Control, α-IL-4 (day -3)

C. lV OT-II T cells, Influenza-OVA, Harvest mLNs

D. Lymph node

E. Endogenous T cells, OT-II T cells

F. Wild type + OT-II, CD1d⁻/⁻ + OT-II

G. Oligomycin, FCCP, Rotenone

H. Oligomycin, FCCP, Rotenone

I. Zika virus infection, Harvest LNs and collect serum

J. Lymph node day 14 vs. Neutralizing antibodies day 14

K. Shared genes

(legend on next page)
Figure S7. Importance of Early IL-4 Production on B Cell Immunity, Related to Figure 7

(A) Representative confocal microscopy images of mediastinal lymph node sections from mice that were treated as in Figure 7E. Sections were stained with antibodies to the B lymphocyte marker, IgD (green) and the germinal center marker, GL7 (magenta). Scale bar, 450 μm. Quantification of the germinal center area in confocal images by ImageJ is shown in the right chart.

(B) Flow cytometry analysis at day 8 of mediastinal lymph nodes from wild type mice that were treated with PBS or anti-IL4 blocking antibody on day −3, and infected with 200 PFU of influenza virus at day 0. Contour plots show Fas+GL7+ germinal center cells. The responses are quantified in right charts.

(C) Experimental scheme showing the strategy for the generation of antigen-specific Tfh cell precursors.

(D–F) Flow cytometry analysis of mediastinal lymph node cells from mice treated as in (C). Representative contour plots display (D) the gating strategy used to differentiate endogenous CD4+ T cells from adoptively transferred OT-II cells, (E) the percentage of endogenous and exogenous cells bearing the Tfh cell markers PD-1 and CXCR5 and (F) the percentage of B cells that acquire GL7 marker. Quantifications are shown on the right charts.

(G and H) OCR at baseline and after sequential treatment with oligomycin, FCCP and Rotenone of B cells that were (G) stimulated ex vivo with anti-IgM and/or IL-4 or (H) isolated from lymph node cells of influenza-infected mice (day 3). Bar charts show the respiratory capacity obtained as (OCR after FCCP) − (basal OCR). Each dot represents an individual measurement.

(I) Experimental scheme showing Zika virus infection strategy and time points at which lymph nodes and serum were collected from macaques.

(J) Linear regression model of gene signatures for NKT cells, Tfh cells, B cells, IL-4, STAT6 and IL-18 from harvested lymph nodes at day 14 with neutralizing antibodies measured in plasma at day 17. Red squares represent positive correlation, blue squares represent negative correlation and white squares represent no significant correlation.

(K) Gene interacting network inference using GeneMANIA software representing interconnectivity of NKT, Tfh cells, B cells, IL-4, STAT6 and IL-18 pathways and the co-expression of their leading genes.

Unless stated, each dot represents a single mouse. Horizontal bars – mean; error bars – SEM. Data are representative of two independent experiments. Statistical analysis, two tailed Student’s t test, *p < 0.05, **p < 0.01 and ****p < 0.0001.