Enhanced IL-2 in early life limits the development of T_{FH} and protective antiviral immunity

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T follicular helper cell (T_{FH})–dependent antibody responses are critical for long-term immunity. Antibody responses are diminished in early life, limiting long-term protective immunity and allowing prolonged or recurrent infection, which may be important for viral lung infections that are highly prevalent in infancy. In a murine model using respiratory syncytial virus (RSV), we show that T_{FH} and the high-affinity antibody production they promote are vital for preventing disease on RSV reinfection. Following a secondary RSV infection, T_{FH}-deficient mice had significantly exacerbated disease characterized by delayed viral clearance, increased weight loss, and immunopathology. T_{FH} generation in early life was compromised by heightened IL-2 and STAT5 signaling in differentiating naive T cells. Neutralization of IL-2 during early-life RSV infection resulted in a T_{FH}-dependent increase in antibody-mediated immunity and was sufficient to limit disease severity upon reinfection. These data demonstrate the importance of T_{FH} in protection against recurrent RSV infection and highlight a mechanism by which this is suppressed in early life.

Introduction

Antibody-mediated immunity is central to long-term protection following natural infection or vaccination. The generation of high-affinity isotype-switched antibody predominantly takes place in germinal centers (GCs), anatomically distinct areas within the B cell zones of secondary lymphoid tissues. Within these GCs, B cells undergo somatic hypermutation and affinity maturation. A specialized subset of CD4 T cells, T follicular helper cells (T_{FH}) control the proliferation, maturation, and survival of GC B cells (reviewed in Vinuesa et al., 2016). T_{FH} cells migrate into B cell follicles via a CXCL13–CXCR5 interaction, where they promote B cell proliferation and differentiation through costimulatory CD40L–CD40 signaling and the secretion of soluble mediators, most notably IL-21 and IL-4.

Expression of the transcriptional regulator Bcl6 is necessary and sufficient for T_{FH} generation (Johnston et al., 2009; Nurieva et al., 2009); deleting Bcl6 in T cells completely ablates T_{FH} formation (Hollister et al., 2013). In adult mice, the initial differentiation of T_{FH} requires multiple signals, including costimulation via CD28 and ICOS (Choi et al., 2011; Weber et al., 2015) and cytokine-induced STAT3 signaling (Ray et al., 2014). Conversely, STAT5 signaling can inhibit this process (Johnston et al., 2012; Nurieva et al., 2012). Sustained signaling through ICOS, CD28, and STAT3 is also involved in the maintenance and function of T_{FH} cells within GCs (Linterman et al., 2014; McIlwain et al., 2015; Weber et al., 2015; Wu et al., 2015).

T_{FH} cells are required for optimal antibody responses in a range of contexts, including murine models of vaccination, allergic inflammation, and acute and chronic viral infection (Dolence et al., 2018; Grezmiel et al., 2017; Harker et al., 2011; Hollister et al., 2013). Modulation of T_{FH} responses, for instance via ICOS–ICOS-L blockade or removal of IL-6 family cytokines, which signal through STAT3, can reduce antibody responses and alter isotype balance (Harker et al., 2011; Harker et al., 2015; Uwadiae et al., 2019). In humans, the expansion of functional circulating CXCR5+ PD1+ CD4+ T cells (surrogates of lymphoid T_{FH}) correlates with the amplitude of antibody responses after influenza vaccination (He et al., 2013) and the emergence of broadly neutralizing antibodies in HIV-infected individuals (Locci et al., 2013). In contrast, individuals with loss-of-function mutations in key genes such as ICOS or STAT3 have reduced humoral immunity and defects in the differentiation and function of their T_{FH} (Grimbacher et al., 2003; Ma et al., 2012).

Early life is often associated with a failure to respond robustly to vaccination or generate long-term protective immunity post-infection (p.i.). This is particularly true in the case of respiratory syncytial virus (RSV) infection, the most common cause of infant bronchiolitis in high-income countries and a virus for which there is no licensed vaccine (reviewed in Openshaw et al., 2017). Infants can be infected multiple times with the same serotype of RSV in their first 3 yr of life, and even in adults,
protective levels of RSV-specific antibody responses are short-lived, and reinfection is common (Habibi et al., 2015). In mouse models, RSV-specific antibody responses are also limited after neonatal RSV infection compared with adults, characterized by reduced IgG (Tregoning et al., 2013). The underlying mechanism of this defect and whether the reduced humoral immune response contributes to disease on reinfection is currently unknown.

In the present study, the mechanisms underlying defects in early-life antibody-mediated immunity were explored. While RSV-infected adult mice generated robust antibody responses, those infected in early life failed to generate RSV-specific antibody responses. This failure to produce antibody was associated with limited differentiation of T<sub>FH</sub> and GC B cells in the lung draining LNs (dLNs) and enhanced susceptibility to secondary infection. T<sub>FH</sub>-deficient mice highlighted the importance of these cells in the development of protective immunity during adult RSV infection. We also demonstrate that IL-2–dependent suppression of T<sub>FH</sub> Responses in early life is partially responsible for limiting antibody production, and reversing this process enhances protective immunity.

**Results**

**RSV infection fails to generate T<sub>FH</sub>, GC B cells, and antibody in early life**

Since GC reactions are essential for a high-quality antibody response, we tested the role of age on the generation of GC responses to RSV. An initial characterization of the humoral immune response to RSV in 8-wk-old adult mice showed that T<sub>FH</sub> and GC B cell frequencies peaked at day 14 p.i., primarily in the lung dLNs (Fig. S1 A). Circulating and airway virus-specific antibodies are detectable by day 7 p.i. and increase between day 7 and 28 p.i. (Fig. S1, B and C).

To investigate the role of age on humoral responses 1-, 2-, and 8-wk-old mice were infected i.n. with the same initial dose of RSV. Age had a significant impact on the generation of circulating RSV-specific Ig, with IgG (Fig. 1 A) and locally secreted IgA (Fig. 1, B and C) significantly lower in mice infected in the first 2 wk of life, with no significant difference seen between mice infected at either 1 or 2 wk of age. Expression of RSV L gene in the lungs at day 4 p.i., the peak of RSV viral replication in mice, was similar among the different age groups (Fig. 1 D). Mice infected at 1 but not 2 wk of age had fewer CD4<sup>+</sup> T cells in their LNs at day 14 p.i. compared with adults (Fig. S1 D). Likewise total antigen-experienced CD4<sup>+</sup> T cells (Foxp3<sup>-</sup>CD11a<sup>+</sup>CD49d<sup>+</sup>) and regulatory Foxp3<sup>+</sup> CD4<sup>+</sup> T (T reg) cells in the LNs p.i. were lower in 1-wk-old mice compared with 8-wk-old mice, although proportions were similar (Fig. 1 E and Fig. S1 D). Both the number and proportion of these antigen-experienced cells that differentiated into CXC5<sup>+</sup> PD1<sup>+</sup> T<sub>FH</sub> was significantly higher when mice were infected as adults, with the proportion that differentiated into T<sub>FH</sub> increasing progressively with age (Fig. 1 E). In contrast, the number and proportion of Foxp3<sup>+</sup> T follicular regulatory cells (T<sub>FH</sub>) in the LNs were similar among the different ages of mice (Fig. S1 D). A similar proportion of cytokine-secreting (IFN-γ, IL-21, or IL-4) LN-derived CD4<sup>+</sup> T cells was observed across all ages after virus-specific peptide stimulation, although there was a trend toward decreased IL-21 production in 1-wk-old mice (Fig. S1 E). A similar pattern was seen in the B cell response, with fewer total B cells in the LN from mice infected at 1 wk of age (Fig. S1 D), and the number and proportion of GC B cells (IgD<sup>-</sup>CD38<sup>+</sup>IgM<sup>-</sup>GL7<sup>+</sup>) increased progressively with age at initial infection (Fig. 1 F). In line with previous studies (Ruckwardt et al., 2011), the proportion of virus-specific CD8<sup>+</sup> T cells present in the lungs in early life was greatly reduced compared with 8-wk-old mice, although proportions were similar in the LNs (Fig. S1 F).

GCs were readily detectable in both the LN and spleen of mice infected at 8 wk of age, while few were observed in mice infected at 1 wk of age (Fig. S1 G). Mice infected in the neonatal period lacked sustained GC activity, with fewer T<sub>FH</sub>, GC B cells, or IgG1<sup>+</sup> memory B cells (IgD<sup>-</sup>IgG1<sup>-</sup>GL7<sup>-</sup>CD38<sup>+</sup>) 8 wk after initial infection compared with mice infected at 8 wk of age (Fig. 1 G). Concomitantly with the reduced antibody-mediated immunity, mice initially infected at 1 or 2 wk of age showed limited protection from reinfection, suffering rapid weight loss that peaked at day 5 p.i. (Fig. 1 H). These data demonstrate a significant defect in the ability of neonatal mice to mount protective antibody-mediated immunity in response to RSV when compared with adult mice. This was associated with a selective failure in the ability of activated CD4<sup>+</sup> T cells to form T<sub>FH</sub> and B cells to differentiate into GC B cells.

**T<sub>FH</sub> are critical in regulating the outcome of secondary RSV infection**

Early-life infection was associated with a failure to generate T cell–dependent antibody responses in the secondary lymphoid compartment and enhanced disease during RSV reinfection. Since T<sub>FH</sub> are essential in generating GC-dependent antibody and T<sub>FH</sub> formation was found to be defective in early life, the importance of T<sub>FH</sub> in protective immunity against RSV was examined.

Bcl6 is essential for the differentiation of T<sub>FH</sub> in vivo (Johnston et al., 2009; Nurieva et al., 2009; Yu et al., 2009). Adult Cd4<sup>-</sup>-cre Bcld<sup>Δ/Δ</sup> (Bcld<sup>ΔCD4</sup>) mice, which are unable to generate T<sub>FH</sub>, and littermate controls (Bcld<sup>ΔWT</sup>) were infected with RSV, and disease progression was monitored. Disease severity as measured by weight loss was unaltered, peaking at day 7 p.i. in both Bcld<sup>ΔWT</sup> and Bcld<sup>ΔCD4</sup> mice (Fig. 2 A). Peak viral load at day 4 p.i. was comparable between WT and Bcld<sup>ΔCD4</sup> mice (Fig. 2 B), as were virus-specific H2-D<sup>B</sup> M<sub>87-195</sub> CD8<sup>+</sup> and I-A<sup>B</sup> M<sub>2-127-137</sub> CD4<sup>+</sup> T cell responses in the lungs (Fig. 2 C). CXCR5<sup>+</sup> PD1<sup>+</sup> T<sub>FH</sub> and CXCR5<sup>+</sup>PD1<sup>+</sup> T<sub>FH</sub> populations, which are thought to be associated with the GC itself, were limited in the lung dLNs of Bcld<sup>ΔCD4</sup> mice at day 14 after RSV infection but abundant in Bcld<sup>ΔWT</sup> controls (Fig. 2 D). T cell–specific deletion of Bcl6 also resulted in a several-fold reduction in the number of GC B cells in the LNs (Fig. 2 E). RSV-specific IgG1 and IgG2c were significantly lower in Bcld<sup>ΔCD4</sup> mice (Fig. 2 F).

While Bcl6 expression in T cells appeared to be redundant in determining the outcome of primary RSV infection, it did impact the antibody response. We therefore investigated the effect on disease when mice were reinfected with the same dose of RSV 5 wk after primary infection. Bcld<sup>ΔCD4</sup> mice had significantly
greater weight loss and delayed recovery compared with Bcl6WT littermate controls (Fig. 3 A). This was associated with an increased viral load at day 4 p.i. (Fig. 3 B) and increased cell numbers in the airways and lungs (Fig. 3, C and D). Eosinophil, but not neutrophil, proportions were increased in the airways of Bcl6ΔCD4 mice compared with Bcl6WT controls (Fig. 3 E), as were natural killer (NK) cell numbers (Fig. 3 F).

The magnitude of the virus-specific CD8 and CD4+ T cell response in the lungs was similar in Bcl6WT and Bcl6ΔCD4 mice (Fig. 3 G), but a greater proportion of virus-specific CD4+ and CD8+ T cells in the lungs of Bcl6ΔCD4 mice produced IFN-γ compared with controls (Fig. S2 A). The frequency of IL-17A+ and IL-13+ CD4+ T cells remained similar (Fig. S2 B). Despite this, and unlike RSV-infected mice, IAV-infected mice did not lose weight during reinfection with the same strain of IAV 4 wk later (Fig. S2 F), fitting with the observation that in IAV infection, other pathways, including tissue-resident T cells and non-TFH-dependent generation of antibody, may be sufficient to protect against re-infection (Miyauchi et al., 2016; Wu et al., 2014).
These data demonstrate that TFH are essential in promoting GC formation and antibody responses in response to respiratory virus infection but did not alter CD8\(^+\) T cell responses. Further, GC B cell activity was associated with protecting RSV-infected, but not IAV-infected, mice from disease on reinfection.

**Disease resolution after RSV infection of neonatally primed mice is TFH dependent**

To determine the role of Bcl6 expression by T cells on the outcome of RSV infection during early life, Bcl6\(^{WT}\) and Bcl6\(^{ΔCD4}\) mice were infected with RSV at day 7 of life. Weight gain was similar in both groups following infection (Fig. S3 A). At day 14 p.i., H2\(^{Db}\) M187-195\(^+\) CD8\(^+\) T cells were identifiable in the lungs of both Bcl6\(^{WT}\) and Bcl6\(^{ΔCD4}\) mice, although at a much lower proportion and frequency than was observed after infection of adult animals, fitting with previous observations of an altered, and reduced, CD8\(^+\) T cell hierarchy in neonatally infected mice (Ruckwardt et al., 2011; Fig. 2 C and Fig. S3 B). The proportion of CD4\(^+\) T cells specific for I-AbM2-127-37\(^+\) in the lungs and dLNs was similar to that seen 14 d p.i. in adults and did not change in TFH-deficient mice (Fig. 4 A). As in adults, Bcl6 deficiency prevented TFH differentiation entirely (Fig. 4 A). Fitting with this, GC B cells were low in frequency after early-life infection, with no significant difference seen between WT littermates and Bcl6\(^{ΔCD4}\) mice (Fig. 4 B). Virus-specific antibody was also limited after early-life infection, with no difference between WT and KO mice (Fig. 4 C).

Reinfection of neonatally primed mice 8 wk later resulted in rapid (within 24 h) weight loss that peaked around day 5 p.i.
Figure 3. Impaired humoral immunity limits protection from RSV reinfection. 7- to 8-wk-old WT (Bcl6fl/fl) or TFH-deficient KO (Cd4cre Bcl6fl/fl) mice were infected with 1 × 10^6 ffu RSV A2 i.n. and 4 wk later reinfected with 1 × 10^6 ffu RSV A2 i.n. (A) Weight change was monitored throughout infection. (B) RSV viral load in the lungs at day 4 p.i. detected by RT-qPCR. (C and D) Airway and lung cell numbers. (E and F) Proportion of eosinophils and neutrophils (E) and number of NK cells in the airways at day 4 p.i. as measured by flow cytometry (F). (G) Virus-specific CD8 and CD4 T cells from the lungs were analyzed by flow cytometry at day 14 p.i. (H) TFH and GC B cell numbers were determined in the lungs and LN by flow cytometry, and total numbers were enumerated. (I) RSV-specific IgG, IgG1, and IgG2a in the serum and IgA in the airways were determined by ELISA. Data are representative of n = 12–13 mice per group from two independent experiments. Data are shown with mean ± SEM. *, P < 0.05; and ***, P < 0.001.
Figure 4. Early-life immune responses to RSV are T<sub>FH</sub> independent. WT (Bcl6<sup>WT</sup>) or T<sub>FH</sub>-deficient KO (Cd4cre Bcl6<sup>KO</sup>) mice were infected as neonates (7 d old) with 2.5 × 10<sup>5</sup> ffu or as adults (7–8 wk old) with 1 × 10<sup>6</sup> ffu RSV A2 i.n. (A) Virus-specific CD4<sup>+</sup> T cells and T<sub>FH</sub> populations in the LNs of mice infected as neonates or adults were analyzed by flow cytometry at day 14 p.i. Gating for CXCR5<sup>+</sup>PD1<sup>+</sup> (purple line) and CXCR5<sup>hi</sup>PD1<sup>+</sup> (orange line) are highlighted. (B) LN GC B cells were analyzed by flow cytometry at day 14 p.i. in mice infected as neonates. (C) Virus-specific serum IgG1 and IgG2c was measured by ELISA.
Weight loss on reinfection was greater in Bcl6ΔCD4 mice than littermate controls, and recovery was delayed (Fig. 4 D). Cell counts in the airways, lungs, and LNs before and after reinfection were comparable between the two groups (Fig. S3 D). The enhanced weight loss and cellular inflammation seen on reinfection of neonatally primed mice is associated with both a T helper type 2 (Th2) skewed immune response not normally seen in adult mice, and exacerbated NK and CD8 T cell responses (Culley et al., 2002; Harker et al., 2014; Tregoning et al., 2008).

Virus-specific CD4+ and CD8+ T cell numbers in the lungs after reinfection were similar between WT and Bcl6-deficient mice (Fig. S3 E). After reinfection, WT mice had lower Th1 (IFN-γ), Th2 (IL-4+ and IL-13+), and T cytotoxic type 2 (IL-13+ CD8+) cell proportions at day 7, and IFN-γ+ CD8 T cells were slightly increased in Tfh-deficient animals at day 4 p.i. (Fig. 4, E–G). IFN-γ, IL-4, and IL-13 concentrations in the airways were similar between WT and Bcl6ΔCD4 mice (Fig. S3 F), as was eosinophilia (Fig. 4 H); however, airway NK cell frequencies were elevated in Bcl6ΔCD4 mice compared with controls (Fig. 4 I). IgG was detectable in the sera of control mice after primary neonatal infection and elevated on reinfection but was largely absent in Tfh-deficient mice (Fig. 4 J). Viral loads during both primary and secondary infection of neonates were similar between control and Bcl6ΔCD4 mice (Fig. 4 K).

The humoral response to neonatal RSV infection was significantly lower than in adults; this was associated with a failure of virus-specific CD4+ T cells to differentiate into Tfh. Genetic ablation of Tfh during neonatal infection had no further impact on the antibody response, indicating their absence is a critical limiting factor, although Bcl6ΔCD4 mice had a more prolonged disease course on viral reinfection.

IFN-γ and IL-2 signaling dampen B cell immunity in early life

Inadequate neonatal Tfh responses to RSV infection contribute to augmented disease on reinfection; we subsequently investigated why this was the case. In mice, IFN-γ has previously been found to limit antibody responses after early-life RSV infection (Tregoning et al., 2013) and has been shown to limit Tfh during severe malarial infection (Ryg-Cornejo et al., 2016). IL-2 signaling through STAT-5 also regulates Tfh differentiation in IAV infection (Ballesteros-Tato et al., 2012).

Anti-IFN-γ and anti-IL-2 treatment resulted in distinct outcomes on the immune response to RSV. At day 14 p.i., both treatments increased cell numbers in the LNs compared with isotype control–treated neonates, but only anti-IFN-γ increased cell numbers in the lungs (Fig. 5 A). Anti-IFN-γ treatment resulted in expanded H2-DR M187.195+–specific CD8+ and I-Aβ M2-127+37+ CD4+ T cell responses in the lungs compared with isotype-treated controls, changes not seen in the anti-IL-2 group (Fig. 5 B). In the LNs, anti-IL-2, but not anti-IFN-γ, treatment resulted in increased numbers of I-AβM2-127-37+ CD4+ T cells (Fig. 5 C). IL-2 signaling is important in the formation and function of Treg cells (reviewed in Malek and Castro, 2010); however, in the current study, transient neutralization of IL-2 did not alter the number of Foxp3+ Treg cells in the lungs, LNs, or spleen, while anti-IFN-γ treatment increased Treg cell numbers in the lungs, in line with its effects on total lung cell count (Fig. S4, A and B).

Depleting IL-2 during neonatal RSV did, however, increase the proportion of CD4+ T cells and I-AβM2-127-37+ CD4+ T cells that were CXCR5+ PD1+ (Fig. 5 D). It also altered the phenotype of the Tfh, increasing the expression of both Bcl6 and ICOS compared with isotype controls (Fig. 5 E and Fig. S4 C). Concomitant with this, the proportion of GL7+ CD38+ GC B cells in the LNs increased after anti-IL-2 treatment (Fig. 5 F). The frequency of GC B cells that were IgG1+ was also significantly increased after both anti-IL-2 and anti-IFN-γ treatment compared with control mice, while IgG2c expression was slightly reduced (Fig. 5 G). Neutralizing either IFN-γ or IL-2 during neonatal RSV infection resulted in increased RSV-specific Ig at day 14 p.i., with anti-IFN-γ also resulting in significantly increased RSV-specific IgM and IgG1 (Fig. 5 H). Together, these data showed that both IFN-γ and IL-2 signaling suppress IgG1 responses during early-life viral infection, with IFN-γ broadly limiting immune responses and IL-2 appearing to specifically attenuate the formation of Tfh and GC B cells.

IL-2–dependent suppression of Tfh prevents protective immunity in early life

As blockade of IL-2 signaling in early life enhanced GC reactions, we next examined its effects on long-term antibody-mediated immunity and the role of Tfh (experimental design shown in Fig. 6 A). Neutralization of IL-2 during primary neonatal RSV infection resulted in significantly increased RSV-specific Ig at day 14 p.i., with anti-IFN-γ also resulting in significantly increased RSV-specific IgM and IgG1 (Fig. 5 H). Together, these data showed that both IFN-γ and IL-2 signaling suppress IgG1 responses during early-life viral infection, with IFN-γ broadly limiting immune responses and IL-2 appearing to specifically attenuate the formation of Tfh and GC B cells.
in the lungs of reinfected Bcl6\textsuperscript{WT} mice, while no difference was observed in Bcl6\textsuperscript{ΔCD4} mice (Fig. 6 E). Anti-IL-2 treatment of neonatal WT mice also reduced the numbers of eosinophils and CD4\textsuperscript{+} and CD8\textsuperscript{+} T cells in the airways but had no significant effect on neutrophil number (Fig. 6 F). Functionally, there was a slight increase in the IFN-γ production capacity of CD8 T cells from the airways of Bcl6\textsuperscript{WT} mice treated with anti-IL-2, but IFN-γ and IL-13 production by CD4\textsuperscript{+} T cells was similar (Fig. 6 G). The proportion of T reg cells in the lungs and the LNs was similar across all groups (Fig. 6 H).
These data support the conclusion that IL-2 suppresses antibody-mediated immunity in early life in a TFH-dependent manner. Neutralization of IL-2 during primary infection promotes antibody responses and can improve protection and reduce disease on reinfection.

Multiple mechanisms contribute to IL-2–dependent suppression of TFH in early life

We next wanted to explore what blocks the development of TFH cells in early life. IL-2–induced STAT5 signaling has been shown to suppress both the differentiation and maintenance of TFH (Ballesteros-Tato et al., 2012; Johnston et al., 2012; Nurieva et al., 2012). Conversely, STAT3, and to a lesser extent STAT1 signaling, through cytokines such as IL-6 can promote TFH differentiation and function (Choi et al., 2013; Harker et al., 2013; Harker et al., 2011; McIlwain et al., 2015; Papillion et al., 2019). TFH differentiation in adult mice occurs within the first few days after acute infection (Choi et al., 2011); 3 d p.i., neonatal WT mice had a population of antigen-experienced, Foxp3−CD4+ T cells that had up-regulated CXCR5 and PD1 in both the LNs and the spleen. These cells were not present in Bcl6-deficient animals but were present in increased numbers in the LNs of neonatal WT mice after anti-IL-2 treatment (Fig. 7 A). Supporting their identity as early TFH, these cells...
also had reduced PSGL1 expression and were not Ly6C+ (Poholek et al., 2010; Fig. S5 A).

There was a significant increase in phosphorylated STAT5 (pSTAT5) and a decrease in pSTAT3 in neonatal antigen-experienced (CD44+Foxp3−GITR−) CD4+ T cells compared with those taken from adults at day 3 p.i., suggesting an environment enriched for STAT5 signaling cytokines in early life (Fig. 7 B). Expression of \( \text{Il2} \) and \( \text{Il6} \) in the spleens of uninfected 1-, 2-, and 8-wk-old mice did not vary depending on age, and gene expression was undetectable in the LNs (Fig. S5 B). However, 3 d after RSV infection, \( \text{Il2} \) transcripts were detectable in the LNs and were significantly higher in 1-wk-old than 8-wk-old mice, while no difference was seen in the spleen (Fig. 7 C). \( \text{Il6} \) expression was similar in both the LNs and spleen of 1- and 8-wk-old mice at 3 d p.i. (Fig. 7 C). Intracellular staining for IL-2 indicated that Foxp3−CD4+ T cells, alongside CD8+ T cells, were the main sources of IL-2 in the LN in response to polyclonal stimulation (Fig. S5 C). IL-2 production by CD4+ and CD8+ T cells was...
significantly increased in uninfected mice at 1 wk compared with 8 wk of age, with Foxp3− CD4+ T cells becoming the major source in the neonatal LN p.i. (Fig. 7 D). Foxp3 expression is known to prevent IL-2 production by T reg cells (Wu et al., 2006), however in neonatal but not adult mice, some Foxp3− CD4+ T cells produced IL-2, although a much lower proportion than the Foxp3+ CD4+ T cells (Fig. 7 D). This may indicate that Foxp3 expression in T reg cells has not fully stabilized at this age, in line with observations there is a delay in Foxp3 expression in early life (Fontenot et al., 2005). IL-2 production by Foxp3− CD4+ T cells remained significantly increased in neonatal mice compared with adults at day 3 after RSV infection, although production by Foxp3− CD4+ T cells and CD8 T cells was similar (Fig. 7 D). In the absence of ex vivo stimulation, IL-2 production by naïve Foxp3− CD4+ T cells was also observed in early life after viral infection, and IL-2+ cells expressed more IL-2 per cell than those from adults (Fig. 7 E).

The capacity of naïve CD44+ Foxp3− splenic CD4+ T cells from both 1- and 8-wk-old uninfected mice to phosphorylate STAT5 in response to recombinant early-life IL-2 (rIL-2) and STAT3 in response to IL-6 in vitro was examined. While pSTAT3 levels were similar in both age groups after IL-6 stimulation, pSTAT5 levels were significantly heightened in 1-wk-old mice compared with 8-wk-old mice when stimulated with IL-2 (Fig. 7 F). To determine if this increased sensitivity was associated with altered expression of IL-2R proteins we measured expression of the three subunits, CD25 (IL-2Ra), CD122 (IL-2Rb), and CD132 (common γc). CD44+ CD62L+Foxp3− cells from 1-wk-old mice had significantly increased expression of CD25 and CD122 compared with adult cells (Fig. S5 D). CD132 was undetectable above the negative control in all cell types. CD25 and CD122 expression was substantially lower than expression of the same proteins on Foxp3+ CD4+ T cells (Fig. S5 D), with only 20% of CD44+CD62L+Foxp3− cells expressing CD25 (Fig. S5 E), indicating that this change in receptor expression is unlikely to fully explain the increased sensitivity seen in neonatal CD4+ T cells.

These data suggest a predisposition of CD4+ T cells for enhanced STAT5 over STAT3 signaling in early life compared with adulthood, with both elevated expression of IL-2 and sensitivity to IL-2 present, potentially limiting TFH cell differentiation in early life.

Discussion
Antibody-mediated immunity is essential to long-term protection following infection or vaccination but is often defective in early life. Here, we found that T FH are essential for protective immunity after RSV infection but fail to differentiate after early-life infection. The absence of T FH in early life was associated with increased severity of disease on reinfection, and this phenotype was reproduced in T FH-deficient adult mice. Strikingly naïve CD4 T cells from neonatal mice had a heightened capacity to phosphorylate STAT5 in response to IL-2, and antibody-mediated immunity can be boosted by removing IL-2 signaling during neonatal infection.

In adult mice, RSV infection and other respiratory infections such as IAV (Yoo et al., 2012) robustly promote T FH and corresponding B cell and antibody responses. These T FH responses are essential in protecting adult mice from disease when RSV reinfection occurs. The disease on reinfection seen in Bcl6−/CD4 mice, which have a targeted depletion of T FH, is very similar to that observed when WT mice receive their first RSV infection as neonates (Culley et al., 2002). Here, we show a critical role for T FH in maintaining immunological balance, and in their presence, viral load, Th1 and Th2 cells, CD8 T cells, eosinophils, and NK cells are limited after RSV reinfection. T FH do not appear to regulate these immune responses during primary infection but likely act predominantly through their role in promoting T-dependent antibody. These antibodies will limit reinfection, in line with the reduced viral loads seen when T FH are present, thus reducing inflammation. In support of this, administration of RSV-specific antibodies before infection has been shown to protect mice from disease (Collarini et al., 2009; Fisher et al., 1999; Habibi et al., 2015; Karron et al., 2013; Radu et al., 2010).

While T FH enable protective immunity during adult RSV infection, their failure to form after early-life infection in mice is associated with an increase in the severity of disease on reinfection. T FH differentiation in neonatal mice has previously been shown to be limited in response to systemic vaccination with ovalbumin/alum or hemagglutinin from IAV (Debock et al., 2013; Mastelic et al., 2012; Mastelic Gavillet et al., 2015). It is important to note, however, that the immunological development of mice does not precisely mirror that observed in humans, with a 1-d-old mouse resembling a premature newborn human infant (Adkins et al., 2004; Simon et al., 2015). At 1 wk of age, mice then more closely resemble human infants at term and rapidly mature, with weaning occurring between 2 and 3 wk of age and reaching adolescence by 6–8 wk of age, as compared with 2 yr and ≥10 yr, respectively, in humans. Thus, the experiments performed here could be considered analogous to comparing infancy to young adolescence in humans. Fitting this comparison, human infants have reduced circulating T FH and plasmablast frequencies compared with older children and poorer response to the RTS,S malaria vaccine (Hill et al., 2020). Lower antibody titers are also seen in response to live attenuated RSV vaccines in infants compared with older age groups (Karron et al., 2013). Infants can also be repeatedly infected with the same serotype of RSV before the age of 3 yr, suggesting infection is insufficient to generate protective titers of antibodies (Openshaw et al., 2017), thus highlighting the significance and importance of defects in T FH differentiation in infancy. Caution should, however, be taken directly translating the findings observed here directly to human RSV infection. A substantially higher dose of RSV is required to infect a mouse than would be necessary to establish an infection in a human (Habibi et al., 2015; Príbul et al., 2008). Further, unlike a proportion of human infants, neonatal mice do not develop severe bronchiolitis upon RSV exposure, and indeed, as also shown here, they display less severe disease, as measured by weight loss, than mice infected with RSV as adults, though this may be a function of how infection interferes with appetite (Groves et al., 2020).

The comparison of neonatal and adult RSV infection in mice does, however, allow a detailed exploration of the mechanisms...
regulating TFH differentiation during respiratory virus infection at different ages and their implications in a well-characterized and relevant setting. In neonatal mice, defective TFH differentiation after protein immunization can be partially overcome by use of a suitable adjuvant (Mastelic-Gavillet et al., 2019; Mastelic-Gavillet et al., 2015; Vono et al., 2018). Here, we show that live RSV infection, despite possessing multiple stimulatory pathogen-associated molecular patterns and damage-associated molecular patterns, does not promote TFH differentiation in neonates. This is despite the presence of a similar frequency of virus-specific CD4+ T cells in neonates to that seen after adult infection, suggesting it is not simply due to reduced antigen presentation. Nor does it appear to be the results of enhanced TFH suppressing formation of TFH. Likewise, preexposure of neonatal mice to CpG before RSV infection is not sufficient to increase antibody responses (Yamaguchi et al., 2012). The limited generation of TFH in early life also appears to be independent of mouse strain, with both BALB/c and C57BL/6 background failing to generate TFH and antibody responses after neonatal RSV infection. Limited protective immunity after early-life RSV infection is a common feature across many, although interestingly not all, different murine genetic backgrounds (Tregoning et al., 2010), in line with this observation.

The few TFH that are induced in neonatal RSV infection display similar features to those observed after protein vaccination of neonatal mice, with reduced expression of Bcl6, CXCR5, and PD-1. This particular expression pattern has been shown to lead to preferential support of T-independent rather than T-dependent antibody responses in other models (Mastelic-Gavillet et al., 2019). In vaccination, this dysfunction has been associated with excess IL-4 signaling in early life (Debock et al., 2013), and fitting with this, neonatal exposure to RSV is also associated with enhanced IL-4 signaling (Culley et al., 2002). Overexposure of either neonatal or adult mice to IL-4 during primary infection does not, however, suppress the total amount of RSV-specific antibody generated (Harker et al., 2007; Harker et al., 2010), suggesting IL-4 signaling is not the key signal dysregulating TFH after early-life infection. Conversely, overexpression of IFN-γ, which antagonizes IL-4 production, does limit virus-specific antibody, and its neutralization during neonatal RSV infection leads to increased antibody responses (Harker et al., 2007; Harker et al., 2010; Tregoning et al., 2013). Interestingly, here, we find that IFN-γ does not limit TFH differentiation, as has been seen in malaria infection (Ryg-Cornejo et al., 2016), but rather suppresses overall adaptive immune responses.

Initial TFH differentiation from naïve CD4 T cells is known to rely on STAT3 signaling; in mice, this is driven through gp130-associated cytokines such as IL-6 (Harker et al., 2013; McIlwain et al., 2015; Ray et al., 2014). STAT5, which is potently activated by IL-2, antagonizes this process preventing TFH differentiation and favoring the differentiation of other CD4+ T cell subsets (Ballesteros-Tato et al., 2012; Johnston et al., 2012; Nurieva et al., 2012). While several other studies have shown the capacity of IL-2 to limit TFH differentiation in adult mice, here, we show for the first time that TFH differentiation during early-life infection is found to be exquisitely suppressed by IL-2, with virtually no TFH found in its presence. The removal of IL-2 during neonatal RSV infection leads to increased TFH formation, GC reactions, and virus-specific antibody and is associated with increased protective immunity on reinfection in adulthood.

IL-2 is among the first cytokines produced upon TCR engagement in a naïve CD4+ T cell. Given the potent ability of IL-2 to suppress T cell-dependent antibody responses, adult mouse models have highlighted that IL-2 signaling is therefore tightly regulated at multiple levels to enable TFH differentiation, including DC-dependent quenching of IL-2 through expression of CD25 (Li et al., 2016) and T cell desensitization (DiToro et al., 2018). We, however, find enhanced IL2 expression in the LNs of neonatal mice after viral infection compared with adults. This is associated with neonatal Foxp3+ CD4+ T cells possessing a significantly increased capacity to produce IL-2 both before and after infection. The increased IL-2 in the neonatal LN appears to impede TFH differentiation, and pSTAT5 is increased in activated neonatal CD4+ T cells compared with adult CD4+ T cells while pSTAT3 is decreased. Suppressing IL-2 leads to increased TFH frequencies at this early time point. We also found an alteration in the ability of neonatal naïve CD4+ T cells to respond to IL-2 signaling, with STAT5 more readily phosphorylated in response to IL-2 as compared with adult-derived CD4+ T cells showing an intrinsic difference between the naïve CD4 T cells at an early age compared with adulthood.

The precise mechanism underlying this enhanced IL-2 signaling in neonatal LNs remains unknown. It has been proposed that T reg cells and self-reactive CD4+ T cells participate in quorum sensing, coordinated by IL-2, whereby self-reactive CD4+ T cells promote T reg cell development and function, suppressing self-reactive T cell activity (Almeida et al., 2012; Amado et al., 2013). IL-2 from non-T reg cells is thought to be essential in the peripheral maintenance of T reg cells (Setoguchi et al., 2005), while IL-2 produced by single-positive CD4+ T cells is thought to be key in regulating T reg cell generation (Hemmers et al., 2019). In mice, the first few weeks of life are associated with rapid thymic output and the establishment of peripheral tolerance, including the progressive generation of Foxp3+ CD4 T cells (Fontenot et al., 2005). Fate-mapping of IL-2 also indicates the rapid accumulation IL-2-producing CD4 T cells in the periphery during this time period (Yamamoto et al., 2013), in line with the enhanced IL-2 production observed in the dLNs. It seems likely, at least in part, therefore that the increase in IL-2, and its suppression of TFH, seen in early life is partially a result of the requirement for IL-2 signaling in the establishment of peripheral tolerance to self-reactive and microbial antigens in early life. This may explain why, unlike in adulthood (Setoguchi et al., 2005), the failure of IL-2 neutralizing antibody administration in early life to alter T reg cell numbers p.i. It also highlights the potential challenges encountered when seeking to modulate this signaling pathway in early life.

Overall, this study demonstrates that protective antibody-mediated immunity against respiratory viral infection is dependent on TFH and is limited in early life. Enhanced IL-2 signaling in CD4+ T cells in neonatal mice prevents the differentiation of TFH, and their absence results in significant disease on reinfection. While it remains to be seen if this
precise mechanism is in action in human infants infected with RSV, this age group frequently experiences recurrent infection with a variety of pathogens in early life, and immunizations often display reduced efficacy. The recent observation of reduced Th1 responses in infants being immunized during the postnatal period further supports a similar process occurring in human babies (Hill et al., 2020). Identifying vaccine strategies that promote STAT3 signals while limiting STAT5 activation and ultimately favor Th1 generation could be an important strategy in enhancing infant immunity.

Materials and methods

Animals

All animal work was conducted in accordance with the UK Home Office and the local AWERB committee (UK Home Office project license P996A24e1). Specific pathogen–free 6- to 8-wk-old or pregnant female BALB/c and C57BL/6 mice were purchased from Charles River Laboratories. 6- to 10-wk-old B6.129S-Bcl6tm1.1Dent mice (Hollister et al., 2013; stock number 023727) were purchased from The Jackson Laboratory and bred with B6.Cg-Tg(Cd4-cre)1Cwi/Blufj (CD4cre) mice (Lee et al., 2001; stock number 022071; kindly provided by Professor Matthias Merkenschlager, Imperial College London, London, UK) at Central Biomedical Services (Imperial College London, UK) and Charles River Laboratories (Margate, UK). Genotypes were determined using DNA extracted from ear biopsies processed using Extract-N-Amp Tissue PCR Kit (Sigma-Aldrich), amplified by PCR, and separated by gel electrophoresis according to the Bcl6 and Generic Cre protocols provided by The Jackson Laboratory; custom PCR primers were purchased from Sigma-Aldrich.

In vivo viral infections

Plaque-purified RSV A2 was obtained either from ATCC or kindly provided by Dr Cecilia Johansson (Imperial College London, London, UK). RSV A2 was grown in HEp-2 cells and viral titer determined by focus-forming assay (Pribul et al., 2008). WT BALB/c mice were infected i.n. at 1, 2, or 8 wk of age with 2.5 × 10^6 focus-forming units (ffu) RSV A2 in 20, 40, and 100 µl PBS, respectively, and rechallenged with 10^6 ffu RSV A2 in 100 µl PBS at 8 wk after primary infection. WT C57BL/6 (Bcl6^fl/fl) or T<sub>Fli</sub>-deficient KO (Cd4cre Bcl6<sup>flox/flox</sup>) mice were infected i.n. at 1 wk or 7-8 wk of age with 2.5 × 10^5 or 10^6 ffu RSV A2, respectively. Mice initially infected as adults were rechallenged 4 wk later with 10^6 ffu RSV A2 i.n.; mice initially infected at 1 wk were rechallenged 8 wk later with 8 × 10^5 ffu RSV A2.

H1N1 IAV PR8 (strain A/Puerto Rico/8/1934 H1N1) was kindly provided by Professor Wendy Barclay (Imperial College London, London, UK). At 7-8 wk of age, mice were infected i.n. with 40 FFU IAV PR8 and rechallenged with the same dose 4 wk later. All infections were performed under isoflurane-induced anesthesia, and animals were subsequently weighed daily to monitor disease severity. Mice were housed in individually ventilated cages, and all procedures were approved by the UK Home Office and conducted in accordance with the Animals (Scientific Procedures) Act 1986.

In vivo cytokine depletion

IL-2 was depleted by i.p. administration of 100 µg anti-mouse IL-2 (clones 54B6-1 and JES6-5H4; BioXCell) on days −1, +2, and +5 p.i. Control animals were dosed to the same schedule with 100 µg rat IgG2a (clones LTF-2 and 2A3; BioXCell). IFN-γ was depleted by i.p. administration of 100 µg anti-mouse IFN-γ (clone XMG1.2; BioLegend) on days −1, +2, and +5 p.i., while control animals received 100 µg rat IgG1 (clone HRPN; BioXCell).

Cell recovery

Mice were exsanguinated via the jugular vein. Blood was collected using heparinized capillary tubes (Hirschmann) and then centrifuged to isolate plasma. Bronchoalveolar lavage was performed by cannulating the trachea and washing the lungs with sterile PBS (1 ml flushed into the airways three times for adult mice and 0.5 ml flushed three times for mice under 5 wk of age). After centrifugation, supernatant was taken for mediator analysis, and cells were resuspended and used for flow cytometry. The right lung lobe was snap frozen and then disrupted in RPMI 1640, 100 U/ml penicillin/streptomycin (Life Technologies) using a handheld homogenizer (Stuart SHM1). Whole homogenate was used to determine viral load and supernatant was used for mediator measurements. All other lung lobes, mediastinal LNs (dLN), and spleen were processed for flow cytometry. Lung tissue was weighed, finely chopped, and incubated for 45 min at 37°C in complete RPMI (RPMI 1640, 10% FCS, 2 mM L-glutamine, and 100 U/ml penicillin/streptomycin; Life Technologies) supplemented with 0.15 mg/ml collagenase D (Roche Diagnostics) and 25 µg/ml DNase I (Roche Diagnostics). Lung, LN, and spleen were passed through 100-µm cell strainers (Greiner Bio-One) to create single-cell suspensions in complete RPMI. Lung and splenic cell suspensions were centrifuged, supernatant removed, and the cells treated with ACK buffer (150 mM ammonium chloride, 10 mM potassium bicarbonate, and 0.1 mM EDTA) for 3 min at room temperature and then washed and resuspended in complete RPMI. Total live cell counts were obtained using trypan blue exclusion staining and FastRed 102 counting slides (Kova International).

Ex vivo cell stimulations

For detection of T cell cytokine production, single-cell suspensions were stimulated with 5-h incubation with 10 ng/ml PMA (Sigma-Aldrich), 0.5 µg/ml ionomycin (WVR), 10 µg/ml Brefeldin A (Sigma-Aldrich), and 50 µl/ml recombinant mouse IL-2 (R&D Systems) in complete RPMI at 37°C. Virus-specific T cell cytokine production was induced by 5-h incubation with RSV peptides, including 3 µg/ml CD8 stimulating M249-249 (SYIG-SINNI) and F55-93 (KYKNAVEL) and 5 µg/ml CD4 stimulating G181-197 (TCWAICKRPNKKPGK), F51-66 (GWYTSVITIELSNIKE), and P19-55 (SIISVNDIEVTKESP) for BALB/c and 3 µg/ml CD8 stimulating M2187-195 (NAITNAKII) with 10 µg/ml CD4 stimulating M309-323 (NGKAFKYIKPQSQF) and M2-126-39 (NY-FEWPPHALLVRQ) for BL/6 samples.

For detection of STAT phosphorylation, ex vivo tissues were harvested directly into warm 1% paraformaldehyde (PFA) in PBS and incubated for 10 min at 37°C. A single-cell suspension was then generated for subsequent staining. For detection of
STAT phosphorylation following in vitro cytokine stimulation, single-cell suspensions were rested in RPMI 1640 and 100 U/ml penicillin/streptomycin (Life Technologies) for 2 h and then stimulated with 150 U/ml rIL-2 or 50 ng/ml rIL-6 (both from BioLegend) for 15 min. Cells were then fixed with 1% PFA (Sigma-Aldrich) in PBS for 15 min. All steps were performed at 37°C. Cells were subsequently frozen in True-Phos Perm Buffer (BioLegend) at −80°C until processed as described for flow cytometry.

**Flow cytometry**

Flow cytometry was performed as described previously (Pyle et al., 2017). All antibodies are from BioLegend unless otherwise stated. Single-cell suspensions were first stained with UV fixable viability dye (BioLegend) in PBS for 20 min at 4°C. The following anti-mouse antibodies were diluted in PBS containing 2% fetal bovine serum (Life Technologies) and 2 mM EDTA and used for surface staining, in combination with a 1 in 100 dilution of TruStain FcX antibody (BioLegend): anti-CD4 (RM4-5), anti-CD3e (145-2C11; eBioscience, Life Technologies); anti-CD8 (53–6.7), anti-B220 (RA3–6B2), anti-CD19 (6D5), anti-CD11c (N418; eBioscience); anti-CD11b (M1/70), anti-ly-6C (HIK1.4), anti-ly-6G (IAE (A7R34), anti-CD45 (30–F11), anti-CD40.2 (53–2.1), anti-Nkp46 (29A1.4), anti-CD64 (X54–5/7.1), anti-IA/le (M5/114.15.2), anti-CD11a (M17/4), anti-CD49d (R1–2; eBioscience); anti-CD44 (IM7), anti-Siglec-F (BD), anti-GITR (DTA–I; BD), anti-CD68 (FA–II), anti-CD62L (MEL–14), anti–PGL1 (2P1H1), anti–PD–1 (29F.1A2), anti–CXCR5 (L139D7), anti–ICOS (C989.4A), anti–CD38 (90; eBioscience); anti–GL7 (GL7), anti–Fas (1A5A7; eBioscience), anti-CD138 (281–2), anti–IgG1 (RMG1–1), anti–IgG2a (RMG2–6a), anti–IgD (11–26c.2a), anti–IgM (RMM–1), anti–KLRG1 (MAFA), anti–CD25 (PC61), CD80 (16–10A1), Kb M242–90 (for BALB/c mice), or Db M187–195 and I–Ab M27–37 (for C57BL/6 mice) tetramers. Tetramers were kindly provided by the NIH Tetrimer Core Facility, either pre–folded or as monomers, which were folded in–house with SA–PE (Invitrogen). For intracellular cytokine staining, cells were fixed with 1% PFA (Sigma–Aldrich) in PBS for 10 min at room temperature and stained with the following antibodies in permeabilization buffer (eBioscience): anti–mouse–I–4 (IIB1), anti–IL–7A (TCL1), anti–IFN–γ (XMGL1.2), anti–IL–2 (JES6–5H4), and anti–IL–13 (eBio13A; eBioscience). For intranuclear staining, cells were permeabilized and fixed with Transcription Factor Fixation/Permeabilization Kit (eBioscience) and stained with the following antibodies in permeabilization buffer (eBioscience): anti–mouse–Foxp3 (FJK–165; eBioscience) and anti–Bcl6 (K12–91; BD). For pSTAT detection, cells were fixed with 1% PFA in PBS for 10 min at room temperature, permeabilized with True–Phos Perm Buffer (BioLegend) at −20°C, incubated with a 1:100 dilution of TruStain FcX antibody for 1 h, and stained with mouse anti–Stat5 (pY694, BD) and anti–Stat3 (pS727; BD) alongside surface–stain antibodies detailed above in permeabilization buffer (eBioscience). Samples were acquired using a five–laser LSRS Fortessa (BD) and analyzed using FlowJo (BD).

**Immunofluorescence microscopy**

Preparation and imaging of dLN and spleen was performed as previously described (Wong et al., 2019). Briefly, tissues were harvested directly into 4% PFA and 5% sucrose in PBS and fixed for 4–6 h and transferred to 10% in PBS for 1 h and then 30% sucrose overnight. All steps were performed at 4°C. Tissues were then frozen in optimal cutting temperature compound, and 8–μm cryostat sections were prepared, fixed with 4% PFA, and blocked with blocking solution (PBS containing 3% normal goat serum, 3% normal donkey serum, 1% BSA, 0.05% Tween 20, and 0.1% Triton X–100). Sections were then stained with rat anti–mouse IgD–Alexa Fluor 647 (11–26c2a), rat anti–mouse/human GL7–Alexa488 (GL7), and anti–mouse CD4–biotin (RM4–5) diluted in blocking solution for 2 h at room temperature, washed, incubated with streptavidin–BV421 diluted in blocking solution for 2 h, washed, refixed with 4% PFA, and mounted. Sections were imaged using a Leica SP5 MP/FLM inverted microscope and images analyzed in using Fiji software.

**Quantitative RT–PCR**

RNA was extracted from lung, dLN, and spleen homogenate using RNeasy Mini Kit columns (Qiagen) according to the manufacturer’s instructions and converted to cDNA using High–Capacity cDNA Reverse Transcription Kit (Thermo Fisher) or GoScript Reverse transcription Kit (Promega). TaqMan assays (Thermo Fisher) were used to test the expression of IL2 and IL6 relative to Gapdh. RSV L gene was quantified relative to 18s housekeeping gene using a 900 nM forward primer (5′–GAA CTCACTGTTAGGTGAATTGTGCA–3′), 300 nM reverse primer (5′–TTCCAGCTATTCTTCTGCAAT–3′), and 175 nM probe (5′–6–carboxyfluorescein–TTGAACTCTTCTCACACATCGGT–6–carboxytetramethylrhodamine–3′) as described in Harker et al. (2014). Quantitative RT–PCR was performed using a Viia7 RT–PCR System (Thermo Fisher).

**Antibody and cytokine measurements**

IgE, IL–4, IL–13, and IFN–γ were measured using sandwich ELISA kits (Invitrogen) according to manufacturer’s instructions. For quantification of virus–specific Ig, IgG1, IgG2a/c, IgA, and IgM, ELISA plates were coated with RSV antigen grown in the laboratory, and bound antibody was detected using the SBA Clonotyping System–HRP (Southern Biotech).

**Statistics**

GraphPad 6.0 (GraphPad) was used for all statistics. For weight loss, area under the curve was calculated and significance tested by a Mann–Whitney U test (data with two groups) or one–way ANOVA (data with more than two groups). For experiments with only two groups, a nonparametric Mann–Whitney U test was used. For groups with three or more groups, a nonparametric Kruskal–Wallis H test with a Dunn’s test was used. In all cases *, P < 0.05; **, P < 0.01; and ***, P < 0.001.

**Online supplemental material**

Fig. S1 demonstrates that adult GC B cell and T_{H4} Responses peak at day 14 after RSV infection, with local and systemic antibody responses increasing between 1 and 3 wk p.i. In contrast, adaptive immune responses and GC formation is significantly reduced in mice infected as neonates. Fig. S2 shows that Bc1 deficiency in T cells restricts antibody responses to IAV after...
primary infection but does not result in enhanced disease upon reinfection. Fig. S3 illustrates that the immune response to primary RSV infection in neonatal mice is not dependent on Bcl6 expression in T cells, using WT (Bcl6+/+ and TFP-deficient KO (Cd4cre Bcl6+/+) mice. Fig. S4 details that neutralization of IL-2 during neonatal RSV infection limits lung and airway inflammation upon rechallenge with RSV in a Bcl6-dependent manner and does not affect numbers of T reg cells or T FR. Fig. S5 demonstrates that CD8+ T cells and Foxp3+CD4+ T cells are the main source of IL-2 in neonatal secondary lymphoid tissue following RSV infection and that IL-2 receptor subunit expression is increased on neonatal naive Foxp3+CD4+ T cells, compared with adults.

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Viral disruption of early-life antibody
Figure S1. Age-related reduction in adaptive immune responses to RSV infection. 8-wk-old BALB/c mice were infected 8 × 10^5 ffu RSV A2 i.n. (A) TFH (CD19−CD8−Foxp3−CXCR5+PD1−CD49d+CD11a+) and GC B cell (IgD−IgM−CD38−GL7+CD19+) numbers were determined in the lungs, mediastinal LNs, and spleen p.i. (B and C) RSV-specific Ig, IgG1, and IgG2a in the serum (B) and IgA in the airways (C). (D–G) 1-, 2-, and 8-wk-old BALB/c mice were infected 2.5 × 10^5 ffu RSV A2 i.n. and sacrificed at day 14 p.i. (D) The total number of CD4 T cells, B cells, and Foxp3+ CD4 T cells was enumerated. (E) Cytokine production by LN CD4 T cells after RSV peptide stimulation. (F) Virus-specific CD8 T cells in the lungs and LNs. (G) LN and spleen (SP) from 1- and 8-wk-old mice were imaged by immunofluorescence microscopy for IgD, CD4, and GL7. Data are from n = 5–10 mice per time point and representative of two independent experiments, except in G, where a representative image from n = 6 mice per group across two independent repeats is shown. *, P < 0.05; and **, P < 0.01.
Figure S2. **Bcl6 deficiency in T cells during IAV primary infection and secondary infection reduces antibody responses, but does not affect weight loss.** 7- to 8-wk-old WT (Bcl6fl/fl) or TFH-deficient KO (Cd4cre Bcl6fl/fl) mice were infected with 10⁶ ffu RSV A2 i.n. (A and B) Virus-specific CD4 and CD8 T cells were stimulated ex vivo with RSV peptides to induce cytokine production and analyzed by flow cytometry. (C–F) 7- to 8-wk-old WT (Bcl6WT) or TFH-deficient KO (Cd4cre Bcl6WT) mice were infected with 40 PFU IAV PR8 and weighed daily to monitor disease severity. (D) TFH and GC B cell populations were analyzed by flow cytometry. (E) IAV-specific Ig, IgG1, and IgG2c in the serum as determined by ELISA. (F) WT or TFH-deficient KO mice were rechallenged with 40 PFU IAV PR8 8 wk after primary challenge and weighed daily. Data are from n ≥ 5 mice per group and representative of two independent experiments. Data are shown with mean ± SEM. *, P < 0.05; and **, P < 0.01.
Figure S3.  **BCL6 deficiency in early life does not affect the acute immune response.** WT (Bcl6^{fl/fl}) or T_{FH}-deficient KO (Cd4cre Bcl6^{fl/fl}) mice were infected as neonates (7 d old) with 2.5 × 10^5 ffu RSV A2 i.n.  **(A)** Mice infected as neonates were weighed daily.  **(B and C)** Virus-specific T cell populations were analyzed in the lungs (B) and LN (C) of infected neonates at day 14 p.i.  **(D-F)** Mice were reinfected 8 wk later with 8 × 10^5 ffu RSV A2 i.n.  **(D)** Airway, lung, and LN cells before (day 0) and after (days 4 and 7) reinfection.  **(E)** Virus-specific T cell populations were analyzed by flow cytometry at all time points.  **(F)** Cytokine concentrations in the airways was measured by ELISA. Data are from n ≥ 4 mice per group and representative of two independent experiments. Data are shown with mean ± SEM.
Figure S4. IL-2 neutralization does not limit T reg cell numbers in early-life RSV infection but limits inflammation on reinfection as adults. Mice were infected with $2.5 \times 10^5$ ffu RSV A2 at 7 d of age and treated with anti-IFN-γ, anti-IL-2, or isotype control i.p.; mice were sacrificed at day 14 p.i. (A) Foxp3+ T reg cells were enumerated in the lung, LNs, and spleen. (B) The number and frequency of Foxp3+CXCR5+PD1+ CD4 T cells in the lung dLNs. (C) ICOS and Bcl6 expression by LN TFH was determined by flow cytometry, and representative histograms are shown. (D) Total airway, lung, and LN cell counts at day 7 after reinfection with $8 \times 10^5$ PFU RSV A2 i.n. in mice initially infected with $2.5 \times 10^5$ PFU RSV A2 i.n. as neonates. Data represent two independent experiments with $n = 8–12$ mice per group. Data are shown with mean ± SEM. **, P < 0.01; and ***, P < 0.001.
Figure S5. Assessment of IL-2 and IL-6 production and signaling in secondary lymphoid tissue. (A) Neonatal LN TFH and non-TFH, as shown in Fig. 7 A, were assessed for expression of PSGL1 and Ly6C. (B) Expression of IL6 and IL2 relative to GAPDH in the spleen of 1-, 2-, and 8-wk-old uninfected C57B/L6 mice was determined by RT-qPCR. (C) IL-2 was measured by flow cytometry in different cell populations from lung dLNs of neonatal C57B/L6 at day 3 after RSV infection. A gating strategy for the cell types and a representative IL-2 histogram following either polyclonal (PMA/I and brefeldin A [BFA]) or unstimulated (brefeldin A alone) are shown. (D) CD25, CD122, and CD132 expression by splenic CD44−CD62L−Foxp3− and Foxp3+ CD4 T cells in 1- and 8-wk-old C57BL/6 mice. FMO, fluorescence minus one. (E) Percentage of CD25+ splenic CD44−CD62L−Foxp3− and Foxp3+ CD4 T cells in 1-wk-old C57BL/6 mice. Data in A and B are from n = 4–5 mice per group and are representative of two independent experiments. Data in D and E are from n = 6 mice and are representative of two independent experiments. (C) Representative gating strategy from an RSV-infected mouse; data are enumerated in Fig. 7, D and E. Data are shown with mean ± SEM. *, P < 0.05; and **, P < 0.01.