

ARTICLE

B cell–intrinsic TBK1 is essential for germinal center formation during infection and vaccination in mice

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The germinal center (GC) is a site where somatic hypermutation and clonal selection are coupled for antibody affinity maturation against infections. However, how GCs are formed and regulated is incompletely understood. Here, we identified an unexpected role of Tank-binding kinase-1 (TBK1) as a crucial B cell–intrinsic factor for GC formation. Using immunization and malaria infection models, we show that TBK1-deficient B cells failed to form GC despite normal Tfh cell differentiation, although some malaria-infected B cell–specific TBK1-deficient mice could survive by GC-independent mechanisms. Mechanistically, TBK1 phosphorylation elevates in B cells during GC differentiation and regulates the balance of IRF4/BCL6 expression by limiting CD40 and BCR activation through noncanonical NF- κ B and AKT^{T308} signaling. In the absence of TBK1, CD40 and BCR signaling synergistically enhanced IRF4 expression in Pre-GC, leading to BCL6 suppression, and therefore failed to form GCs. As a result, memory B cells generated from TBK1-deficient B cells fail to confer sterile immunity upon reinfection, suggesting that TBK1 determines B cell fate to promote long-lasting humoral immunity.

Introduction

Germinal centers (GCs) are highly dynamic structures formed in secondary and tertiary lymphoid organs and play a key role in humoral immune responses against pathogens and during vaccination. Induction of GCs allows antigen-activated B cells to be selected for the production of affinity-matured antibodies through the process of somatic hypermutation and clonal selection and then to exit GCs to become antigen-specific long-lived plasma cells and memory B cells. The anatomical structure and function of GCs have been studied for a long time (Victoria and Nussenzweig, 2012), but only recently has information been gathered about how B cells are guided through the development of GCs. Upon antigen encounter, B cells rapidly proliferate and differentiate into multipotent precursor cells that have the potential to diverge into two distinct fates: either GC or extra-follicular (EF) responses (Taylor et al., 2012). Collective studies suggested that the combination of multiple factors, including BCR, costimulatory receptors, and cytokines, leads to differential

transcriptional regulation that determines the fate of antigen-activated B cells to commit into GCs (Cyster and Allen, 2019; Song and Matthias, 2018). Induction of the GC reaction is often the desirable outcome of an infection or vaccine to provide antigen-specific long-lasting humoral immunity. Conversely, the induction of GC-independent EF responses in several disease models and vaccinations results in early acute production of short-lived plasmablasts (Elsner and Shlomchik, 2020). The intrinsic and extrinsic factors that determine the fate of antigen-activated B cells, whether to enter GC reactions or to commit to EF responses, are not yet well defined.

B cells recognize antigens through the BCR and migrate to the B–T cell border to present the antigen to cognate CD4⁺ T cells with the engagement of costimulatory molecules, such as CD40L, CD80, CD86, and inducible T cell costimulator (ICOS) ligand (ICOSL), to drive the differentiation of T follicular helper (Tfh) cells. The interaction between antigen-activated B cells

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and cognate Tfh cells, through physical interaction and cytokines, leads to maturation of Tfh cells and GC formation. The strength of BCR affinity and T cell help through CD40 signaling has been demonstrated to regulate GC initiation and determine the fate of GC B cells (Shinnakasu and Kurosaki, 2017; Laidlaw and Cyster, 2021). GC B cells, with intermediate BCR affinity and CD40 help, tend to differentiate into memory B cells, while high BCR affinity with strong CD40 signaling drives B cells to differentiate into plasma cells fate. Recent studies also showed that CD40 and BCR signaling is altered as naive B cells differentiate into GC B cells to regulate clonal selection in the light zone (Luo et al., 2019, 2018). However, the B cell-intrinsic factors and molecular mechanisms that regulate the dynamics of CD40 and BCR signaling to initiate GC differentiation are not fully understood.

TANK-binding kinase-1 (TBK1) is a serine/threonine kinase that is known for its role in innate immunity, especially for type I IFN (IFN-I) production upon nucleic acid recognition (Hemmi et al., 2004; Ishii et al., 2008). Apart from its role in IFN-I production via phosphorylation of IFN regulatory factor 3/7 (IRF3/7), TBK1 is also involved in various signaling pathways in a cell type-specific manner (Xiao et al., 2017; Jin et al., 2012; Zhao et al., 2018) and in adaptive immunity (Jin et al., 2012; Yu et al., 2015; Pedros et al., 2016; Marichal et al., 2011). Importantly, silencing *Tbk1* expression in CD4 T cells using shRNA suggested the role of TBK1 in Tfh cell maturation through ICOS signaling (Pedros et al., 2016), while TBK1 signaling in B cells was shown to negatively regulate IgA antibody class switching (Jin et al., 2012). Moreover, *Tbk1* is one of the genes up-regulated in GC B cells upon 4-hydroxy-3-nitrophenylacetyl (NP)-CGG immunization (Kaji et al., 2012), suggesting that TBK1 may have a role in GC formation. However, whether TBK1 plays a critical role in Tfh-B cell communications remains unclear.

Unlike most infectious diseases against which long-lasting protection is conferred through induction of immune memory after recovery from primary infection, malaria is exceptional. Human malaria infection only induces partial immunity after subsequent repeated infections (Portugal et al., 2013). An individual remains susceptible to repeated malaria infection despite gaining partial immunity that protects from severe complications (Coban et al., 2018). How the GC is regulated during malaria infection and impairs immune memory that confers sterile protection is not fully understood (Akkaya et al., 2020b). Both host and parasite factors are thought to contribute to the development or suppression of protective humoral immunity (Saito et al., 2017; Akkaya et al., 2020a).

Here, using both malaria infection and immunization models, we identified B cell-intrinsic TBK1 as an important molecule involved in regulating GC differentiation. We found that phosphorylation of TBK1 increased in B cells during GC differentiation. Notably, *Tbk1* deletion in B cells led to impaired GC formation and therefore increased susceptibility to a nonlethal malaria infection. Memory B cells generated in the absence of TBK1-dependent GC failed to confer sterile immunity upon reinfection challenge. Although TBK1-deficient B cells could be activated to differentiate into GC precursor cells (Pre-GC), the fate of these cells was distinct from that of TBK1-sufficient

B cells. We found that TBK1 negatively regulates both CD40 and BCR signaling in Pre-GC. Mechanistically, increased CD40-induced RelB expression in TBK1-deficient Pre-GC enhanced IRF4 expression that subsequently suppressed B-cell lymphoma 6 (*Bcl6*) expression, the master regulator of GC formation. Unlike naive B cells, BCR activation further enhanced CD40-induced IRF4 expression in Pre-GC in a TBK1-dependent manner. This study revealed a crucial role of B cell-intrinsic TBK1 as a determinant of GC commitment through the fine tuning of CD40 and BCR signaling.

Results

TBK1 in B cells is indispensable for GC formation

Murine malaria infection with *Plasmodium yoelii* nonlethal (PyNL) has been shown to induce robust Tfh cell-dependent GC formation (Arroyo and Pepper, 2020; Pérez-Mazliah et al., 2015; Fig. S1, A and B). In fact, Tfh cell-deficient mice (*Cd4^{cre}Bcl6^{fl/fl}*) infected with PyNL could not generate Tfh cells and form GCs and thus could not recover from the infection, suggesting GC-dependent humoral immunity is responsible for the self-recovery from PyNL infection (Fig. S1, C-E). To investigate how TBK1 plays a critical role in Tfh-B cell communications, we first investigated the role of TBK1 at distinct stages of GC differentiation from naive B cells to B220⁺ IgD⁻ CD38⁻ GL7⁺ Pre-GC and B220⁺ IgD⁻ CD38⁻ GL7⁺ mature GC B cells. To examine the phosphorylation activity of TBK1 at different stages of GC differentiation, we purified splenic naive B cells, Pre-GC, and mature GC B cells on day 0, day 9, and day 30 after PyNL infection, respectively, and performed Western blot analysis. We found that TBK1 was constitutively phosphorylated in naive B cells from mouse (Fig. 1 A) and human peripheral blood mononuclear cells (PBMCs; Fig. S2 A). Interestingly, both TBK1 protein expression and TBK1 phosphorylation increased in Pre-GC and GC B cells compared with naive B cells (Fig. 1 A).

Next, to investigate the role of TBK1 in GC formation, B cell-intrinsic TBK1-deficient mice (*Mbl^{cre} Tbk1^{fl/fl}* mice [*B Tbk1^{-/-}*]) were generated (Fig. S2 B). The deletion of *Tbk1* in B cells did not impair normal B cell development in the spleen, peritoneum, and bone marrow (Fig. S2, C and D). However, TBK1 deficiency in B cells led to impaired GC formation (Fig. 1, B-D) and significantly poor recovery from PyNL infection; eventually, 60% of the mice succumbed to death (Fig. 1, E and F). To address whether the involvement of TBK1 in GC formation is limited to malaria infection, we immunized mice with 4-hydroxy-5-iodo-3-nitrophenyl (NIP)-OVA with alum. After immunization, TBK1-deficient B cells, similarly to infection, were found to have severely impaired GC formation in the draining LNs and spleens (Fig. 1, G and H).

However, despite the lack of GC formation in *B Tbk1^{-/-}* mice, robust class-switched antibody responses were still detected in the serum upon either malaria infection or immunization, albeit significantly less and with lower IgG1 antibody avidity (Fig. S3, A-C). In addition, the absence of GC in those *B Tbk1^{-/-}* mice did not affect IgG class switching, which is in line with the fact that class-switch recombination happens before B cells enter GC reactions (Roco et al., 2019).

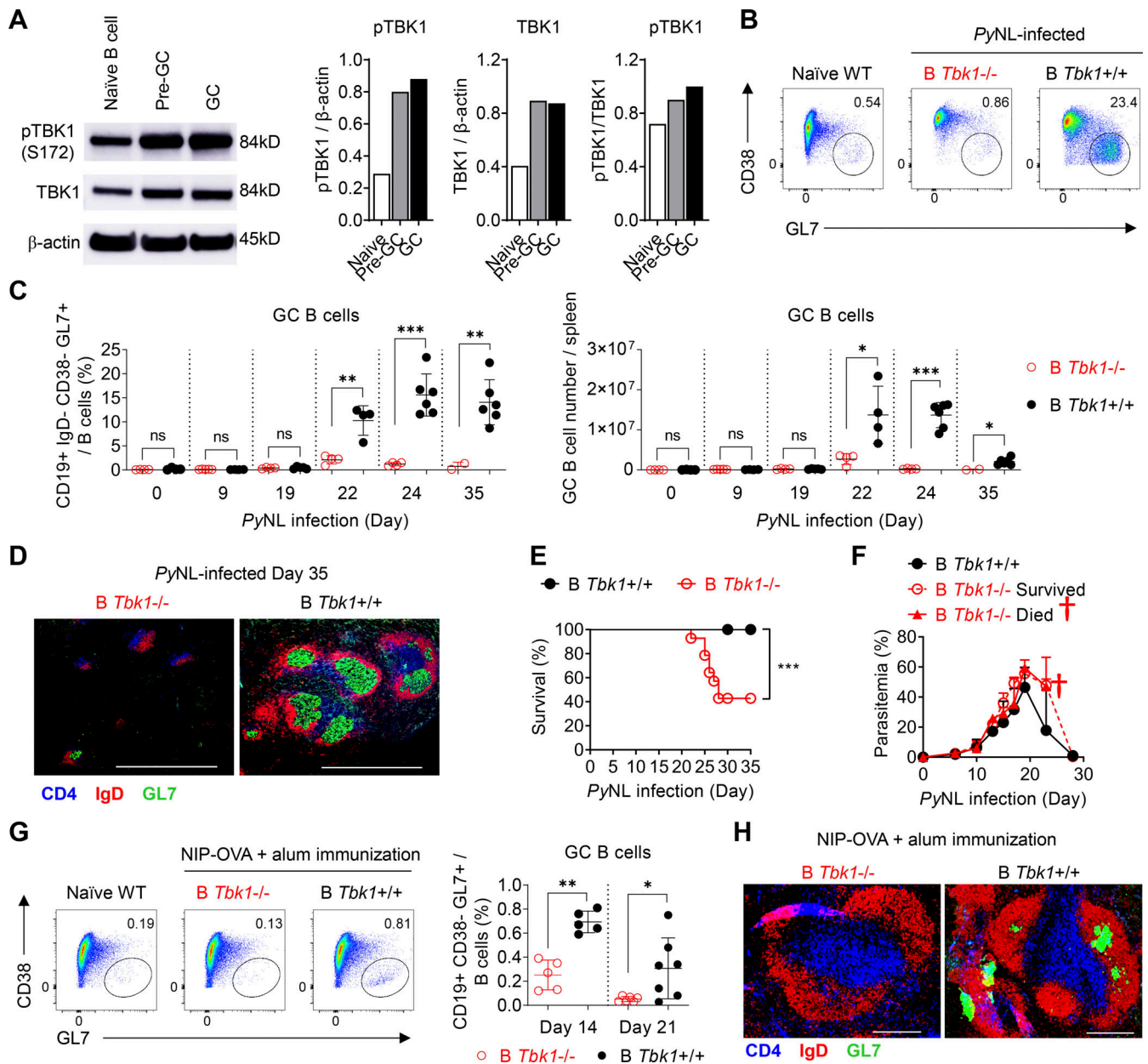


Figure 1. TBK1 is involved in GC B cell differentiation. (A) Protein expression and phosphorylation of TBK1 in purified naive B cells from naive mice, Pre-GC B cells from day 9 PyNL-infected mice, and GC B cells from day 30 PyNL-infected mice, pooled from two mice per group. Graph shows the quantitative band intensity normalized to actin control or total TBK1 protein. (B) Representative FACS analysis of GC B cell population in the spleen on day 24 after PyNL infection. Plots are gated on CD19⁺ IgD⁻ cells. (C) Proportions and cell numbers of the GC B cell population in the spleen of PyNL-infected mice over time. *n* = 4–6 mice/group at each time point from day 0 to day 24; *n* = 2–6 mice/group at day 35. (D) Immunohistochemical analysis of spleens from *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice that recovered from PyNL infection on day 35 after infection. Staining of CD4 (blue), expressed in CD4 T cells; IgD (red), expressed in mature follicular B cells; GL7 (green), expressed in the GC. Scale bars, 1 mm. (E) Survival of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice upon PyNL infection. *n* = 11–14 mice/group. (F) Parasitemia of *B Tbk1*^{-/-} mice that died, *n* = 8; *B Tbk1*^{-/-} mice that survived, *n* = 6; and *B Tbk1*^{+/+} mice, *n* = 11; after infection with PyNL. †, time point of death. (G) Representative FACS plots and percentages of splenic GC B cell population among total B cells 14 d and 21 d after immunization with NIP-OVA with alum adjuvant. *n* = 5–7 mice/group. (H) Immunohistochemical analysis of spleens from *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice 21 d after immunization. Scale bars, 200 μm. Each dot represents an individual mouse. Data are representative of two (A) or three (G) independent experiments and pooled from five (C) or three (E and F) independent experiments, representative of at least two (D) or three (H) biological replicates. Data are shown as mean ± SD. *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001; unpaired Student's *t* test (C), Mann-Whitney *t* test (G), or log-rank (Mantel-Cox) test (E). Source data are available for this figure: SourceData F1.

Although B cell-intrinsic TBK1 was not required for antibody-secreting cell (ASC) formation (Fig. S3 D), the lack of GC in *B Tbk1*^{-/-} mice suggests that those ASCs in *B Tbk1*^{-/-} mice were of GC-independent origin and therefore had a

lower capacity of antibody production. Collectively, these data suggested that B cell-intrinsic TBK1 is absolutely required for GC formation but is dispensable for ASC differentiation.

Tfh cell formation is independent of B cell-intrinsic TBK1

GC formation relies on cytokines and costimulatory signals from cognate Tfh cells (Nutt and Tarlinton, 2011). Interaction between antigen-activated B cells and cognate CD4⁺ T cells at the border of the B cell follicle drives Tfh cell differentiation. Tfh cells then signal cognate activated B cells to further differentiate into GC B cells, while signals from GC B cells are important to sustain Tfh cell maintenance. We therefore sought to determine whether the defect in GC formation in the absence of TBK1 in B cells was due to impaired B cell signals to drive Tfh cell formation. Surprisingly, the population of Tfh cells in B *Tbkl*^{-/-} mice during PyNL infection was comparable to the WT counterparts in a time-course analysis up to day 22, despite the lack of GC in B *Tbkl*^{-/-} at this time point (Fig. 2 A). However, the Tfh cell population was unsustainable at a later time point on day 24 in B *Tbkl*^{-/-} mice (Fig. 2 A). To address the functionality of these Tfh cells, we further examined the expression of BCL6, the master regulator of Tfh cells, IL-21, and ICOS, which determine the differentiation and functionality of Tfh cells (Nurieva et al., 2009; Vogelzang et al., 2008), on day 24 after infection. While the expression of BCL6 and IL-21 increased as T cells differentiated and matured into a Tfh cell population (Fig. 2 B), Tfh cells induced by *Tbkl*-deficient B cells expressed BCL6 and IL-21 (Fig. 2 C) comparable to that in WT B cells, suggesting *Tbkl* deficiency in B cells did not impair cognate T cell priming for Tfh cell differentiation. Furthermore, ICOS expression on activated CD4⁺ T cells (CXCR5⁻ PD1⁺), pre-Tfh cells (CXCR5^{lo} PD1⁺), and mature Tfh cells (CXCR5^{hi} PD1⁺) was not suppressed in B *Tbkl*^{-/-} mice (Fig. 2 C), suggesting that *Tbkl*-deficient B cells were able to interact with and activate CD4⁺ T cells. The induction of BCL6, IL-21, and ICOS expression in Tfh cells in B *Tbkl*^{-/-} mice suggests that the lack of GC in B *Tbkl*^{-/-} mice was not due to impaired Tfh cell formation. The T follicular regulatory (Tfr) cell is a new player that limits GC formation by suppressing Tfh cell function (Wing et al., 2017). However, the populations of Tfr cells were comparable between B *Tbkl*^{-/-} and the WT counterpart (Fig. 2 D), suggesting the lack of sustainable Tfh cells was not due to inhibition by Foxp3⁺ Tfr cells. The maintenance of Tfh cells is known to be dependent on constant interaction with GC B cells. Therefore, we investigated the costimulatory molecule expression on B cells. We found that ICOSL and CD86 expression on B cells was significantly reduced in B *Tbkl*^{-/-} mice on day 24 after infection (Fig. 2 E), suggesting that TBK1 sustains B-T cell interaction for Tfh cell maintenance. The increase of ICOS expression on Tfh cells in PyNL-infected B *Tbkl*^{-/-} mice was likely due to the higher parasitemia, because immunization-induced up-regulation of ICOS expression on Tfh cells was not higher in B *Tbkl*^{-/-} mice than in the WT mice (Fig. 2 F), indicating the suppressed ICOSL on B cell was not indirectly affected by ICOS expression on Tfh cells.

To understand whether the impaired GC formation in B *Tbkl*^{-/-} mice was due to an unsustainable Tfh cell population, we immunized mixed bone marrow chimeric mice containing 50:50 *Tbkl*^{-/-} and WT B cell populations (Fig. 2 G). The supposed Tfh cell population can at least be maintained by WT B cells within the same mouse; yet, *Tbkl*^{-/-} B cells failed to effectively differentiate into GC B cells (Fig. 2, H and I), indicating that TBK1

controls B cell-intrinsic signaling for GC B cell differentiation. Consistently, we also observed a marked reduction of ICOSL expression on the small population of *Tbkl*^{-/-} GC B cells formed in the chimeric mice. Taken together, all data suggest that *Tbkl*-deficient B cells were able to activate cognate CD4 T cells and initiate Tfh cell differentiation; however, in return, the lack of GC formation could not sustain Tfh cell maintenance in the long term due to unsustainable ICOS-ICOSL interaction.

CD4⁺ T cell-intrinsic TBK1 is dispensable for Tfh cell maturation

We next sought to directly investigate whether CD4⁺ T cell-intrinsic TBK1 is required for Tfh cell maturation and GC formation, because TBK1 was reported to be involved in the maturation of Tfh cells using adoptive transfer of CD4⁺ T cells transduced with shRNA to silence the *Tbkl* gene (Pedros et al., 2016). We generated *Cd4*^{cre} *Tbkl*^{fllox} (T *Tbkl*^{-/-}) mice in which *Tbkl* was specifically deleted in CD4⁺ T cells (Fig. 3 A). We found that *Tbkl*-deficient CD4⁺ T cells could differentiate into mature Tfh cells with BCL6, IL-21, and ICOS expression comparable to that of the WT counterparts in both PyNL infection and immunization models (Fig. 3, B–D). We further examined the functionality of these Tfh cells by evaluating the ability of *Tbkl*-deficient Tfh cells to induce GC formation in both PyNL infection and immunization models. Surprisingly, unlike the previous report that *Tbkl* knockdown in CD4⁺ T cells was unable to induce GC formation (Pedros et al., 2016), we could detect comparable GC B cell populations in T *Tbkl*^{-/-} mice and the WT counterparts (Fig. 3, E and F). Overall, collectively, these data show that B cell-intrinsic but not CD4⁺ T cell-intrinsic TBK1 plays an important role in the regulation of GC formation.

TBK1 determines B cell fate by controlling the transcriptional profile of Pre-GC

As TBK1 phosphorylation increased in CD38⁺ GL7⁺ Pre-GC cells compared with naive B cells (Fig. 1 A), we examined whether TBK1 is essential for B cells to proliferate and differentiate into a Pre-GC population. There was no impaired proliferation of activated B cells, as malaria-activated B cells from B *Tbkl*^{-/-} mice had equal proliferation capacity in response to CD40 or IgM stimulation as the B *Tbkl*^{+/+} mice (Fig. 4 A). Moreover, a substantial amount of the Pre-GC population could be detected in the spleen on day 9 after PyNL infection, regardless of the presence of TBK1 in B cells, albeit less in B *Tbkl*^{-/-} mice (Fig. 4 B), with significantly reduced IgG1 class switching (Fig. 4 C). To determine whether the lack of GC formation was due to impaired Pre-GC proliferation, we performed an in vivo 5-ethynyl-2'-deoxyuridine (EdU) assay; however, we did not find differences in cell cycle and proliferation between *Tbkl*^{-/-} and *Tbkl*^{+/+} Pre-GC (Fig. 4 D), nor did cell death increase in Pre-GC due to *Tbkl* deficiency (Fig. 4 E).

Next, to identify the TBK1-mediated factors that determine differentiation of Pre-GC into mature GC, we sorted Pre-GC populations from B *Tbkl*^{-/-} and B *Tbkl*^{+/+} mice for RNA sequencing (RNA-seq). Gene set enrichment analysis (GSEA; Subramanian et al., 2005) using hallmark gene sets from the Molecular Signatures Database (MSigDB) revealed that gene signatures highly enriched in *Tbkl*^{-/-} Pre-GC were related to Myc target genes and NF-κB signaling (Fig. 4 F), which are

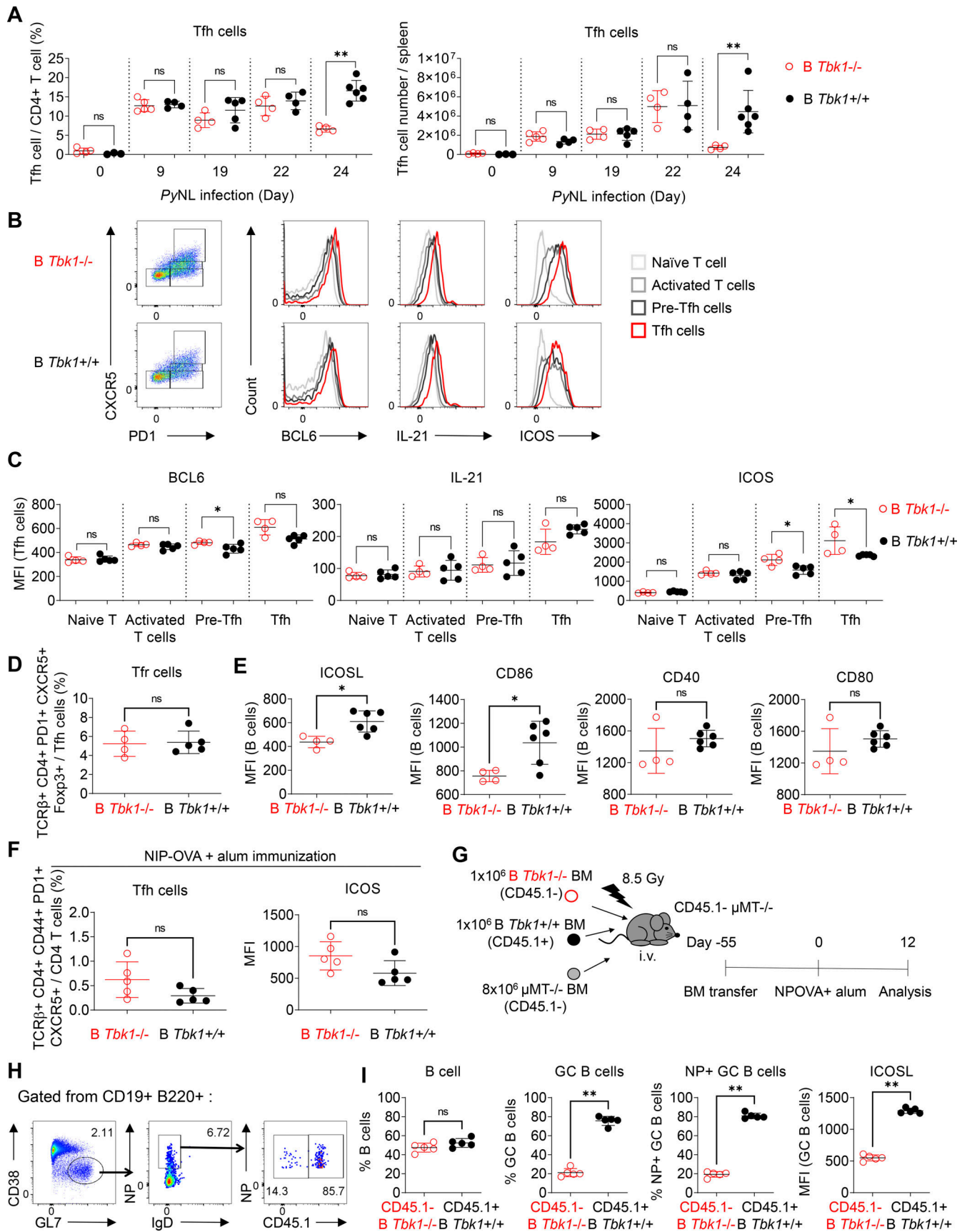


Figure 2. **TBK1-deficient B cells drive Tfh cell differentiation despite the lack of GC B cells.** (A) Time-course analysis of Tfh cell proportions and cell numbers in the spleens of B *Tbk1*^{-/-} and B *Tbk1*^{+/+} mice infected with PyNL analyzed by FACS. *n* = 4–6 mice/group. (B) Representative FACS plots and

histograms of BCL6, IL-21, and ICOS expression in naive T cells (TCR β^+ CD4 $^+$ PD1 $^-$ CXCR5 $^-$), activated T cells (TCR β^+ CD4 $^+$ PD1 $^+$ CXCR5 $^-$), pre-Tfh cells (TCR β^+ CD4 $^+$ PD1 $^+$ CXCR5 lo), and mature Tfh cells (TCR β^+ CD4 $^+$ PD1 $^+$ CXCR5 hi) in the spleens of B *Tbkl* $^{-/-}$ and B *Tbkl* $^{+/+}$ mice on day 22 after infection with PyNL. **(C)** Mean fluorescence intensity (MFI) of BCL6, IL-21, and ICOS expression in subsets of T cells in the spleens of B *Tbkl* $^{-/-}$ and B *Tbkl* $^{+/+}$ mice on day 24 after infection with PyNL. $n = 4-5$ mice/group from two independent experiments. **(D)** Tfr population in the spleens of B *Tbkl* $^{-/-}$ and B *Tbkl* $^{+/+}$ mice on day 24 after infection with PyNL. $n = 4-5$ mice/group. **(E)** MFI of ICOSL, CD86, CD40, and CD80 expression on B cells on day 24 after infection with PyNL. $n = 4-6$ mice/group. **(F)** Tfh cell population and ICOS expression on Tfh cells in the dLN on day 12 after immunization with NIP-OVA and alum. $n = 5$ mice/group. **(G-I)** Mixed bone marrow (BM) chimeric mice were immunized with NIP-OVA and alum. Schematic diagram of the experimental protocol (G), flow cytometry gating strategy (H), flow cytometry analysis of CD45.1 $^-$ B *Tbkl* $^{-/-}$ B cells and CD45.1 $^+$ WT B cells (I). $n = 5$ mice. Representative of two independent experiments. Each dot represents an individual mouse. Data are from four experiments (A) or are representative of two experiments (B-I). Data are shown as mean \pm SD. *, $P < 0.05$; **, $P < 0.01$; Mann-Whitney t test (A, C-F, and I).

signaling downstream of BCR and CD40 activation. Of the 588 significantly up-regulated genes and 557 down-regulated genes in B *Tbkl* $^{-/-}$ Pre-GC cells compared with B *Tbkl* $^{+/+}$ Pre-GC, we focused on those that were up-regulated or down-regulated by 1.33-fold (adjusted $P < 0.05$), which were 131 genes and 147 genes, respectively (Fig. 4 G). Gene ontology (GO) analysis for biological process consistently showed that most up-regulated genes were related to regulation of signal transduction, including *Nfkbid*, *Nfkbia*, *Ebi3*, *Irf4*, and *Myc*, which are associated with CD40 and BCR signaling (Fig. 4 H and Fig. S4 A). Therefore, we further analyzed gene signatures of BCR and CD40 signaling (Victora et al., 2010), and we found significant up-regulation of CD40- and BCR-induced genes in *Tbkl*-deficient Pre-GC (Fig. 4 I and Fig. S4 B), suggesting that TBK1 acts as a negative regulator of CD40 and BCR signaling in Pre-GC. A *Myc* $^+$ Pre-GC-specific gene signature (Calado et al., 2012) further confirmed that TBK1 negatively regulates *Myc*-related genes (Fig. 4 I). Consistent with the phenotype of previously reported *Myc* $^+$ Pre-GC (Calado et al., 2012), we found that *Tbkl*-deficient Pre-GC had increased *Nfkbia* and *Nfkbid* expression indicative of enhanced NF- κ B activation. Notably, enhanced *Myc* and *Irf4* expression in Pre-GC was shown to suppress GC-related genes (Dominguez-Sola et al., 2012). Collectively, these findings suggest that TBK1 regulates *Myc* and NF- κ B signaling via CD40 and BCR signaling for GC fate decision.

TBK1 negatively regulates IRF4 in Pre-GC

IRF4 is an early transient transcription factor induced by CD40 signaling that is essential for the transcriptional regulation of GC genes such as *Bcl6* (Ochiai et al., 2013). However, IRF4 has been shown to act as a double-edged sword with opposing roles in the regulation of GC differentiation, depending on its expression levels at different stages of GC differentiation (Zhang et al., 2017). Although IRF4 drives the expression of *Bcl6*, sustained up-regulation of IRF4 expression in turn suppresses the transcription of *Bcl6* (Saito et al., 2007). Consistent with our RNA-seq data (Table S1), we additionally found that *Tbkl*-deficient Pre-GC had increased expression of IRF4 and decreased BCL6 by flow cytometric analysis (Fig. 5, A and B). *Tbkl*-deficient B cells preferentially differentiated into a IRF4 $^+$ Bcl6 $^-$ population, in contrast to the increase of the Bcl6 $^+$ IRF4 $^-$ Pre-GC population in B *Tbkl* $^{+/+}$ mice (Fig. 5 C). The decreased ratio of BCL6/IRF4 expression in Pre-GC (Fig. 5 D) indicates that *Tbkl*-deficient Pre-GC has a lower potential to further differentiate into mature GC B cells. Indeed, the population of BCL6 $^+$ B cells was consistently suppressed in B *Tbkl* $^{-/-}$ mice, while BCL6 $^+$ B cells expanded

ninefold and differentiated into GC B cell in B *Tbkl* $^{+/+}$ mice on day 24 after infection (Fig. 5 E). The Pre-GC population has been shown to have the potential to either differentiate into GC or GC-independent plasmablast/memory B cells (Taylor et al., 2012). Notably, IRF4 is also a transcription factor that drives the gene expression of *Prdm1* that encodes BLIMP-1, the master regulator of plasmablast differentiation (Klein et al., 2006). Our RNA-seq data also showed that *Prdm1* gene expression was significantly up-regulated in *Tbkl*-deficient Pre-GC (Table S1), indicating that the differentiation commitment of this population was skewed to a plasmablast fate instead of a GC B cell fate. Consistently, with the increased *Prdm1* expression in *Tbkl*-deficient Pre-GC, the population of CD138 $^+$ ASCs was significantly higher in B *Tbkl* $^{-/-}$ mice at the acute phase of infection (Fig. 5 F). These findings strongly support that TBK1 dictates the fate of the Pre-GC population to predominantly differentiate into GC by controlling the expression of IRF4.

TBK1 limits the activation of CD40 signaling in B cells via TNF receptor-associated factor 2 (TRAF2)

CD40 stimulation of B cells induces both canonical and non-canonical NF- κ B signaling with differential roles in B cell survival, proliferation, isotype switching, and differentiation fate (Zarnegar et al., 2004; Heise et al., 2014). IRF4 is a downstream target of NF- κ B upon CD40 stimulation in B cells (De Silva et al., 2016). Indeed, we found that CD40 stimulation, but not BCR stimulation, strongly induced IRF4 expression in naive B cells (Fig. 5 G). Furthermore, noncanonical NF- κ B signaling was enhanced upon CD40 stimulation (Fig. S5). To examine whether TBK1 modulates IRF4 expression via NF- κ B in Pre-GC, we examined the expression of noncanonical NF- κ B molecules RelB and p52 in a Pre-GC cell population in B *Tbkl* $^{-/-}$ and B *Tbkl* $^{+/+}$ mice. We found that the expression of RelB and p52 was significantly higher in *Tbkl*-deficient Pre-GC than in the WT Pre-GC (Fig. 5 H). Collectively, our findings suggest that TBK1 negatively regulates CD40 signaling via noncanonical NF- κ B that skews the balance of IRF4/BCL6 to promote GC formation.

We next sought the role of CD40 stimulation on the induction of TBK1 phosphorylation; therefore, we stimulated purified naive B cells with anti-CD40 and then performed Western blot analysis. To our surprise, we found that CD40 stimulation down-regulated TBK1 phosphorylation as early as 10 min after stimulation (Fig. 5 I). This prompted us to hypothesize that TBK1 limits the activation of CD40 signaling in steady state to prevent CD40 overactivation spontaneously in the absence of CD40L and that a reduced phosphorylation level of TBK1, but not complete

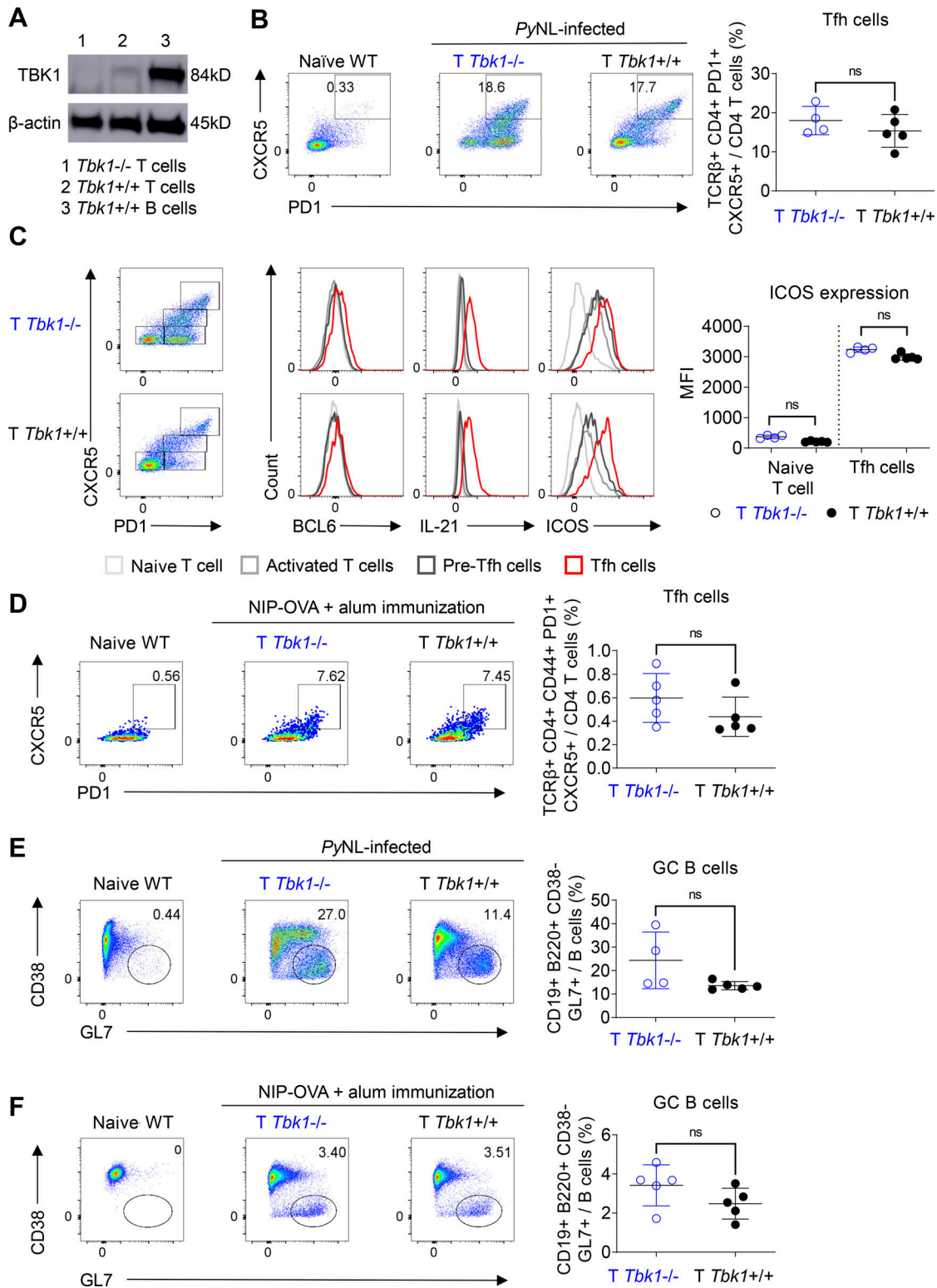
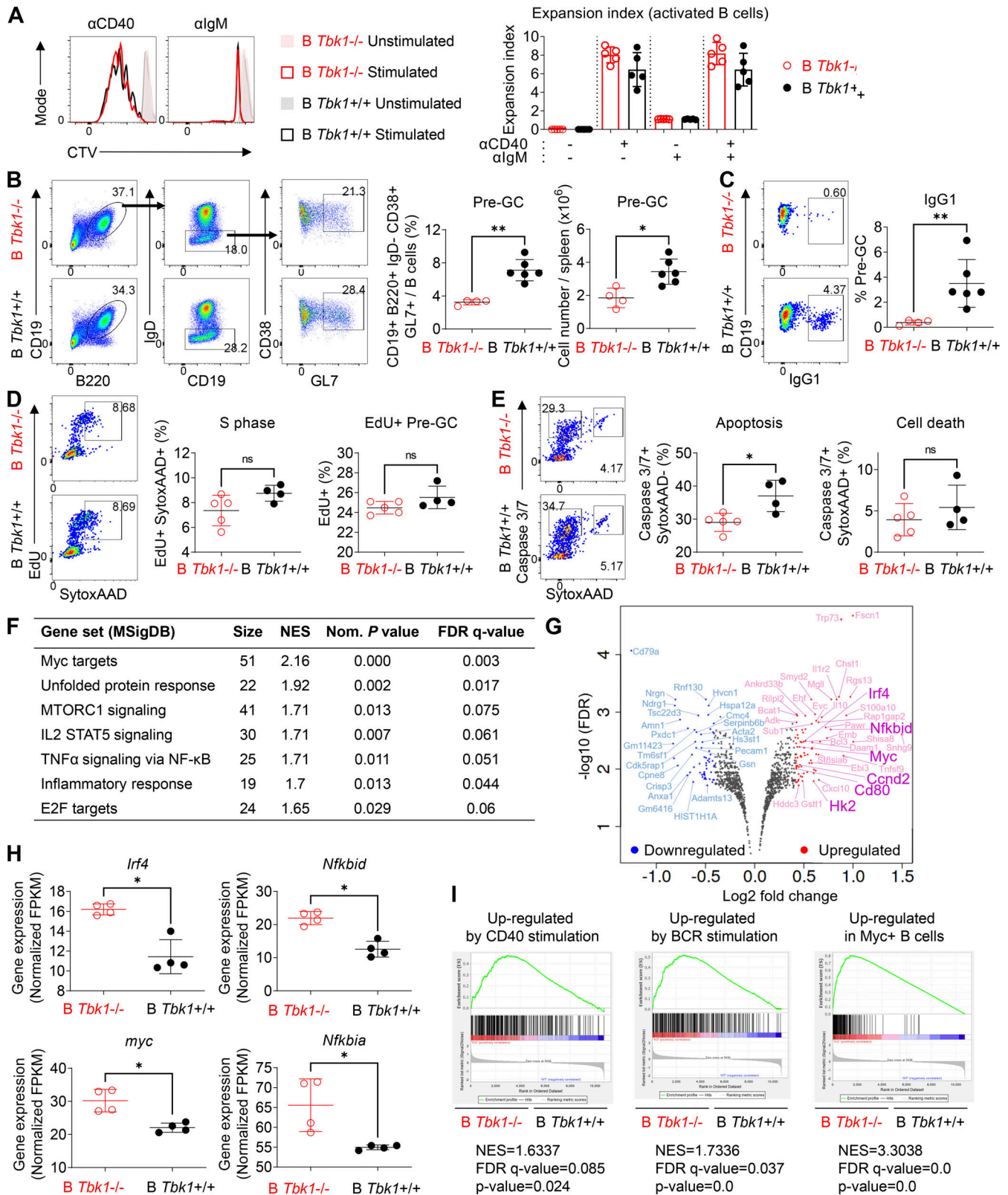


Figure 3. CD4⁺ T cell-intrinsic TBK1 is dispensable for Tfh cell maturation. (A) Depletion efficiency of *Tbk1* in splenic CD4⁺ T cells of *Cd4^{cre} Tbk1^{fl/fl}* mice (*Tbk1*^{-/-}) and *Cd4^{wt} Tbk1^{fl/fl}* mice (*Tbk1*^{+/+}) was analyzed by immunoblotting. B cells from WT mice were used as a control. (B) Population of Tfh cells in the spleens of *Tbk1*^{-/-} and *Tbk1*^{+/+} mice on day 24 after infection with PyNL analyzed by FACS. *n* = 4–5 mice/group. (C) Representative FACS plots and histograms of BCL6, IL-21, and ICOS expression in naive T cells, activated T cells, pre-Tfh cells, and mature Tfh cells in the spleens of *Tbk1*^{-/-} and *Tbk1*^{+/+} mice on day 24 after infection with PyNL. *n* = 4–5 mice/group. MFI, mean fluorescence intensity. (D) Population of Tfh cells in the spleens of *Tbk1*^{-/-} and *Tbk1*^{+/+} mice on day 12 after immunization with NIP-OVA with alum. *n* = 5 mice/group. (E) GC population in the spleens of *Tbk1*^{-/-} and *Tbk1*^{+/+} mice on day 24 after infection with PyNL. *n* = 4–5 mice/group. (F) GC population in the spleens of *Tbk1*^{-/-} and *Tbk1*^{+/+} mice on day 12 after immunization with NIP-OVA with alum. *N* = 5 mice/group. Each dot represents an individual mouse. Data are representative of two independent experiments (A–F). Data are shown as mean \pm SD. Mann-Whitney *t* test (B–F). Source data are available for this figure: SourceData F3.



sets enriched for DEGs between B *Tbk1*^{-/-} Pre-GC and B *Tbk1*^{+/+} Pre-GC analyzed by GSEA using gene signatures from MSigDB Hallmark (false discovery rate [FDR] q value <0.25%; $P < 0.05$). NES, normalized enrichment score. **(G)** Volcano plot of DEGs (1.33-fold; $P < 0.05$) between B *Tbk1*^{-/-} Pre-GC and B *Tbk1*^{+/+} Pre-GC isolated from day 9 after infection with PyNL analyzed by RNA-seq. $n = 4$ mice/group. **(H)** Gene expression of *Irf4*, *myc*, *Nfkbid*, and *Nfkbia* presented as fragments per kilobase of exon per million reads mapped (FPKM) values in RNA-seq of B *Tbk1*^{-/-} Pre-GC and B *Tbk1*^{+/+} Pre-GC. **(I)** Enrichment of genes up-regulated by CD40 and BCR and expressed in c-Myc⁺ B cells in B *Tbk1*^{-/-} Pre-GC compared with B *Tbk1*^{+/+} Pre-GC analyzed by GSEA. NES, normalized enrichment score. Each dot represents data from an individual mouse. Data are representative of two (A and C–E) or three independent experiments (B) or from one experiment (F–I). Data are shown as mean \pm SD. *, $P < 0.05$; **, $P < 0.01$; Mann-Whitney t test (A–E and H). AAD, SYTOX AADvanced.

abrogation of TBK1, may be required to allow optimal CD40 activation. TRAF2 is a molecule downstream of CD40 that regulates noncanonical NF- κ B. Therefore, we examined the phosphorylation of TRAF2 at serine 11 upon CD40 stimulation and found that phosphorylation of TRAF2 S11 was partially dependent on TBK1 (Fig. 5 I). Collectively, our findings suggest that TBK1 negatively regulates CD40 signaling to skew the balance of IRF4/BCL6 to promote GC formation via TRAF2 phosphorylation and noncanonical NF- κ B signaling.

TBK1 fine-tunes CD40 together with BCR signaling in Pre-GC

The alteration of CD40 and BCR signaling as naive B cells differentiate into GC suggests that fine tuning of CD40 and BCR signaling may be important to determine the differentiation fate of Pre-GC (Luo et al., 2018, 2019). Therefore, we sought to examine the role of TBK1 in the regulation of CD40 and BCR signaling, specifically in Pre-GC. Due to the difficulty of isolating the small population of Pre-GC for ex vivo stimulation, we stimulated splenocytes from PyNL-infected mice on day 9 and gated on the GL7⁺ B cells as Pre-GC and the GL7⁻ B cells as non-Pre-GC control (Fig. 6 A). Unlike in naive B cells (Fig. 5 G), BCR activation alone strongly up-regulated IRF4 expression in GL7⁺ Pre-GC, and it was further enhanced by CD40 costimulation (Fig. 6 B). We also found that the synergistic effect of BCR and CD40 stimulation on IRF4 is boosted in *Tbk1*-deficient GL7⁺ Pre-GC and GL7⁻ non-Pre-GC (Fig. 6 B). This finding supports our RNA-seq data showing that TBK1 negatively regulates IRF4 via both CD40 and BCR signaling in Pre-GC. Notably, IRF4 and c-Myc expression was slightly but significantly higher in unstimulated *Tbk1*-deficient Pre-GC and was enhanced by CD40 and BCR costimulation (Fig. 6 C). Although BCR stimulation is a strong inducer of c-Myc expression, we found that CD40 stimulation alone, but not BCR stimulation, strongly enhanced c-Myc expression in Pre-GC. CD40 stimulation further augmented c-Myc expression in *Tbk1*-deficient Pre-GC (Fig. 6 C). This finding indicates that the dynamics of CD40 and BCR signaling in Pre-GC were tightly regulated by TBK1.

AKT is a signaling molecule downstream of BCR activation that plays an important role in GC formation (Zhu et al., 2019). BCR activation in naive B cells induces AKT phosphorylation at the S473 hydrophobic motif by mammalian target of rapamycin complex 2 (mTORC2). However, strong AKT phosphorylation at the T308 activation loop by PDK1 in GC B cells was shown to inhibit AKT phosphorylation at S473 (Luo et al., 2019). TBK1 has been shown to phosphorylate AKT at S473 and T308 sites independent of mTORC2 and PDK1, depending on the stimulus and cell types (Ou et al., 2011; Xie et al., 2011); however, the involvement of TBK1 in AKT phosphorylation in Pre-GC B cells is unknown. We examined AKT phosphorylation in Pre-GC B cells

from PyNL-infected mice on day 9 and found that *Tbk1* deficiency significantly enhanced phosphorylated AKT (pAKT) at T308 but only slightly at S473 (Fig. 6 D). Collectively, these data suggest that TBK1 acts as a limiting factor to regulate CD40 and BCR signaling in Pre-GC through the regulation of c-Myc, IRF4, and AKT for GC B cell differentiation.

TBK1 is essential for long-lasting protective humoral immunity

While GC formation gives rise to affinity-matured memory B cells for long-term protection from infection, memory B cells could be generated in the absence of GC (Kaji et al., 2012; Taylor et al., 2012). To determine the effect of the lack of TBK1 in B cells on memory B cells, we treated both B *Tbk1*^{-/-} and WT mice with hydroxychloroquine (HCQ) antimalarial daily when they reached the peak parasitemia on day 15 after infection, and we reinfected them with PyNL after they had fully recovered from the primary infection (Fig. 7 A). We found that B *Tbk1*^{-/-} mice were fully susceptible to reinfection, while WT controls were protected (Fig. 7 B). We examined GC formation and the generation of affinity-matured memory B cells identified by CD73 expression, which correlates with somatic hypermutation (Anderson et al., 2007). While GC was not formed in B *Tbk1*^{-/-} mice (Fig. 7 C), CD73 expression was up-regulated on IgG1⁺ memory B cells in WT mice, but not in B *Tbk1*^{-/-} mice (Fig. 7 D). Moreover, B *Tbk1*^{-/-} mice had significantly fewer IgG1⁺ memory B cells (Fig. 7 E). This suggests that TBK1 is crucial for the generation of affinity-matured IgG1⁺ memory B cells through GC reaction. Notably, PyNL infection mainly induced IgG1⁺ GC B cells, while IgG2c⁺ GC B cells were hardly detectable (Fig. 7 F), suggesting that IgG1⁺ memory B cells were mostly derived from GC, while IgG2c⁺ memory B cells were of EF origin. Therefore, CD73 expression on IgG2c⁺ memory B cells was up-regulated in neither WT nor B *Tbk1*^{-/-} mice (Fig. 7 D). Even though B *Tbk1*^{-/-} mice had increased IgG2c⁺ memory B cells compared with WT mice and were capable of producing IgG2c and IgM antibodies upon reinfection (Fig. 7 G), B *Tbk1*^{-/-} mice were still susceptible to reinfection due to the lack of IgG1 and perhaps IgG3 antibodies, suggesting the importance of GC-derived memory B cells to produce affinity-matured antibodies. Taken together, these findings clearly reveal the importance of TBK1 for the generation of long-lasting and high-quality memory B cells through GC formation, which is important for sterile immunity.

Discussion

B cells rely on various extrinsic signals from T cells and follicular dendritic cells, as well as cell-intrinsic signals, to determine their fate of differentiation (Cyster and Allen, 2019). Here, we identified a novel role of B cell-intrinsic TBK1 as a determining factor

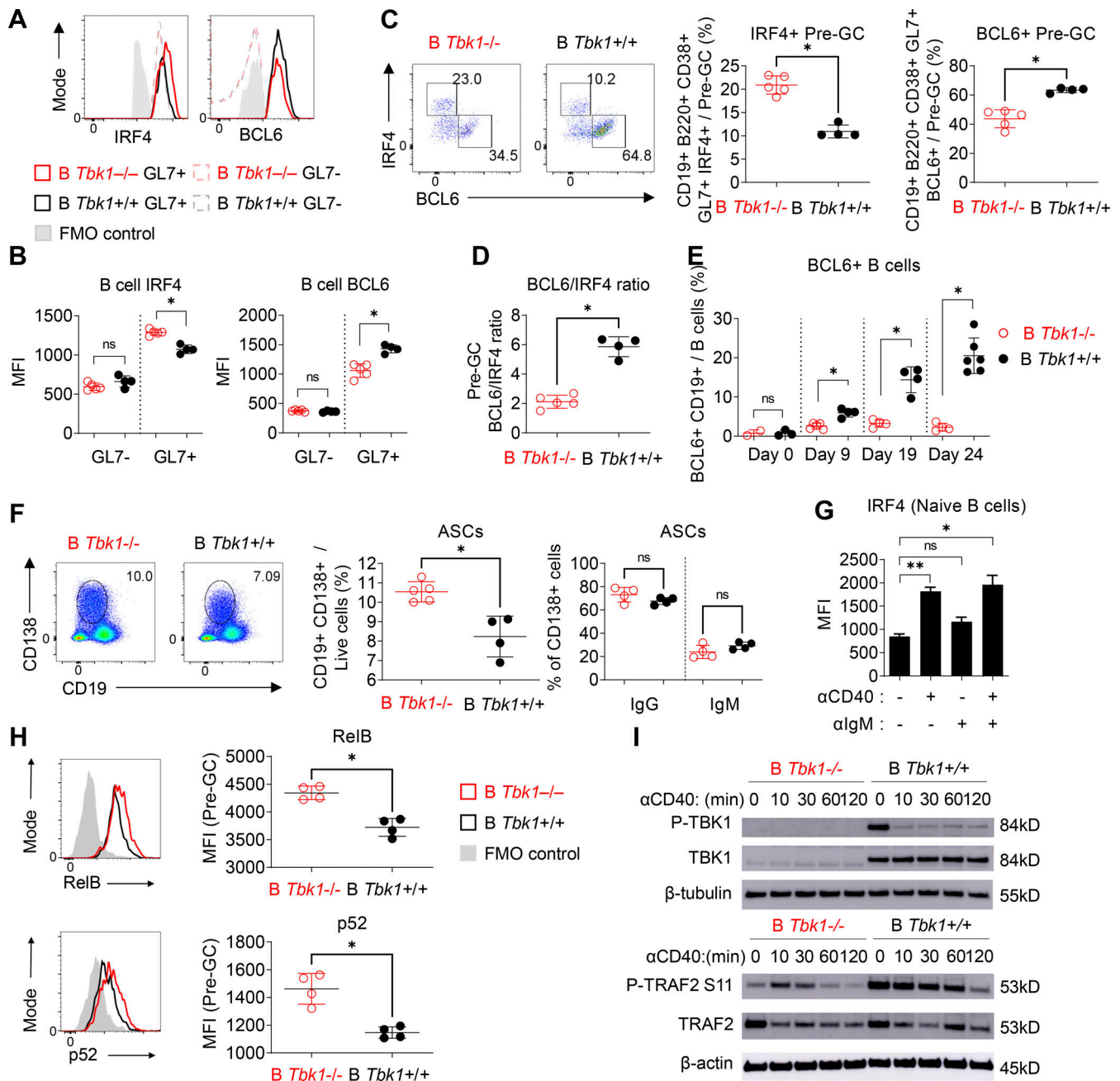


Figure 5. TBK1 negatively regulates IRF4 by limiting CD40 signaling in Pre-GC. (A and B) Representative histograms (A) and mean fluorescence intensity (MFI) of IRF4 and BCL6 expression (B) in GL7⁻ B cells and GL7⁺ Pre-GC of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 9 after infection with PyNL analyzed by FACS. FMO, fluorescence minus one control. n = 4–5 mice/group. **(C)** Representative FACS plots and proportion of IRF4⁺ Pre-GC and BCL6⁺ Pre-GC populations in the spleen of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 9 after infection with PyNL. n = 4–5 mice/group. **(D)** Ratio of BCL6⁺ Pre-GC and IRF4⁺ Pre-GC populations in the spleen on day 9 after infection with PyNL. n = 4–5 mice/group, representative of two independent experiments. **(E)** Time-course analysis of BCL6⁺ B cell population during PyNL infection. n = 4–6 mice/group. **(F)** Population of ASCs in the spleens of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 9 after infection with PyNL. n = 4–5 mice/group. **(G)** IRF4 expression in WT naive B cells stimulated for 16 h with 10 μg/ml anti-CD40 and/or with anti-IgM analyzed by FACS. n = 3 mice. **(H)** RelB and p52 expression in splenic GL7⁺ Pre-GC (CD19⁺ B220⁺ GL7⁺) of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 9 after infection with PyNL. N = 4 mice/group. **(I)** Immunoblot analysis of the time course of in vitro stimulation of naive B cells from *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice with 1 μg/ml anti-CD40. Each dot represents data from an individual mouse. Data are representative of two independent experiments (A–D and F–I) or from three independent experiments (E). Data are shown as mean ± SD. *, P < 0.05; **, P < 0.01; Mann-Whitney t test (B–F and H) or unpaired Student's t test (G). Source data are available for this figure: SourceData F5.

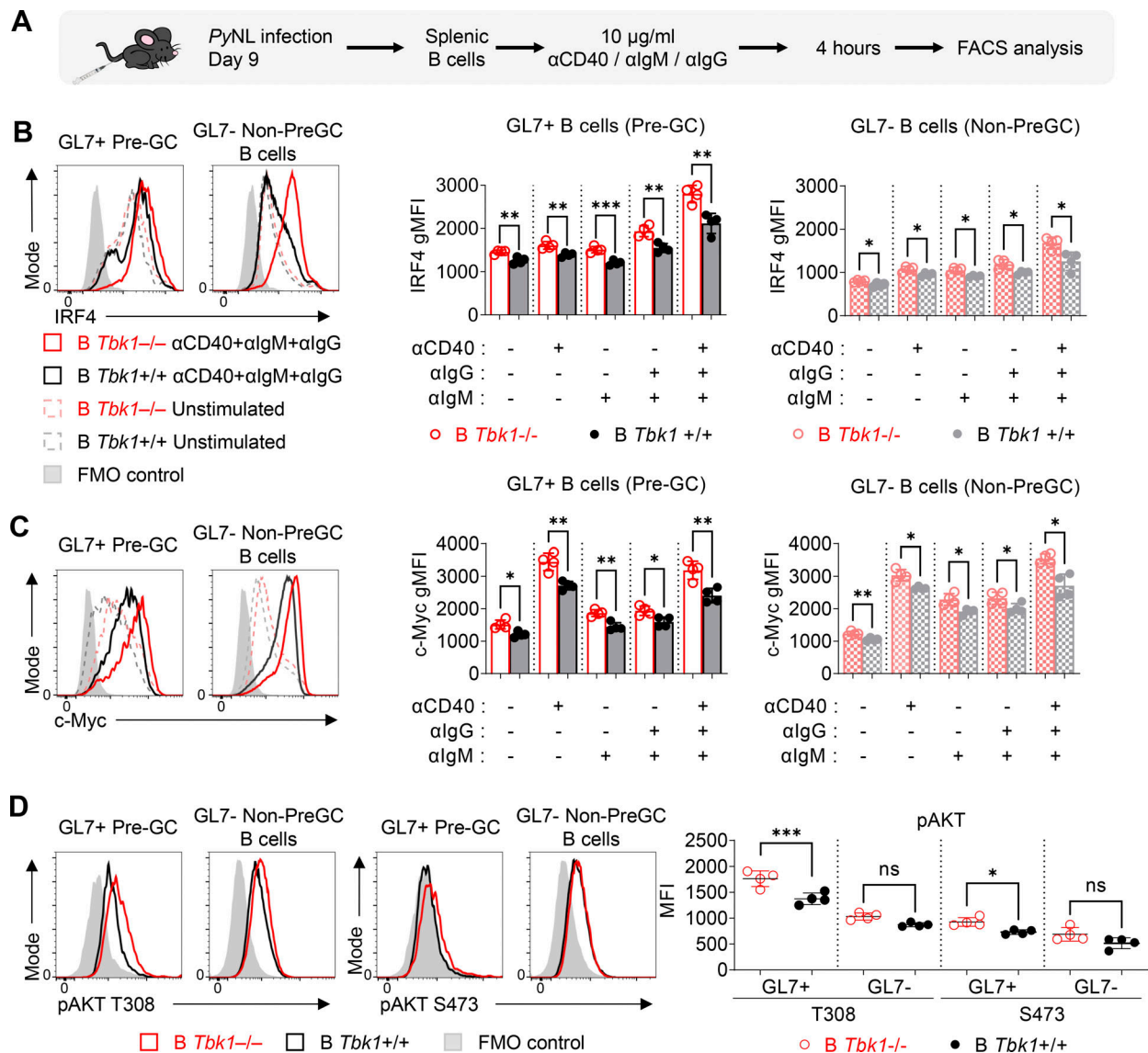


Figure 6. TBK1 is a limiting factor that fine tunes CD40 and BCR signaling in Pre-GC. (A) Schematic diagram of ex vivo splenic B cells isolated from day 9 PyNL-infected $B Tbk1^{-/-}$ and $B Tbk1^{+/+}$ mice with CD40 and/or BCR stimulation for the evaluation of IRF4 and c-Myc expression on gated B220⁺ GL7⁺ Pre-GC and B220⁺ GL7⁻ non-Pre-GC B cell population by FACS. (B and C) Representative histograms and graphs showing IRF4 expression (B) and c-Myc expression (C) of GL7⁺ Pre-GC and GL7⁻ non-Pre-GC upon CD40 and/or BCR stimulation. $n = 4$ mice/group. (D) Representative histograms and graph showing AKT phosphorylation at S473 and T308 on Pre-GC cells on day 9 after infection with PyNL. $n = 4$ mice/group. Each dot represents data from an individual mouse. Data are representative of two independent experiments (A–D). Data are shown as mean \pm SD. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; one-way ANOVA (B and C), Mann-Whitney t test (D). gMFI, geometric mean fluorescence intensity.

for GC commitment, and we also confirmed that TBK1 in CD4⁺ T cells plays a limited role in GC formation. *Tbk1*-deficient B cells can differentiate up to the Pre-GC stage and drive Tfh cell differentiation; however, they failed to further differentiate into GC. Instead, *Tbk1*-deficient Pre-GC committed to a GC-independent fate to differentiate into plasmablasts and memory B cells, but with lack of affinity maturation, and therefore they could not protect the host from an otherwise nonlethal malaria infection. Mechanistically, TBK1 negatively regulates CD40 and BCR signaling to down-regulate IRF4 and c-Myc expression in Pre-GC that leads to the up-regulation of the GC master regulator, BCL6. We further showed that B cell-intrinsic TBK1 fine tunes both CD40 and BCR signaling by negatively regulating

noncanonical NF- κ B, TRAF2, and AKT T308 phosphorylation and thereby synergistically controls IRF4 expression in Pre-GC.

Although known for its role in phosphorylating molecules involved in nucleic acid sensing to produce IFN-I, TBK1 is a pleiotropic kinase that is involved in the regulation of various signaling pathways (Louis et al., 2018). The lack of a consensus binding sequence gave rise to the hypothesis that the involvement of TBK1 depends on its cellular location and abundance (Helgason et al., 2013). This hypothesis is supported by a recent study showing the differential localization of phospho-TBK1 in response to the same stimulant in a cell type-specific manner (Suzuki et al., 2013). Therefore, we would expect TBK1 to have

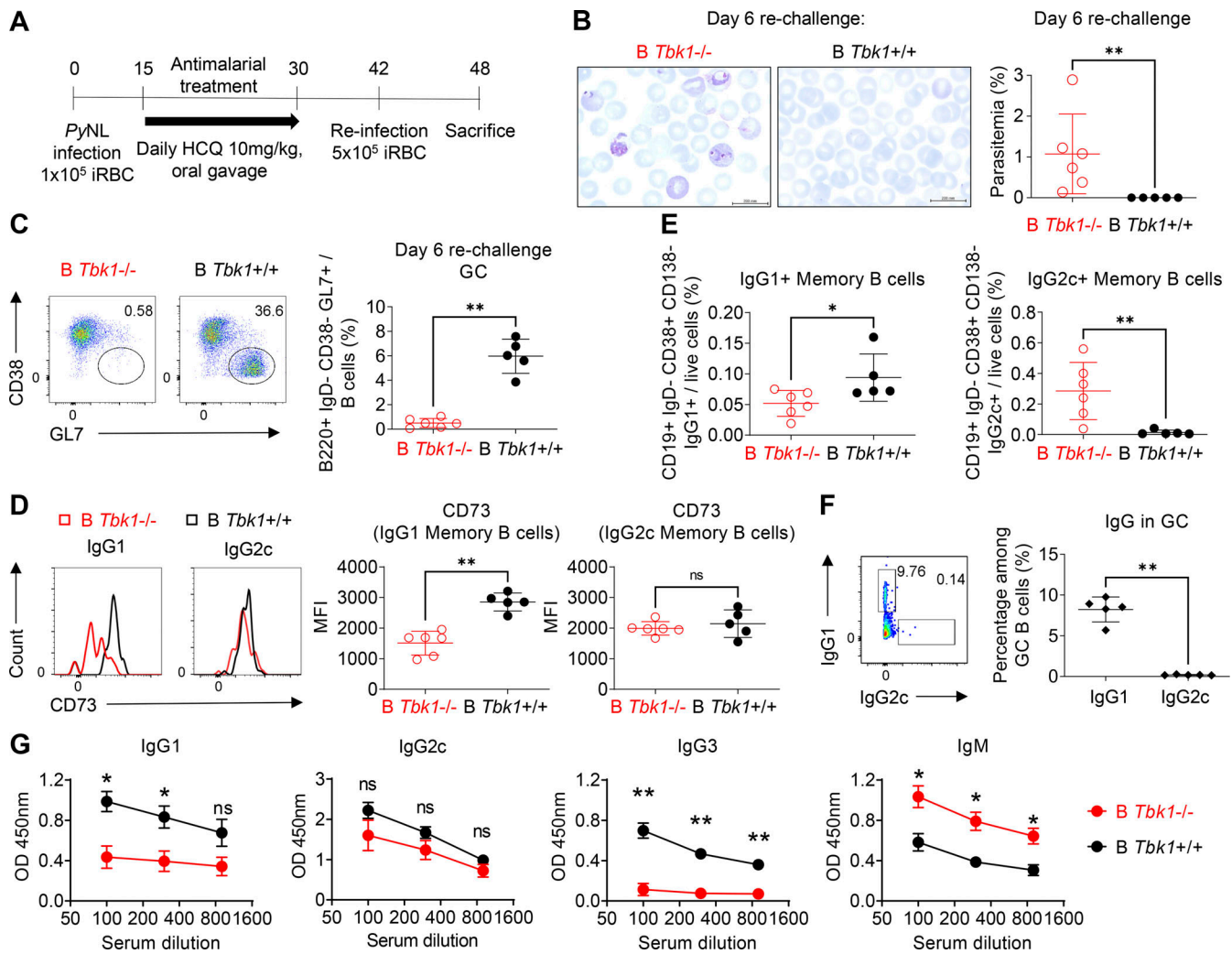


Figure 7. B cell-intrinsic TBK1 deficiency impairs immunity to reinfection. (A) Schematic diagram of PyNL infection, treatment, and reinfection for the evaluation of memory B cells. iRBC, PyNL-infected erythrocytes. (B) Giemsa-stained blood smears of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 6 after challenge with PyNL infection. Scale bar, 200 mm. (C) Population of GC B cells in the spleen of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 6 after challenge with PyNL infection analyzed by FACS. (D) Expression of CD73 on IgG1⁺ or IgG2c⁺ memory B cells (CD19⁺ IgD⁻ CD138⁻ CD38⁺) of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 6 after challenge with PyNL infection analyzed by flow cytometry. MFI, mean fluorescence intensity. (E) Population of IgG1⁺ and IgG2c⁺ GC B cells in the spleens of *B Tbk1*^{+/+} mice on day 6 after challenge with PyNL infection. (F) Population of IgG1⁺ memory B cells in the spleens of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 6 after challenge with PyNL infection. (G) Serum antibody levels of *B Tbk1*^{-/-} and *B Tbk1*^{+/+} mice on day 6 after challenge with PyNL infection. Each dot represents data from an individual mouse. *n* = 5–6 mice/group. Data are representative of two independent experiments (A–G). Data are shown as mean ± SD. *, *P* < 0.05; **, *P* < 0.01; Mann-Whitney *t* test (A–G).

multiple roles in each cell type, depending on the stimulant and differentiation stage. TBK1 is a downstream molecule of cyclic GMP-AMP synthase (cGAS)-STING activation that leads to the induction of IFN-I. Enhanced GC formation in IFNAR KO mice infected with PyNL suggests that IFNAR is not essential for GC formation, but rather has a negative regulatory role (Sebina et al., 2016). Furthermore, cGAS in B cells was shown not to be responsible for GC formation; instead, its expression in innate immune cells mediates B cell responses (Hahn et al., 2018). TBK1 is also involved in autophagy by phosphorylating autophagy receptors to facilitate the recruitment of LC3 molecules (Richter et al., 2016). A study using B cell-specific Atg7-deficient mice suggested that autophagy is not essential for GC formation and maintenance, although it is crucial for memory B cell

maintenance (Chen et al., 2014). Collectively, these studies suggest that the mechanism of TBK1 regulation of GC differentiation is beyond its role in nucleic acid recognition, IFN-I signaling, and autophagy.

It was previously shown that TBK1 regulates IgA class switching, while it did not affect IgG class switching or antibody production (Jin et al., 2012). Therefore, our study clearly suggests the possibility that the increased IgA production in B cell-intrinsic TBK1-deficient mice shown by Jin et al. originated from GC-independent responses. However, whether IgA⁺ GC in Peyer's patches is regulated by TBK1 in a manner similar to that in spleen and LNs requires further investigation. Consistently, we found that TBK1 deficiency did not abolish serum IgG production upon immunization or infection; however, these antibodies

arose from GC-independent plasmablasts. Without affinity maturation in the GC, the quality of antibodies produced was compromised and could not protect the mice from malaria infection. It was long believed that IgG class switching occurs in the GC, but a recent study found that IgG class switching actually happens before B cells enter the GC (Roco et al., 2019), which explains the detection of serum IgG in the absence of GC. Despite the robust IgG2c antibody induction upon PyNL infection, the lack of GC-derived IgG1 in B cell-intrinsic TBK1-deficient mice suggests the importance of IgG1 for sterile protection against malaria and that antibody quality is more crucial than the magnitude of antibody production. Our finding that IgG1⁺ B cells instead of IgG2c⁺ B cells were selected to enter GC during PyNL infection implies the difference in their antigen affinity, because generally higher-affinity B cells outcompete lower-affinity B cells for GC seeding (Schwickert et al., 2011).

A recent study developed a small molecule inhibitor of TBK1 and IKK ϵ , WEHI-112, to treat antibody-dependent arthritis (Louis et al., 2019). It was suggested that the drug might target the innate immune cells to suppress IFN-I and IL-6 and may directly or indirectly suppress Tfh cells, which led to GC inhibition. With our current findings, it is plausible that the TBK1 inhibitor targets B cells directly to suppress GC formation. Surprisingly, we found that CD4⁺ T cell-intrinsic TBK1 had a redundant role in GC differentiation, which is in contrast to a report showing that adoptive transfer of lymphocytic choriomeningitis virus (LCMV)-specific CD4⁺ T cells knocked down for TBK1 failed to induce GC formation upon acute LCMV infection due to impaired Tfh cell maturation (Pedros et al., 2016). The discrepancy could possibly be due to the homing ability of the adoptively transferred TBK1-deficient CD4⁺ T cells to the T cell zone, as *Tbki*-deficient T cells were shown to exhibit impaired migration (Yu et al., 2015). Adoptive transfer of CD4⁺ T cells into a *Cd4^{cre}Bcl6^{fl/fl}* host constitute only a small proportion of T cells among the pool of endogenous host T cells, which may limit the interaction with cognate B cells due to competition with endogenous *Bcl6*-deficient T cells. Complete deletion of *Tbki* in CD4⁺ T cells using *Cd4^{cre}Tbki^{fllox}* mice allowed us to overcome these limitations in this study.

Initial antigen encounter through the BCR is essential for both GC-dependent and GC-independent humoral responses. Although high-affinity B cells outcompete low-affinity B cells for GC seeding, even B cells with low BCR affinity are able to migrate to the T-B border to receive T cell help to form the GC in the absence of high-affinity B cells (Dal Porto et al., 2002; Schwickert et al., 2011), indicating that T cell help plays a crucial role in determining GC commitment. However, recent evidence also suggested that overactivation of BCR signaling induces plasmablast and GC B cell death (Yam-Puc et al., 2021). The ability of *Tbki*-deficient B cells to induce normal Tfh cell differentiation suggests that antigen presentation was not impaired. The amount of T cell help through CD40 signaling is particularly important for B cell fate decision. GC formation is abolished in CD40-deficient mice (Kawabe et al., 1994); however, CD40-mediated gene signatures were only found in Pre-GC and post-GC memory B cells and were absent during GC expansion (Basso et al., 2004). We found that TBK1 is not required in the

initiation of Pre-GC differentiation, but rather involved in the fate decision of Pre-GC, by limiting CD40 and BCR signaling for the progression of GC differentiation. Excessive CD40 signaling was shown to halt Pre-GC further differentiation into GC through sustained noncanonical NF- κ B-activated IRF4 expression (Zhang et al., 2017). Although CD40-induced noncanonical NF- κ B and IRF4 are essential for GC initiation, the expression of RelB and IRF4 is transient during GC initiation and absent from mature GC (De Silva et al., 2016; Willis et al., 2014). Increased noncanonical NF- κ B and IRF4 expression in TBK1-deficient Pre-GC inhibits GC differentiation by inhibiting BCL6 expression, as high IRF4 expression directly binds to the promoter site of the *Bcl6* gene as a transcription repressor (Saito et al., 2007). CD40-induced IRF4 was also shown to down-regulate ICOSL expression in B cells (Ochiai et al., 2018); therefore, it is likely that the high IRF4 expression in *Tbki*^{-/-} B cells caused unsustainable ICOSL-ICOS interaction between Tfh cells and cognate B cells. Additionally, we found that TBK1 is involved in the synergy of CD40 and BCR activation to negatively regulate IRF4 expression and BCR-mediated AKT T308 phosphorylation in Pre-GC.

AKT phosphorylation at T308 and S473 was shown to be TBK1 dependent in fibroblasts upon stimulation with glucose or growth factors (Ou et al., 2011); however, the loss of TBK1 unexpectedly promotes pAKT at S473 in CD4⁺ T cells upon TCR activation, suggesting cell type specificity and stimulant dependency (Yu et al., 2015). Therefore, although TBK1 has been shown to phosphorylate AKT at T308 in other non-B cells, there has been no information indicating whether TBK1 also phosphorylates AKT at T308 in B cells. Our data show that TBK1 acts as a negative regulator of AKT phosphorylation at T308 in Pre-GC B cells. Notably, a recent paper showed that increased pAKT T308 inhibits pAKT S473 to allow migration of GC dark zone B cells to reenter the GC light zone to interact with T cells (Luo et al., 2019). Therefore, the increased pAKT T308 in *Tbki*^{-/-} B cells before GC maturation might inhibit pAKT S473 in early BCR activation and sustain Pre-GC interaction with T cells and thus inhibit Pre-GC progression into GC. Furthermore, our data also showed increased CD40 signaling in *Tbki*^{-/-} Pre-GC, suggesting that the increased pAKT T308 in *Tbki*^{-/-} Pre-GC might promote T cell interaction and thus enhanced CD40 signaling.

CD40 activation recruits TRAF proteins to its cytosolic domain, leading to downstream signaling (Elgueta et al., 2009). TRAF2 is involved in CD40 signaling for the induction of noncanonical NF- κ B. Dephosphorylation of TRAF2 at S11 was shown to be required for NF- κ B-inducing kinase (NIK) accumulation to drive noncanonical NF- κ B in CD40-stimulated naive B cells (Workman et al., 2020). Consistently, we found that CD40 stimulation gradually dephosphorylates TRAF2 over a time course in a TBK1-dependent manner. Interestingly, the trend of TRAF2 dephosphorylation coincides with TBK1 dephosphorylation upon CD40 stimulation. Notably, TBK1 is constitutively phosphorylated in unstimulated naive B cells, but not in other non-B cells, in the spleen. Together, these findings further suggest that phospho-TBK1-mediated TRAF2 S11 phosphorylation is required to constantly degrade NIK to prevent spontaneous noncanonical NF- κ B activation in the absence of CD40 stimulation. TRAF2-deficient B cells also exhibited a phenotype

similar to that of TBK1-deficient B cells with impaired GC formation (Gardam et al., 2011), suggesting the importance of TBK1 in CD40 regulation via TRAF2 for optimal GC formation.

The GC provides stringent quality control for B cells through clonal selection, coupled with several rounds of somatic hypermutation to produce antigen-specific high-affinity B cells that exit as either long-lived plasma cells or memory B cells. Upon reencountering antigen, GC-derived memory B cells can immediately differentiate into ASCs or reenter the GC for further rounds of somatic hypermutation to increase its affinity maturation. Memory B cells are heterogeneous, and recently a few markers have been used to identify subsets of antigen-specific memory B cells (Tomayko et al., 2010). CD73 is an ecto-5'-nucleotidase expressed by a subset of memory B cells that correlates with somatic hypermutation (Anderson et al., 2007; Kaji et al., 2012) and therefore is considered to be a good candidate to distinguish GC-derived memory B cells. CD80⁺ memory B cells were shown to immediately differentiate into plasmablasts upon antigen reencounter instead of entering the GC (Zuccarino-Catania et al., 2014), while CD80⁺ memory B cells were also induced in B cell-intrinsic *Bcl6*-deficient mice that lack the GC (Kaji et al., 2012), suggesting its GC-independent origin. Notably, we noticed that CD80 expression was up-regulated in TBK1-deficient Pre-GC, which is consistent with the notion that GC-independent memory B cells arise early before GC formation. Challenge with PyNL infection suggests that these GC-independent memory B cells confer limited protection for sterile immunity.

Plasmodium spp. parasites evolve various strategies to evade host immunity, making it difficult to develop a vaccine that can provide sterile immunity (Rénia and Goh, 2016). In humans, recovery from malaria infection does not induce long-lasting immune memory to protect from subsequent infection, although the partial immunity is able to reduce parasite burden and disease complications (Lee and Coban, 2018). Recent studies showed that the impaired humoral immunity in malaria patients is associated with impaired Tfh cell phenotypes (Obeng-Adjei et al., 2015) that give rise to short-lived atypical memory B cells (Weiss et al., 2009; Obeng-Adjei et al., 2017) that produce transient antibodies. However, there is a lack of evidence whether malaria has a direct pressure on B cells to suppress GC differentiation to give rise to long-lived plasma cells and memory B cells. The gene expression of *Tbkl* in human PBMCs was found to correlate with the severity of acute *Plasmodium falciparum* infection (Yamagishi et al., 2014). However, *Plasmodium* parasites were shown to evade host immunity by antagonizing TBK1 activity. Indeed, host receptor transporter protein 4 and FOS-like antigen 1 induced by malaria infection were shown to inhibit TBK1 phosphorylation and TBK1 interaction with IFN- γ signaling molecules, exacerbating the severity of infection (He et al., 2020; Cai et al., 2017). In addition, several studies have shown the inhibitory effects of parasite factors or host factors on B cell immunity during malaria, such as *Plasmodium*-expressing repetitive interspersed families of polypeptides (Saito et al., 2017), ApiAP2 (Akkaya et al., 2020a), and plasmepsin-4 (Spaccapelo et al., 2010), as well as host LAIR1 and LILRB1, which are expressed on host immune cells, including B cells

(Saito et al., 2017), or indirectly through the dysregulation of CD4⁺ T cells (Butler et al., 2011; Zander et al., 2015; Kurup et al., 2017; Obeng-Adjei et al., 2017). Together, this evidence suggests that malaria evades B cell immunity through multiple mechanisms, including the inhibition of TBK1 activity in B cells to limit GC formation, thereby suppressing long-lasting immune memory. Here, for the first time, we address the crucial role of TBK1 in GC formation and highlight its importance in long-lasting humoral responses required for sterile immunity against malaria infection.

In conclusion, our study demonstrated the role of B cell-intrinsic TBK1 in the B cell fate decision for GC differentiation and the importance of GC-dependent antibody production and long-lived memory B cell formation for the protection against pathogen reinfection. The involvement of TBK1 in various signaling pathways makes it act as a double-edged sword in different diseases. TBK1 is required for IFN- γ -mediated antiviral immunity, while here we additionally showed its importance in GC-mediated humoral immunity. However, TBK1 is also involved in the pathogenesis of autoimmune diseases (Louis et al., 2018). TBK1 inhibitors such as amlexanox have been used in clinical treatment for asthma (Inagaki et al., 1992) and have recently been proposed for the treatment of obesity-related metabolic disease (Reilly et al., 2013; Oral et al., 2017). Our findings suggest that TBK1 inhibitors should be administered with caution, considering their possible effect on humoral immunity in case of infection and vaccination. Moreover, the timing of TBK1 inhibitor administration may be important to target TBK1 in specific cell types during disease progression and to prevent its suppressive effects on vaccine efficacy.

Materials and methods

Mice

WT C57BL/6J mice were purchased from CLEA Japan. Female CD45.1⁺ C57BL/6J mice were purchased from Sankyo Laboratory Service. μ MT mice were purchased from The Jackson Laboratory. *Cd4*-cre mice (The Jackson Laboratory) were crossed with *Bcl6*^{fllox/fllox} to generate *Cd4*^{cre} *Bcl6*^{fl/fl} mice and *Cd4*^{wt} *Bcl6*^{fl/fl} mice (Lee et al., 2019; Ise et al., 2014). *Mbl*-cre mice, kindly provided by M. Reth (University of Freiburg, Freiburg, Germany; Hobeika et al., 2006), and *Cd4*-cre mice were crossed with *Tbkl*^{fllox/fllox} mice (Taconic) to generate *Mbl*^{cre/wt} *Tbkl*^{fl/fl} and *Cd4*^{cre/wt} *Tbkl*^{fl/fl} mice, respectively. *Tbkl*^{fl/fl} littermates were used as control WT counterparts. Animal experiments were conducted in accordance with institutional guidelines and approved by the review board for animal experiments of The Institute of Medical Science, The University of Tokyo (approval PA19-48).

Malaria infection and treatment

Mice were infected with 10⁵ PyNL-infected erythrocytes i.p. or otherwise as mentioned in the figure legends (Lee et al., 2017). Parasitemia during the course of infection was quantified by Giemsa-stained blood smears or flow cytometry (Lelliott et al., 2014). In Fig. 7, PyNL-infected mice were treated with 10 mg/kg HCQ orally from day 15 to day 30 after infection. Parasitemia resolution was confirmed by Giemsa-stained blood smears.

HCQ-treated mice recovered from PyNL infection were re-infected with 5×10^5 PyNL-infected erythrocytes 1 wk after the end of HCQ treatment to prevent the prophylactic effect of HCQ.

Immunization

Mice were immunized with 100 μ g NIP-OVA (Biosearch Technologies) mixed with alum (InvivoGen) intradermally (i.d.). Sera were collected for antibody assay, inguinal LNs were collected for cellular analysis by flow cytometry, and spleens were collected for immunohistochemical analysis on day 12 or day 21 after immunization.

Mixed bone marrow chimera

Bone marrow mixtures from female Ly5.1⁻ *Mbl^{cre/wt}Tbkl^{f/f}* mice, Ly5.1⁺ WT mice, and μ MT mice were mixed at a 1:1:8 ratio and injected i.v. into lethally irradiated (8.5 Gy) male μ MT mice. Mice were fed enrofloxacin (Elanco) in drinking water for 5 wk. Mice were immunized with NIP-OVA and alum i.d. 7 wk after bone marrow transfer. Spleens were analyzed by flow cytometry on day 12 after immunization.

B cell isolation

Human PBMC purified B cells were isolated by Ficoll-Paque, followed by anti-CD19 magnetic beads (Miltenyi Biotec). Splenic naive B cells were purified using a B cell isolation kit by negative selection (Miltenyi Biotec). Splenic Pre-GC and GC B cells were first enriched by streptavidin magnetic beads conjugated with a mixture of biotinylated anti-CD11b, anti-CD11c, anti-IgD, anti-CD138, anti-CD3, and anti-Ter119 (Miltenyi Biotec), followed by cell sorting using a BD FACSAria III Cell Sorter. Splenic Pre-GC cells defined as B220⁺ IgD⁻ CD38⁺ GL7⁺ cells were isolated from PyNL-infected mice on day 9 after infection. Splenic mature GC B cells defined as B220⁺ IgD⁻ CD38⁻ GL7⁺ cells were purified from mice recovered from PyNL infection after day 28 infection.

RNA-seq

5×10^4 cells were directly sorted into TRIzol reagent (Invitrogen). RNAs were purified following the manufacturer's instructions with the addition of RNA grade glycogen (Thermo Fisher Scientific). RNA integrity number >9.7 was confirmed using the RNA 6000 Nano kit on an Agilent 2100 Bioanalyzer. DNA libraries were constructed using the TruSeq Stranded mRNA Sample Prep Kit. RNA-seq was performed by the Genome Information Research Center of Osaka University using NovaSeq 6000 (Illumina) in a 100-bp single-end read mode. Sequence reads were mapped to mouse reference genome sequence mm10 using TopHat version 2.0.13, Bowtie2 version 2.2.3, and SMtools version 0.1.19. Fragments per kilobase of exon per million reads mapped were calculated by Cufflinks version 2.2.1. Data were analyzed using iDEP.91 and GSEA software version 4.0.3 (Subramanian et al., 2005). GSEA results were analyzed based on the collection of MSigDB gene sets and gene signatures up-regulated or down-regulated by CD40 and BCR stimulation as described previously (Victoria et al., 2010). GO enrichment analysis was performed using iDEP.91. Heatmaps for differentially expressed genes (DEGs) were generated by iDEP.91 or Morpheus after filtering out genes with low statistical significance

($P > 0.05$). Genes presented in the heatmaps were arranged from top to bottom with descending differential expression. The RNA-seq data are available in the Gene Expression Omnibus database (accession no. GSE188659).

Antibody assay

Antibody titers were measured from sera collected from PyNL-infected or immunized mice by ELISA. High-binding 96-well microtiter plates were coated overnight with 10 μ g/ml PyNL parasite crude extract, uninfected RBC extract, or 4-Hydroxy-3-nitrophenylacetyl hapten conjugated BSA ratio 30 (Biosearch Technologies) and blocked with 5% skim milk or 1% BSA (Nacalai Tesque). Sera were serially diluted, and antibodies were detected using HRP-conjugated goat anti-mouse IgG1, IgG2c, IgG3, and IgM antibodies (SouthernBiotech). Anti-PyNL antibody levels were calculated by subtracting autoantibodies against uninfected RBCs. Antibody avidity assays were performed as described (Coban et al., 2004). Briefly, after overnight incubation of sera in antigen-coated plates, antibodies were dissociated with various concentrations of sodium thiocyanate at 0, 1, 2, or 4 M for 15 min. Plates were washed extensively six times before proceeding to subsequent antibody detection by ELISA as described above.

In vitro and ex vivo B cell stimulation

B cells were cultured in RPMI 1640 (Nacalai) supplemented with 10% FBS (Sigma-Aldrich), penicillin and streptomycin antibiotics (Nacalai), Hepes (Nacalai), glutamine (Nacalai), sodium pyruvate (Nacalai), nonessential amino acids (Nacalai), and β -mercaptoethanol (Nacalai). Purified naive B cells were stimulated with 1 μ g/ml anti-CD40 (HM40-3; eBioscience) or 1 μ g/ml F(ab')₂ goat anti-IgM (16-5092-85; eBioscience) or 1 μ g/ml F(ab')₂ goat anti-IgG (16-5098-85; eBioscience). For ex vivo stimulation of Pre-GC, splenocytes from day 9 PyNL-infected mice were stimulated with 1 μ g/ml anti-CD40 (eBioscience) and/or with 1 μ g/ml anti-IgG (eBioscience) and 1 μ g/ml anti-IgM (eBioscience). Cells were harvested at the time points indicated in the figure legends.

Cell proliferation and apoptosis assay

5×10^4 cells were stained with prewarmed CellTrace Violet (CTV; Invitrogen) in serum-free RPMI for 10 min at 37°C and washed twice with RPMI containing 10% FCS. Cells were stimulated with 1 μ g/ml anti-CD40 (eBioscience) and/or with 1 μ g/ml anti-IgM (eBioscience). After 4 d, cells were washed with cold PBS and proceeded to flow cytometric analysis. Cell proliferation was indicated by the dilution of CTV fluorescence intensity. For the in vivo EdU pulse assay, mice were injected with 1 μ g EdU in PBS, and spleens were collected 30 min later. Cell cycle and proliferation were detected using the Click-iT EdU assay kit (Invitrogen) and SYTOX AADvanced (Invitrogen). For cell death and apoptosis assay, cells were stained with the CellEvent Caspase-3/7 Green Flow Cytometry Assay Kit (Invitrogen).

Flow cytometry

Spleen or inguinal LNs were crushed on ice and were passed through 70- μ m mesh to collect single-cell suspensions. Erythrocytes were lysed using ammonium-chloride-potassium lysis

buffer (Lonza) and washed twice with PBS supplemented with 2% heat-inactivated FCS. Cells were incubated with Fc blocker (BioLegend) for 5 min at room temperature before staining with fluorophore-conjugated antibody mixtures at 4°C for 30 min for surface antigen staining. Dead cells were excluded using LIVE/DEAD Fixable Dead Cell Stain (Invitrogen). Cells were fixed and permeabilized with the Foxp3/Transcription Factor Fixation/Permeabilization kit (eBioscience) for intracellular staining. Detection of phosphoproteins and signaling molecules was performed as described previously (Rip et al., 2020). Briefly, cells were fixed with 2% paraformaldehyde (Nacalai Tesque) for 10 min, permeabilized with permeabilization buffer (eBioscience), and stained with Fc blocker followed by rabbit anti-mouse primary antibodies (Cell Signaling Technology) against pAKT T308 (D25E6), pAKT S473 (193H12), c-Myc (D84C12), p65 (D14E12), and p52 or RelB (C1E4) for 30 min at 4°C. After washing with permeabilization buffer, samples were further stained with Alexa Fluor 488-conjugated goat anti-rabbit secondary antibody (Invitrogen) together with the mixture of fluorophore-conjugated antibodies for surface antigen staining. Fluorescence minus one control contains all antibodies except primary antibodies. Samples were analyzed using an LSRFortessa flow cytometer (BD Biosciences). Data were analyzed with FlowJo 10.7.1. Anti-B220 (RA3-6B2), IgD (11-26c.2a), IgG1 (A85-1), IgM (RMM-1), CD138 (281-2), CD38 (90), TCR β (H57-597), CD4 (RM-4-5), CD44 (IM7), CXCR5 (L138D7), BCL6 (7D1), CTLA-4 (UC10-4B9), ICOS (C398.4A) and IRF4 (IRF4.3E4) were purchased from BioLegend. Anti-GL7 (GL-7), PD-1 (J43), Foxp3 (FJK-16s) and IL-21 (FFA21) were purchased from eBioscience. Goat anti-mouse IgG2c human ads-PE were purchased from SouthernBiotech.

Western blotting

Purified B cell populations were lysed with radioimmunoprecipitation assay buffer (Nacalai Tesque) supplemented with protease inhibitor cocktail (Sigma-Aldrich), phosphatase inhibitor cocktail III (Sigma-Aldrich), and PMSF (Cell Signaling Technology). Subcellular fractionation was performed using the Cell Fractionation Kit (Cell Signaling Technology). Protein concentrations were quantified using the Pierce BCA Protein Assay Kit (Thermo Fisher Scientific). Cell lysates were boiled at 95°C for 5 min with NuPAGE LDS Sample Buffer (Invitrogen) and NuPAGE Sample Reducing Agent (Invitrogen). Proteins were separated by SDS-PAGE electrophoresis in NuPAGE 4–12% Bis-Tris Mini Gel (Invitrogen) at 100 V. Proteins were transferred to polyvinylidene fluoride membrane using the iBlot2 gel transfer device (Thermo Fisher Scientific). Membranes were blocked with 5% skim milk or BSA in Tris-buffered saline with Tween 20 (Nacalai Tesque) before overnight incubation with the following rabbit anti-mouse primary antibodies (Cell Signaling Technology): phospho-TBK1 (D52C2), TBK1 (D1B4), phospho-TRAF2 Ser11 (E2B6L), TRAF2 (C192), NIK, phospho-NF- κ B p65 Ser536, NF- κ B p100/p52, RelB (C1E4), β -actin (D6A8), β -tubulin (D2N5G), or lamin B1 (D4Q4Z). Goat anti-rabbit IgG HRP (Cell Signaling Technology) was used as a secondary antibody. Luminescence was captured and measured using an Amersham Imager 600 (GE Healthcare). The protein amount was normalized

with the band intensity of the loading control. WB stripping solution (Nacalai Tesque) was used to remove membrane-bound antibodies before subsequent antibody staining.

Immunohistochemical analysis

Spleens were collected directly into 4% paraformaldehyde on ice and fixed at 4°C overnight. Samples were transferred into 30% sucrose solution at 4°C overnight with constant shaking. Samples were then embedded in Tissue-Tek O.C.T. Compound (Sakura Finetek) and instant frozen at -80°C . Samples were then sectioned with a cryostat at 10- μm thickness and -20°C . Sections were washed with PBS and blocked for 30 min with 3% BSA in Tris-buffered saline with Tween 20. Sections were stained with rat anti-mouse Alexa Fluor 488-GL7 (GL7; BioLegend) overnight at 4°C, washed, and further incubated overnight at 4°C with biotin-CD4 (RM4-5; BioLegend), followed by staining with streptavidin Alexa Fluor 405 (Invitrogen) for 2 h at room temperature and finally with Alexa Fluor 647-IgD (11-26c.2a; BioLegend) for 1 h at room temperature. Stained tissue sections were mounted on coverslips using fluorescence mounting medium (Dako). Tissue sections were observed and captured with an all-in-one fluorescence microscope (Keyence).

Statistical analyses

Reported *n* values are the total number of samples per group. Each sample represents an individual mouse. Graphs were plotted and analyzed with GraphPad Prism 9.0 software. The Mann-Whitney *t* test was used for comparing two groups, while one-way ANOVA was used for comparing more than two groups. Survival between two groups was compared by log-rank Mantel-Cox test.

Online supplemental material

Fig. S1 shows Tfh and GC B cell formation during the course of PyNL infection and parasitemia and survival of Tfh cell-deficient mice. Fig. S2 shows phospho-TBK1 in purified human B cells and mouse splenic B cells; depletion efficiency of B *Tbkl*^{-/-} mice; and B cell subsets in spleen, peritoneum, and bone marrow. Fig. S3 shows a comparison of antibody responses and ASC formation between B *Tbkl*^{-/-} mice and B *Tbkl*^{+/+} mice following PyNL infection or NIP-OVA + alum immunization. Fig. S4 shows heatmaps of RNA-seq data for the GO biological process pathway related to regulation of signal transduction and TBK1-dependent genes up-regulated by CD40 or BCR stimulation in Pre-GC. Fig. S5 shows a comparison of CD40-induced NF- κ B activation between *Tbkl*^{-/-} and *Tbkl*^{+/+} naive B cells. Table S1 shows RNA-seq data of *Tbkl*^{-/-} and *Tbkl*^{+/+} Pre-GCs isolated from PyNL-infected mice.

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Supplemental material

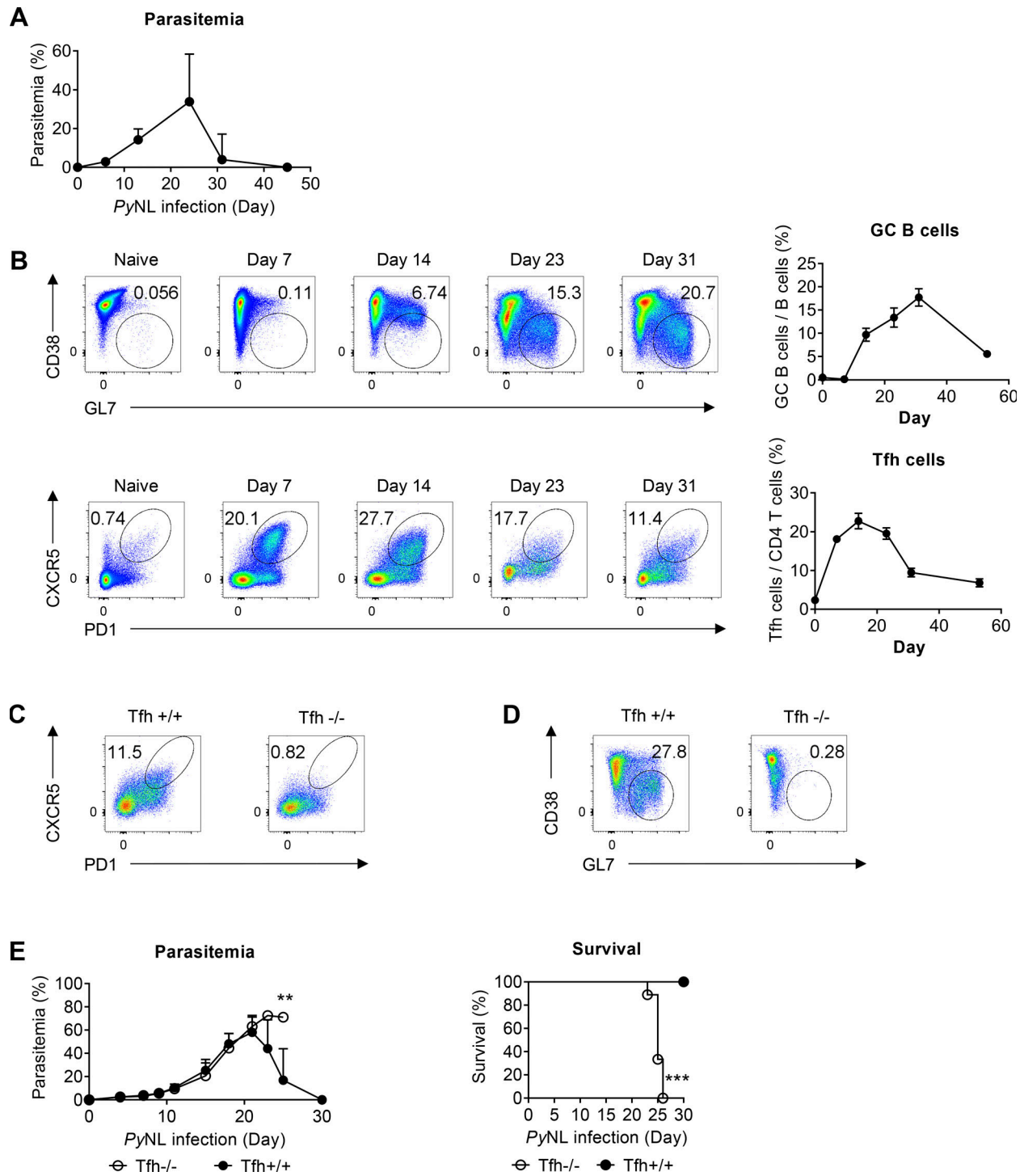


Figure S1. **Tfh cells and GC formation are essential for recovery from PyNL infection.** (A) Parasitemia of WT mice infected with PyNL. $n = 24$ mice pooled from four independent experiments. (B) Time-course analysis of Tfh cells and GC formation in the spleens of WT mice infected with PyNL. $n = 4-5$ mice per time point from four independent experiments. (C and D) Representative FACS plots of Tfh (C) and GC (D) populations in the spleens of $Cd4^{cre}Bcl6^{fl/fl}$ ($Tfh^{-/-}$) mice and $Cd4^{cre}Bcl6^{fl/fl}$ ($Tfh^{+/+}$) mice infected with PyNL. $n = 4$ mice/group representative of two independent experiments. (E) Parasitemia and survival of $Tfh^{-/-}$ mice and $Tfh^{+/+}$ mice infected with PyNL. $n = 9-10$ mice/group pooled from two independent experiments. Each dot represents data from an individual mouse. Data are pooled from four independent experiments (A and B) or representative of two independent experiments (C and D). Data are shown as mean \pm SD. **, $P < 0.01$; ***, $P < 0.001$; Mann-Whitney t test (E), log-rank (Mantel-Cox) test (E).

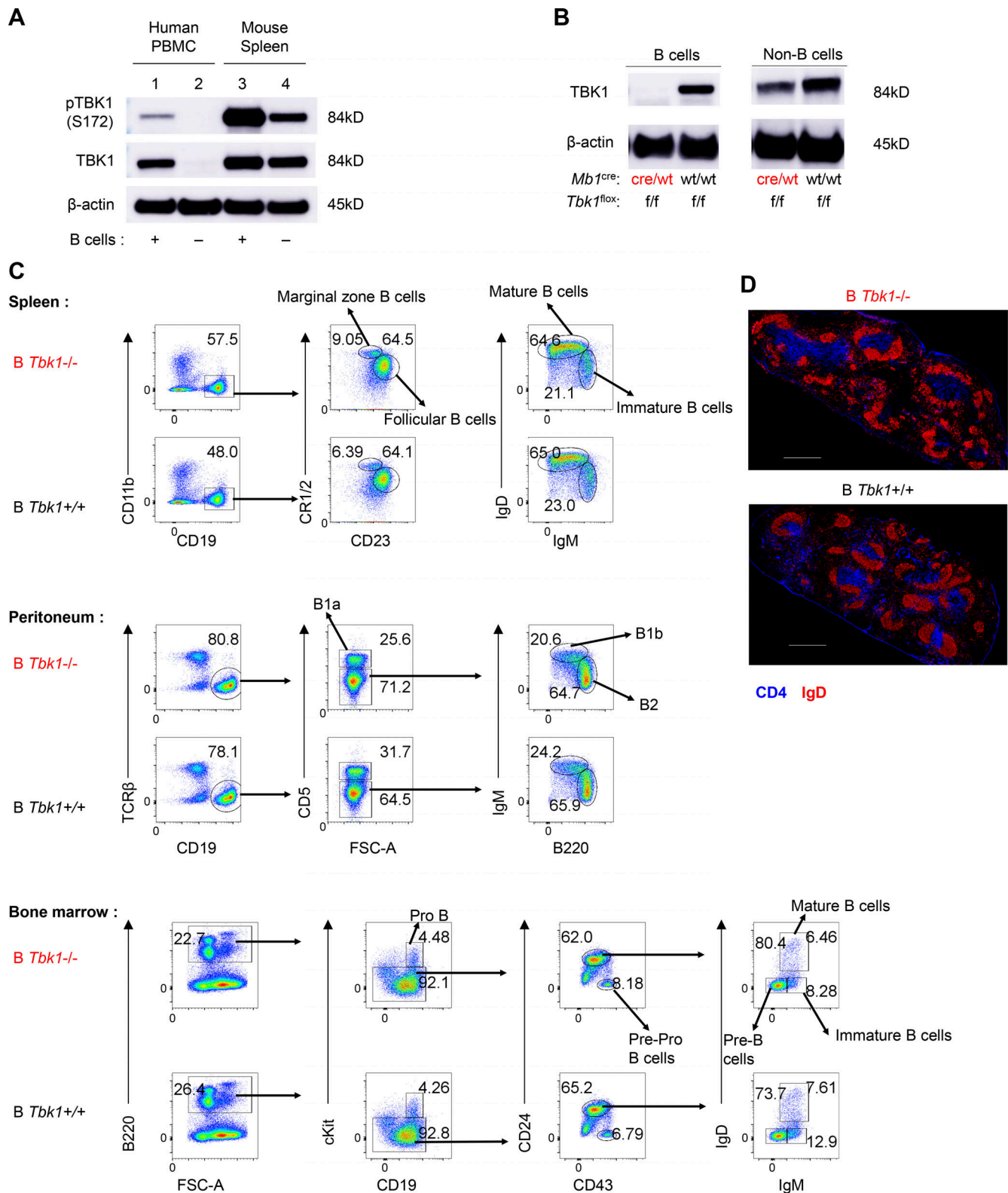
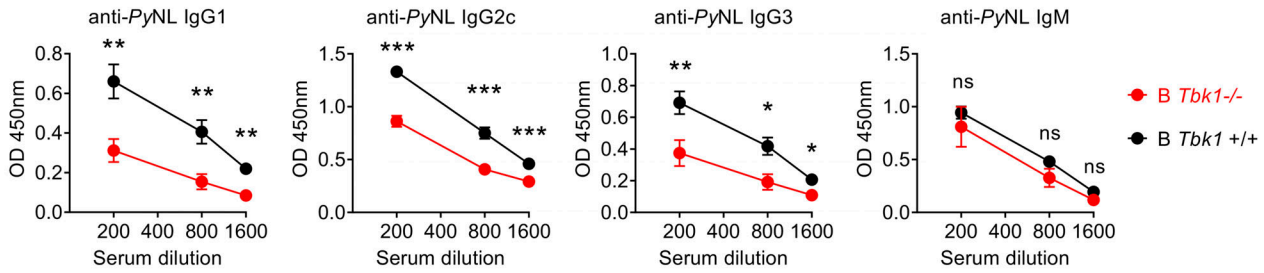
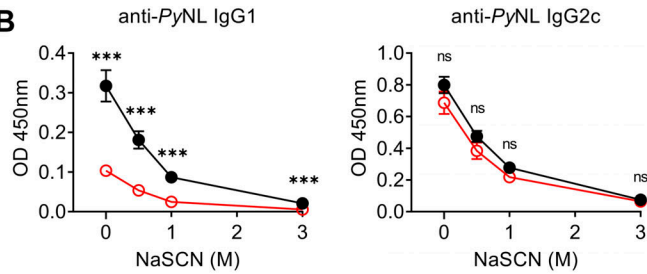


Figure S2. **TBK1 expression in B cell and deletion efficiency in B cells and T cells using a Cre-loxP system.** (A) Expression and phosphorylation of TBK1 in human B cells from PBMCs and mouse splenic B cells by immunoblotting. (B) Depletion efficiency of TBK1 in splenic B cells was analyzed by immunoblotting. (C) Analysis of subpopulations of B cells in spleen, peritoneum, and bone marrow isolated from *B Tbk1^{-/-}* and *B Tbk1^{+/+}* mice by flow cytometry. FSC-A, forward scatter area. (D) Immunohistochemical staining of B cell zone (red; IgD) and T cell zone (blue; CD4) in the spleens of *B Tbk1^{-/-}* and *B Tbk1^{+/+}* mice. Data are representative of two (A) or five (B) independent experiments or from one experiment with a representative of three independent biological samples (C and D). Scale bars, 500 μm. Source data are available for this figure: SourceData FS2.

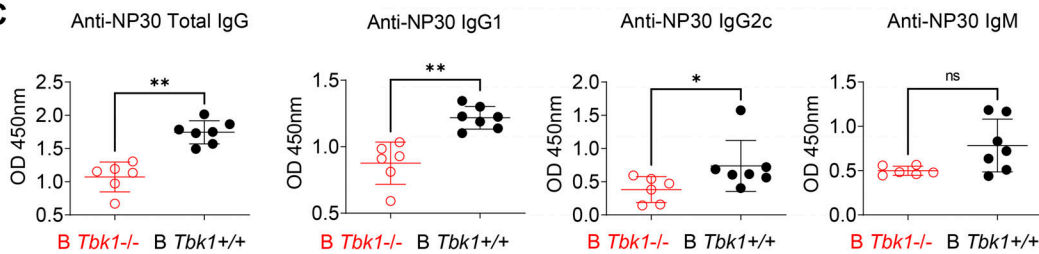
A Primary PyNL infection:



B



C



D PyNL-infected Day 24

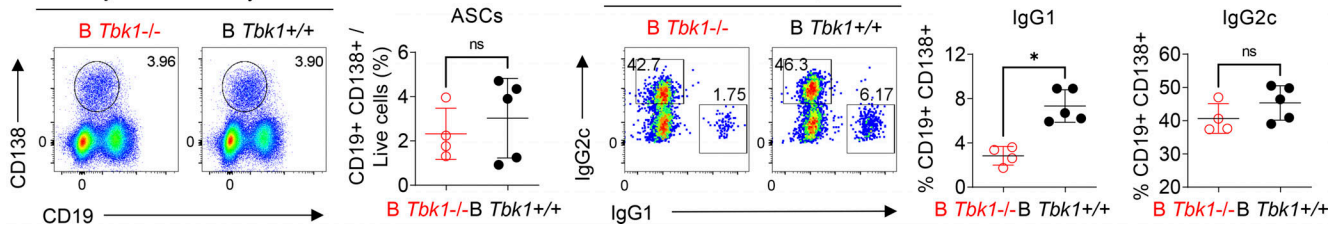


Figure S3. **B** *Tbk1*^{-/-} mice could produce antigen-specific antibodies, albeit less of them, despite the lack of GC formation. **(A)** PyNL-specific serum antibody measurement by ELISA on days 28–35 after infection with PyNL. *n* = 6–10 mice/group from three independent experiments. **(B)** Avidity assay of PyNL-specific serum antibody on days 19–35 after infection with PyNL. *n* = 10–12 mice/group from five independent experiments. **(C)** NP₃₀ (4-hydroxy-3-nitrophenylacetyl hapten conjugated BSA ratio 30)-specific serum antibody measurement by ELISA on day 21 after immunization with NIP-OVA with alum i.d. *n* = 6–7 mice/group representative of two independent experiments. **(D)** ASC population in the spleens of PyNL-infected mice on day 24 analyzed by FACS. *n* = 4–5 mice/group representative of two independent experiments. Each dot represents data from an individual mouse. Data are pooled from three (A) or five (B) independent experiments and representative of two independent experiments (C and D). Data are shown as mean ± SD. *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001; Mann-Whitney *t* test (A–D).

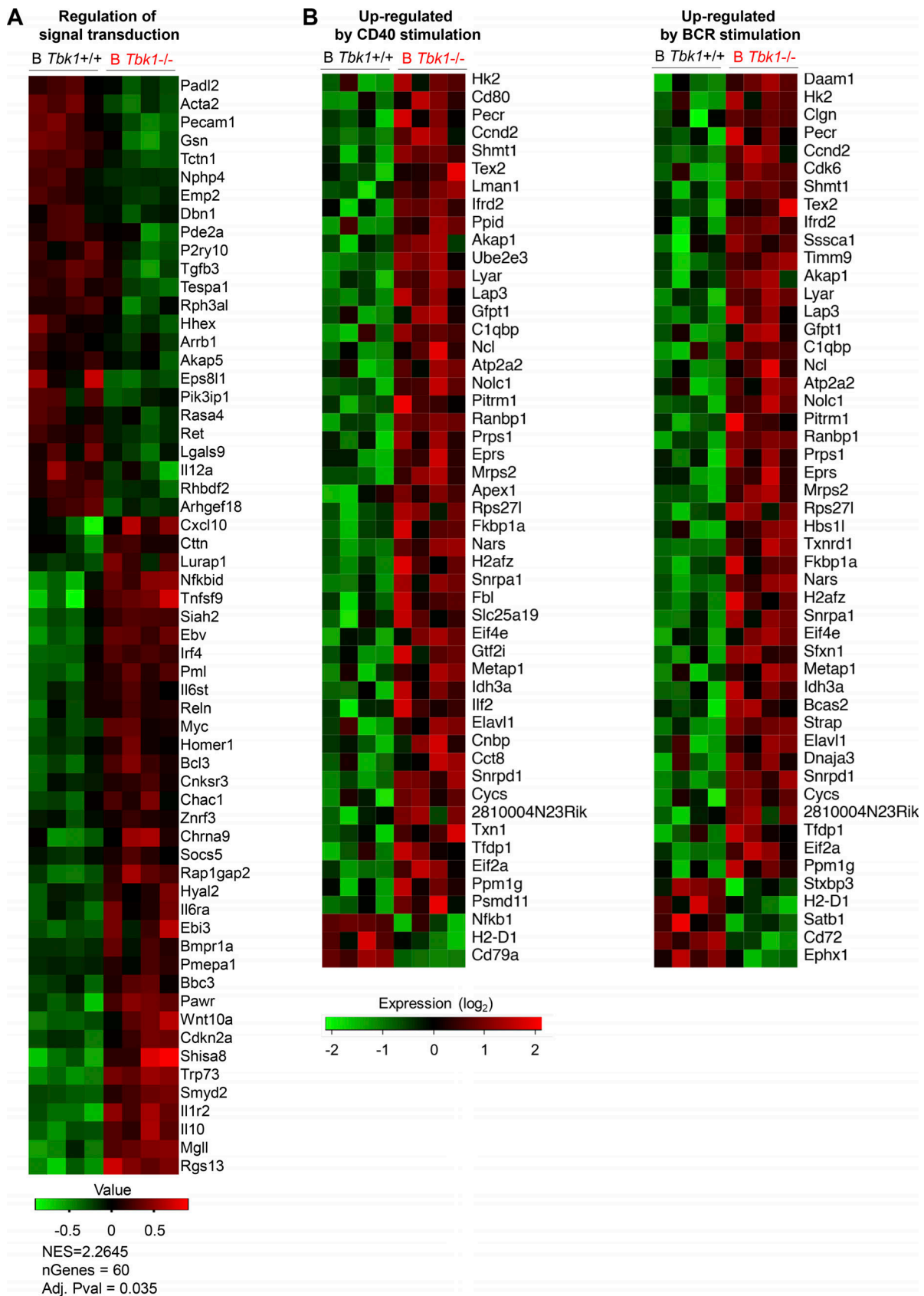


Figure S4. **TBK1 negatively regulates CD40 and BCR signaling.** (A) Heatmap of GO biological process pathway analysis by GSEA (preranked fast GSEA) based on RNA-seq analysis. (B) TBK1-dependent genes up-regulated by CD40 or BCR stimulation in Pre-GC analyzed by GSEA based on RNA-seq data (related to Fig. 4 I). Data are from one experiment. $n = 4$ mice/group. NES, normalized enrichment score.

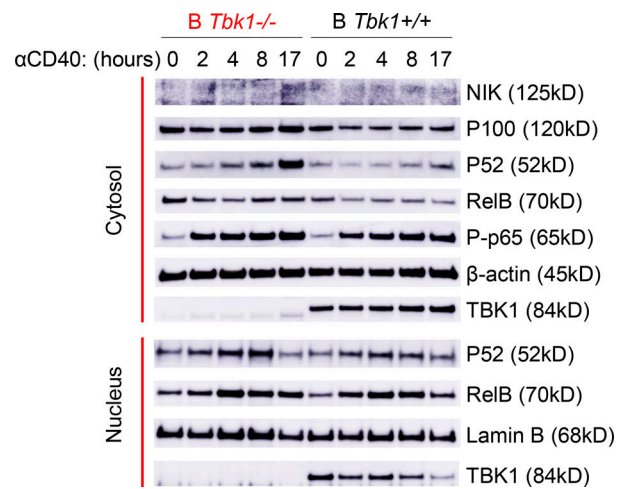


Figure S5. **TBK1 negatively regulates CD40-induced noncanonical NF- κ B signaling in B cells.** Time-course analysis of cytosolic and nuclear fractions of *Tbk1*^{-/-} and B *Tbk1*^{+/+} naive B cells stimulated with 1 μ g/ml anti-CD40 and analyzed by immunoblotting. Representative of two independent experiments. Source data are available for this figure: SourceData FS5.

Provided online is one table. Table S1 provides RNA-seq data of Pre-GC isolated from B *Tbk1*^{-/-} mice versus B *Tbk1*^{+/+} mice on day 9 post-PyNL infection.