

B-cell-specific conditional expression of *Myd88^{p.L252P}* leads to the development of diffuse large B-cell lymphoma in mice

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Introduction

Toll-like receptors (TLRs) belong to a class of pattern-recognition receptors.¹ Ten distinct human TLRs have been described.¹ Except for TLR3, all TLRs require the adaptor protein MYD88 to initiate

downstream signaling.^{1,2} Upon TLR activation, MYD88 is recruited to the Toll/interleukin-1 receptor (TIR) domain of the activated TLR via its own TIR domain.³ MYD88 recruits IRAK1, -2, and -4 to ultimately

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form the myddosome.⁴ IRAK4-mediated phosphorylation of IRAK1 and IRAK2 promotes TRAF6 recruitment,⁵ which subsequently ubiquitylates and activates TAK1,⁶ leading to NF- κ B activation.²

Inappropriate TLR signaling through somatic *MYD88* mutations has been described in numerous hematologic malignancies, such as chronic lymphocytic leukemia (CLL),⁷ Waldenström macroglobulinemia,⁸ and diffuse large B-cell lymphoma (DLBCL).⁹ Specifically, 29% of activated B-cell-type DLBCLs (ABC-DLBCLs), which typically display constitutive NF- κ B activation, carry the p.L265P mutation (position according to the protein accession NP_002459) in the hydrophobic core of the MYD88 TIR domain.⁹ ABC-DLBCL was shown to display *MYD88*^{p.L265P}-dependent NF- κ B activation and STAT3 phosphorylation.⁹ The *MYD88*^{p.L265P} mutation is exceedingly rare in non-ABC-DLBCLs, such as germinal center B-cell (GCB) and primary mediastinal B-cell lymphoma.⁹ Moreover, *MYD88*^{p.L265P}-mutant ABC-DLBCL appears to be dependent on continued *MYD88*^{p.L265P} expression.⁹ Together, these data implicate MYD88-dependent oncogenic NF- κ B signaling as an integral contributor to ABC-DLBCL pathogenesis and as a potential therapeutic target. However, despite the clinical relevance of the *MYD88*^{p.L265P} mutation, no autochthonous mouse model that faithfully mimics this critical genomic aberration has yet been generated.

Materials and methods

Experimental mice

B-cell-specific Cre expression was achieved by using *Cd19*^{Cre},¹⁰ *Aid*^{Cre},¹¹ and *Cd21*^{Cre12} deleter mouse strains (The Jackson Laboratory). The targeting vector depicted in Figure 1A was constructed by using standard techniques. The gene targeting strategy was based on the National Center for Biotechnology Information transcript *NM_010851.2* in which exon 1 contains the translation initiation codon. Wild-type (wt) exons 2-6, including the entire 3' untranslated region were flanked with *LoxP* sites. An additional polyadenylation signal (human growth hormone polyadenylation [hGHpA] signal) has been inserted between the 3' untranslated region and the distal *LoxP* site to prevent downstream transcription of the mutated exon 5. The size of the *LoxP*-flanked region is 4.7 kb. Exons 2-6, including the splice acceptor site of intron 1, have been duplicated and inserted downstream of the distal *LoxP* site. The p.L252P mutation has been introduced into the duplicated exon 5. Positive selection markers were flanked by *Frt* (neomycin resistance; *Neo*^R) and *F3* (puromycin resistance; *Puro*^R) sites and inserted into intron 1 and downstream of the hGHpA, respectively. The targeting vector was generated by using BAC clones from the *C57BL/6J*RPCIB-731 BAC library and was transfected into the *C57BL/6N* embryonic stem (ES) cell line. Targeted clones were isolated by using double positive (*Neo*^R and *Puro*^R) selection, and correct integration was verified by Southern blotting. Upon blastocyst injection, germ line-transmitting transgenic mice were received. The conditional *Myd88*^{p.L252P} allele was obtained after in vivo Flp-mediated removal of the selection markers. This allele expresses the wild-type Myd88 protein, because the presence of the hGHpA cassette downstream of wild-type exon 6 prevents transcription of the mutant exons 2-6. The constitutive *Myd88*^{p.L252P} allele is obtained after in vivo Cre-mediated deletion of wild-type exons 2-6 and hGHpA. This allele expresses the mutant Myd88^{L252P} protein. The remaining recombination sites are located in nonconserved regions of the genome. To generate conditional *BCL2* knockin mice (*Rosa26*^{LSL-BCL2.IRES.GFP}), a *Rosa26* locus-targeting vector was used in which CAGS (cytomegalovirus early enhancer/chicken β actin) promoter-driven expression of the transgene (*BCL2*) and downstream expression of an internal ribosomal entry site-driven green fluorescent protein (IRES-GFP, as a reporter for Cre-mediated recombination) is prevented by a *LoxP*-flanked STOP cassette (supplemental Figure S5, available on the Blood Web site). The transgenic mice expressed the transgenes only after Cre-mediated excision of the *LoxP*-flanked STOP cassette. The targeting vector was electroporated into ES cells (BRUCE4), which were screened

for correct integration by standard Southern blot methods. Correctly targeted ES cells were used to generate chimeras, which were backcrossed onto a pure *C57BL/6N* background and examined for germ line transmission.

The *E μ :Myc* model has been described earlier and was used, as previously described.¹³ The *E μ :Tcl1* model of CLL was described earlier.¹⁴ All the experiments that involved the breeding and/or treatment of rodents were approved by the local animal care committee and the relevant authorities (Landesamt für Natur, Umwelt und Verbraucherschutz Nordrhein-Westfalen, AZ: 84-02.04.2014.A146/A083).

Transplantation experiments

Animal protocols based on the *E μ :Myc* transgenic mouse model as used in this study were approved by the local governmental review board (Landesamt Berlin) and conformed to regulatory standards. Isolation, retroviral infection, transplantation of *E μ :Myc* transgenic fetal liver cells (ie, hematopoietic stem cells), and subsequent monitoring of the recipient mice regarding lymphoma onset were performed as previously described.¹⁵ Specifically, numerous independently isolated fetal liver cell populations were stably transduced with a murine stem cell virus-based *Myd88*^{p.L252P}-IRES-GFP retrovirus or a murine stem cell virus-empty GFP construct as a control before their propagation in recipient mice.

Immunohistochemistry

Formalin-fixed paraffin-embedded (FFPE) murine samples were sliced at 2 to 4 μ m. Sections were stained with hematoxylin and eosin (H&E) and antibodies against Ki-67 (Cell Marque), B220 (RA3-6B2BD), p65 (C-20; Santa Cruz), Irf4 (M-17; Santa Cruz Biotechnology), Bcl6 (C-19; Santa Cruz Biotechnology), and Cd138 (553712; BD). Staining intensities were scored by 2 independent observers who used a 4-tier scale. FFPE human samples were sliced at 2 to 4 μ m. Sections were stained with H&E and antibodies against CD10 (NCL-L-CD10-270; Novocastra), BCL6 (M7211; Dako), IRF4 (M7259; Dako), BCL2 (M0877; Dako), and CD138 (138M-16; Cell Marque).

Sequence analysis of immunoglobulin genes and cloning of polymerase chain reaction products

RNA was extracted from sections of frozen biopsies with TRI Reagent (Sigma, Taufkirchen, Germany). One microgram of RNA was converted to complementary DNA with QuantiTect Reverse Transcription Kit (Qiagen, Hilden, Germany). Mouse *IgH* rearrangements were analyzed by using previously published methods.¹⁶ Each polymerase chain reaction (PCR) was analyzed on a Qiaxcel Advanced Instrument (Qiagen) by using ScreenGel Software v1.2. Direct sequencing was performed with the ABI Prism Dye Terminator Cycle Sequencing Ready Reaction Kit v3.1 (Life Technologies, Darmstadt, Germany) on an ABI 3130 sequencer (Applied Biosystems, Foster City, CA). Sequences were analyzed by using 4Peaks Software v1.7.2 (The Netherlands Cancer Institute, Amsterdam, The Netherlands), Lasergene software (DNASar, Madison, WI), and by manual review. Sequences were compared with mouse germ line *Ig* gene sequences with International ImMunoGeneTics database.¹⁷ In addition, the PCR products of 1 case were cloned with the TOPO TA Cloning Kit (Life Technologies), which resulted in 32 sequences. To generate genealogic trees, sequence analysis focused on the *IGHV* segment, starting within FR1. To exclude possible Taq errors, only mutations shown more than once were considered for analysis (see supplemental Data).

Results

To assess the role of *MYD88*^{p.L265P} in B-cell lymphomagenesis, we generated a conditional *Myd88*^{p.L252P} allele (*Myd88*^{c-p.L252P}) that is expressed from the endogenous locus upon Cre-mediated recombination (Figure 1A). Murine *Myd88*^{p.L252P} is at the orthologous position of human *Myd88*^{p.L265P}. To verify Cre-inducible *Myd88*^{p.L252P} expression, we derived murine embryonic fibroblasts (MEFs) from wild-type

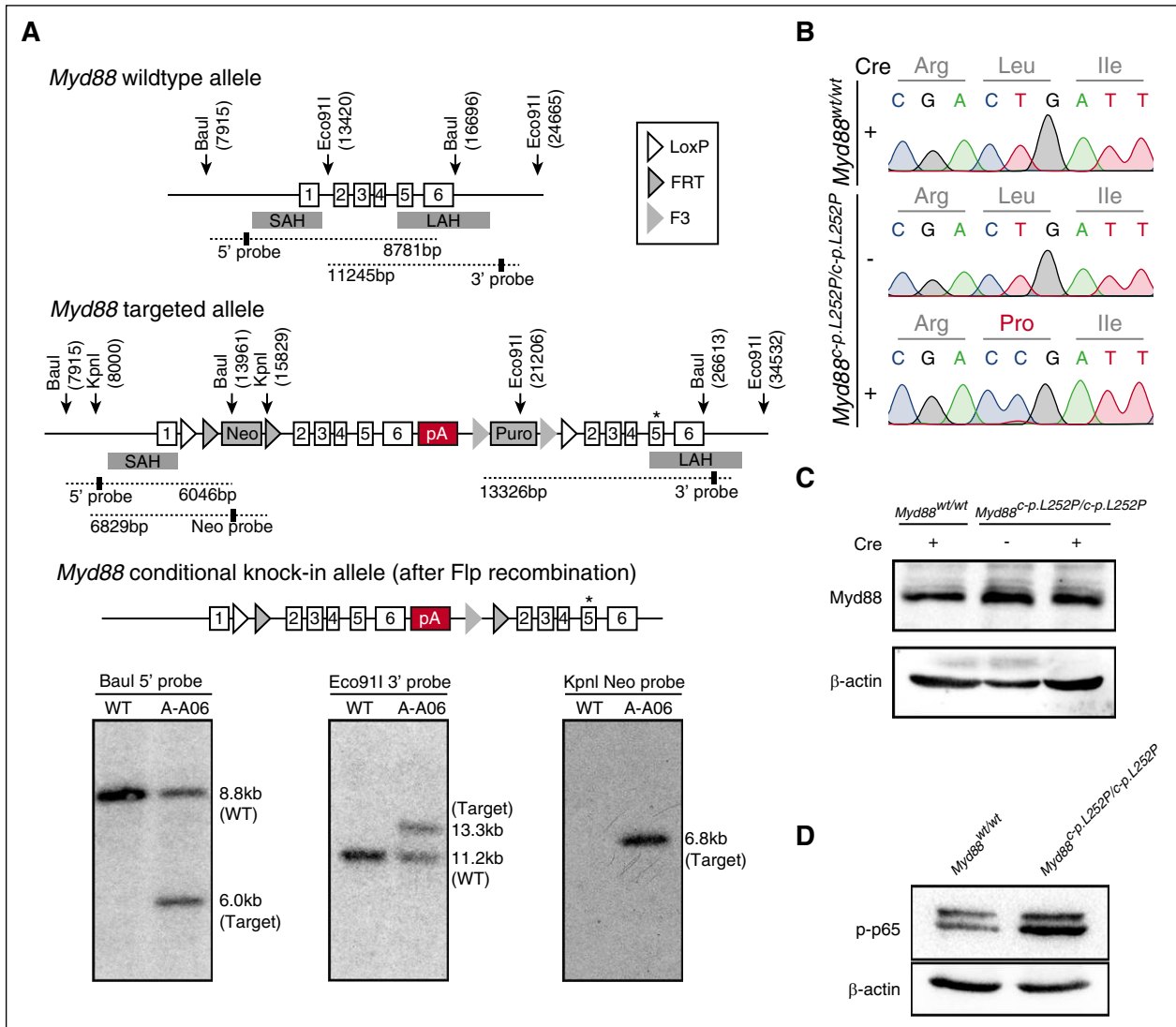


Figure 1. Construction of a conditional *Myd88*^{p.L252P} allele. (A) Targeting of the *Myd88* locus in *C57BL/6N* Tac ES cells. The endogenous *Myd88* locus was targeted with the linearized vector described in the supplemental Data. The targeted allele before (top panel) and after Flp-mediated recombination of FRT and F3 sites (bottom panel) is schematically depicted. The Southern blots of Baul, Eco911, and KpnI digested genomic DNA probed with a 5', a 3', and a Neo probe, respectively, are shown below the schematic illustration of the targeting strategy. Positions of restriction sites and probes are shown in the schematic drawing above. (B) *Myd88*^{p.L252P} mRNA is expressed upon Cre-mediated recombination in MEFs. *Myd88*^{wt/wt} and *Myd88*^{c-p.L252P/c-p.L252P} MEFs were isolated. RNA was isolated from both cell lines before LentiCre application (Sanger sequencing chromatograms, top and middle panels) and the *Myd88* mRNA sequence was determined after reverse transcription. The wild-type sequence was recovered from both cell lines. After LentiCre application and puromycin selection, only the p.L252P sequence could be recovered from *Myd88*^{c-p.L252P/c-p.L252P} MEFs (Sanger sequencing chromatogram, bottom panel). (C) The *Myd88*^{p.L252P} isoform is expressed in *Myd88*^{c-p.L252P/c-p.L252P} MEFs after LentiCre-mediated recombination. *Myd88*^{wt/wt} and *Myd88*^{c-p.L252P/c-p.L252P} MEFs were LentiCre exposed and puromycin selected, as in (B). Whole-cell lysates were separated on sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) and blotted onto polyvinylidene difluoride membranes before Myd88 and β-actin, which served as loading controls, and were visualized by immunoblotting. Both *Myd88*^{wt} and *Myd88*^{p.L252P} proteins were expressed at equal levels. (D) Conditional LentiCre-mediated *Myd88*^{p.L252P} expression leads to p65 Ser-536 phosphorylation. *Myd88*^{wt/wt} and *Myd88*^{c-p.L252P/c-p.L252P} MEFs were transduced with LentiCre and puromycin selected, as in (B). Upon selection, cells were lysed, proteins were separated on SDS-PAGE, and pSer-536 p65 was visualized by immunoblot. SAH, short arm of homology; LAH, long arm of homology; pA, polyadenylation signal sequence.

and *Myd88*^{c-p.L252P/c-p.L252P} mice. By using Sanger sequencing, we detected expression of *Myd88*^{wt} messenger RNA (mRNA) in both *Myd88*^{wt/wt} and *Myd88*^{c-p.L252P/c-p.L252P} MEFs, indicating lack of background expression of *Myd88*^{p.L252P} in the absence of Cre recombinase. In contrast, cytomegalovirus promoter-driven Cre expression mediated by lentiviral transduction of *Myd88*^{c-p.L252P/c-p.L252P} MEFs induced recombination in the *Myd88* locus, leading to expression of *Myd88*^{p.L252P}-mutant mRNA (Figure 1B). *Myd88* protein abundance was unaffected by stable Cre expression, indicating that the *Myd88*^{p.L252P} isoform is expressed at endogenous levels (Figure 1C). This *Myd88*^{p.L252P} expression translated

into robust p65 Ser-536 phosphorylation, indicating activation of NF-κB signaling (Figure 1D).

To determine whether B-cell-specific *Myd88*^{p.L252P} expression promotes lymphomagenesis, we compared *Cd19*^{Cre/wt};*Myd88*^{c-p.L252P/wt} (hereafter M-Cd19) and *Cd19*^{Cre/wt} (hereafter Cd19) mice. The mice were longitudinally monitored by magnetic resonance imaging (MRI). Although no gross lymphoproliferative disease (LPD) was detectable in Cd19 mice, M-Cd19 mice developed splenomegaly and occasional lymphadenopathy starting at the age of ~60 weeks (Figure 2B). Similar MRI morphologic signs of LPD were observed when B-cell-specific *Myd88*^{p.L252P} expression was induced by Cre

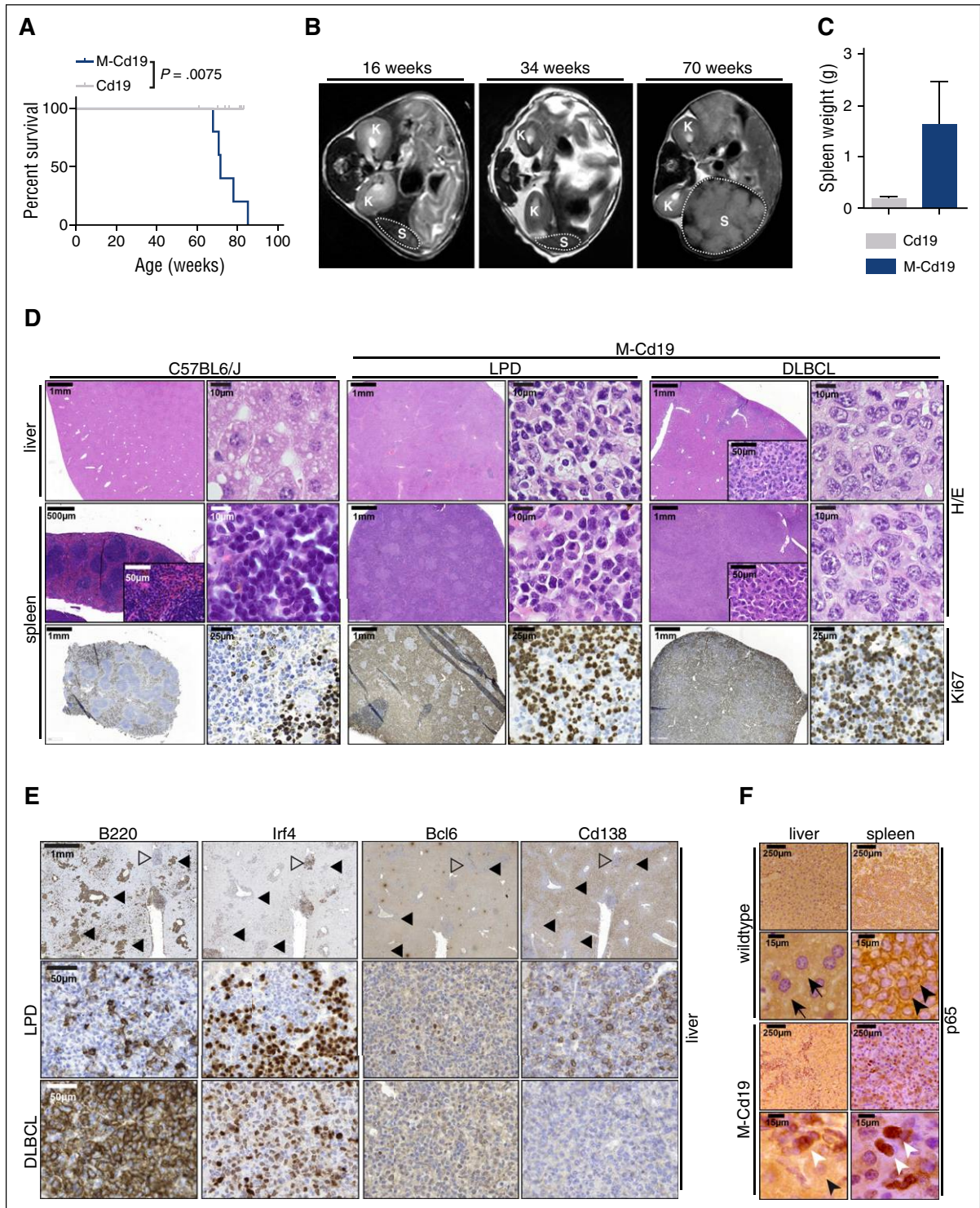


Figure 2. B-cell-specific *Myd88^{p.L252P}* expression drives lymphoproliferation and lymphomagenesis in vivo. (A) B-cell-specific expression of *Myd88^{p.L252P}* significantly reduces overall survival in vivo. Kaplan-Meier curves illustrate the overall survival of M-Cd19 mice. M-Cd19 mice display a significantly reduced survival compared with the respective controls (log-rank test). (B) Serial MRI scans in 16-, 34-, and 70-week-old M-Cd19 mice revealed the occurrence of splenomegaly in 70-week-old M-Cd19 mice. (C) M-Cd19 mice display splenomegaly at the time of death. Preterminal M-Cd19 and Cd19 mice were sacrificed and spleen weight was recorded. Bars represent the average ($n = 3$); error bars represent standard deviations. (D) M-Cd19 mice develop lymphoproliferative disease and occasional lymphoma. The top panels show H&E staining of spleens and livers isolated from *C57BL/6* and M-Cd19 mice at the time of death. Although the organ architecture appeared normal in *C57BL/6* wild-type mice, the architecture of spleens isolated from M-Cd19 mice was largely disrupted by infiltration of small mature lymphocytes (LPD columns) or large blastoid cells (DLBCL columns). The bottom panel shows the partial and complete disruption of the spleen by infiltrates with high proliferative indices. (E) Immunohistochemical characterization of the liver infiltrates of M-Cd19 mice. Areas of infiltrates morphologically resembling DLBCL (marked with solid triangles) showed a homogeneous staining pattern of B220 and Irf4 positivity, whereas they were negative for Bcl6 and Cd138 (bottom panel). Infiltrated areas of small, mature lymphocytes (marked with open triangles) displayed a more heterogeneous staining pattern of positive and negative cells for B220, Irf4, and Cd138, whereas staining was largely negative for Bcl6 (middle panel). (F) The lymphoma cells infiltrating spleens and livers of M-Cd19 mice displayed a largely nuclear localization of p65, indicating NF- κ B activation. Black arrows indicate hepatocytes, black arrowheads indicate cytoplasmic p65 staining in splenic lymphocytes in *C57BL/6* wild-type mice, and white arrowheads indicate nuclear p65 staining in lymphoma cells in M-Cd19 mice.

expression that was driven off the *Aid* promoter (hereafter M-Aid), which is active in GCB cells, during both T cell–dependent and –independent immune responses (supplemental Figure 1A).^{11,18} Furthermore, *Cd21* promoter-driven Cre, which is expressed when immature transitional B cells differentiate into mature long-lived peripheral B cells,¹² induced an LPD in *Cd21^{Cre/wt};Myd88^{c-p.L252P/wt}* mice (hereafter M-Cd21) that was indistinguishable from that observed in M-Cd19 mice (supplemental Figure 1A).

The MRI morphologic occurrence of lymphoproliferative lesions translated into a significantly ($P = .0075$) reduced overall survival of M-Cd19 mice (median survival, 501 days) compared with Cd19 mice (median survival, not reached in 10 mice observed for 427 to 581 days [average, 539 days]) (Figure 2A). Similar data were obtained when Cre expression was driven off the *Aid* or *Cd21* promoter in M-Aid and M-Cd21 mice (median survival, 574 and 610 days, respectively) (supplemental Figure 1B). We noted that the majority of M-Cd19, M-Aid, and M-Cd21 mice included in this study displayed splenomegaly (4 of 4, M-Cd19; 2 of 3, M-Aid; and 2 of 3, M-Cd21), occasionally accompanied by macroscopic lymphadenopathy (1 of 4, M-Cd19; 2 of 3, M-Aid; and 1 of 3, M-Cd21). The strong oncogenic potential of the *Myd88^{p.L252P}* mutation was also verified in an alternative mouse model of aggressive B-cell lymphoma in which *Myd88^{p.L252P}* dramatically shortened tumor onset driven by oncogenic *Myc* (*Eμ:Myd88^{p.L252P}* transgenic fetal liver cells transplanted into lethally irradiated recipient mice, stably transduced with either empty vector [n = 40; median onset not reached] or *Myd88^{p.L252P}* [n = 12; median onset, 36 days]; log-rank Mantel-Cox $P < .0001$) (supplemental Figure 2).

To characterize the disease occurring in M-Cd19 mice, we initially performed histologic examinations. H&E staining of spleens and livers of M-Cd19, M-Aid, and M-Cd21 mice revealed infiltrates of lymphoid cells in these organs. In particular, the architecture of the spleens of these mice was largely disrupted by these infiltrates (4 of 4, M-Cd19; 3 of 3, M-Aid; 3 of 3, M-Cd21). Of note, we did not detect any infiltration of the bone marrow (n = 3 per genotype; supplemental Figure 3). Further analyses of the splenic and hepatic infiltrates revealed that these lesions constituted a largely monomorphic lymphoid cell population with indolent appearance, consistent with an LPD. Intriguingly, in a subset of mice (1 of 4, M-Cd19; 1 of 3, M-Aid; and 1 of 3, M-Cd21), diffuse large lymphoid infiltrates resembling DLBCL were detectable (Figure 2D; supplemental Figure 1D-E). Of note, the Ki-67 indices did not substantially differ between areas with LPD and DLBCL morphology (Figure 2D). To determine the clonality of these infiltrates, we next performed Southern blot analyses to detect clonal immunoglobulin (Ig) rearrangements. We were not able to detect clonal Ig rearrangements in genomic DNA isolated from areas with LPD-like appearance that were detected in M-Cd19, M-Aid, and M-Cd21 mice (supplemental Figure 4A). Strikingly, oligoclonal Ig rearrangement patterns were detected in DNA isolated from infiltrates that displayed DLBCL-like morphology in M-Cd19, M-Aid, and M-Cd21 mice (indicated by white triangles in supplemental Figure 4A). DNA isolated from spleens of an *Eμ:Tcl1*-driven CLL mouse model served as a positive control for the development of a clonal lymphoid neoplasm.¹⁴ Altogether, our data strongly indicate that B-cell–specific *Myd88^{c-p.L252P}* expression largely leads to an LPD, with only occasional development of mono-/oligoclonal lymphoma.

To further characterize the LPD and lymphomas, we next performed immunohistochemistry in liver infiltrates of M-Cd19 mice. As shown in Figure 2E, the areas with DLBCL-like morphology displayed a strikingly uniform staining pattern. These lesions were B220 and Irf4 positive and were Bcl6 and Cd138 negative, resembling the immunophenotype of a post-GCB lymphoid malignancy.¹⁹ In contrast, the areas of LPD displayed a rather heterogeneous staining pattern for

B220, Irf4, and Cd138 and were largely negative for Bcl6, further underscoring the nonclonal characteristics of these lesions. We noted that areas of LPD and DLBCL-like morphology in M-Aid and M-Cd21 mice displayed similar staining patterns (supplemental Figure 1F-G).

To further interrogate the nature of the DLBCL-like lesions, we next used immunohistochemistry to assess subcellular NF-κB localization, because constitutive NF-κB activation, indicated by nuclear enrichment, is a hallmark feature of ABC-DLBCL.⁹ These experiments revealed a uniform nuclear staining of the NF-κB subunit p65 in the DLBCL-like lesions detected in spleens and livers of M-Cd19, M-Aid, and M-Cd21 mice (Figure 2F; supplemental Figure 1C). Given that ABC-DLBCL, which is characterized by constitutive NF-κB activation, carries the p.L265P mutation in 29% of the cases,⁹ and given that the clonal lymphoproliferation we observed in M-Cd19 mice displayed morphologic features of DLBCL, we hypothesized that the B-cell–specific *Myd88^{c-p.L252P}* expression drives ABC-DLBCL development in our model.

To cross-validate our observations, we analyzed *MYD88* mutation frequency in human DLBCLs (n = 24, GCB-DLBCL; n = 21, ABC-DLBCL). We performed targeted sequencing of FFPE-extracted DNA by multiplex PCR covering the *MYD88*, *ATM*, *BTK*, *CD79B*, *DDX3X*, *FBXW7*, *MAPK1*, *NOTCH1*, *PIK3CA*, *PIK3CD*, *PTEN*, *PTPN6*, *SF3B1*, *TP53*, and *XPO1* genes (Figure 3A). DLBCL subtypes were determined based on immunohistochemical stains for CD10, BCL6, and IRF4 following the Hans algorithm (Figure 3B).¹⁹ The non-GCB of Hans algorithm was designated as “ABC” in our study. Confirming previously published data, we found a substantial overrepresentation of *MYD88* and *CD79B* mutations in ABC-DLBCLs compared with GCB-DLBCLs (Figure 3A,C; supplemental Table 1).^{9,20,21} Furthermore, the anti-apoptotic oncogene *BCL2* is frequently deregulated in human DLBCL.^{22,23} Whereas the chromosomal translocation t(14;18)(q32;q21), which juxtaposes *BCL2* and the *IGHV* enhancer, is commonly observed in GCB-DLBCL,²³ focal copy number gains are more commonly observed in ABC-DLBCL (~30% to 40%).²²⁻²⁵ Indeed, mining of the whole-genome sequencing data on DLBCL from the German International Cancer Genome Consortium Molecular Mechanisms in Malignant Lymphoma by Sequencing Project (<https://icgc.org/> and C.L., M.S., and R.S., manuscript in preparation) revealed that 6 DLBCLs carry the *MYD88^{p.L265P}* mutation. Remarkably, all of these samples belonged to the ABC-DLBCL subtype and carried a copy number gain of 18q detected by both fluorescent in situ hybridization with a *BCL2*-specific probe and somatic copy number variant mapping.^{26,27} A critical role for *BCL2*, particularly in activated B cells, is further corroborated by published transcriptome analyses. Specifically, *BCL2* mRNA was not expressed in GCB cells but was massively induced during activation of peripheral blood B cells.²⁸ Consistent with this, the majority of ABC-DLBCLs displayed *BCL2* mRNA levels more than fourfold higher than GCB cells.²⁸ Furthermore, exogenous expression of *MYD88^{p.L265P}* in B cells was recently shown to induce the accumulation of self-reactive B cells in vivo only when apoptosis was opposed by *Bcl2* overexpression.²⁹ Additional circumstantial evidence suggesting that ABC-DLBCL in particular might depend on functional *BCL2* activity could be derived from recently published, large-scale in vitro drug sensitivity profiling experiments. Specifically, re-assessment of a data set published by Stegmeier and colleagues revealed that *MYD88*-mutant DLBCLs in particular display an actionable *BCL2* addiction, which could be exploited by the targeted high-affinity *BCL2* inhibitor ABT-263 (supplemental Figure 5A-B).³⁰ The BTK inhibitor ibrutinib was recently shown to synergistically interact with the *BCL2* family inhibitor navitoclax (ABT-263) in killing ABC-DLBCL cells.³¹ We

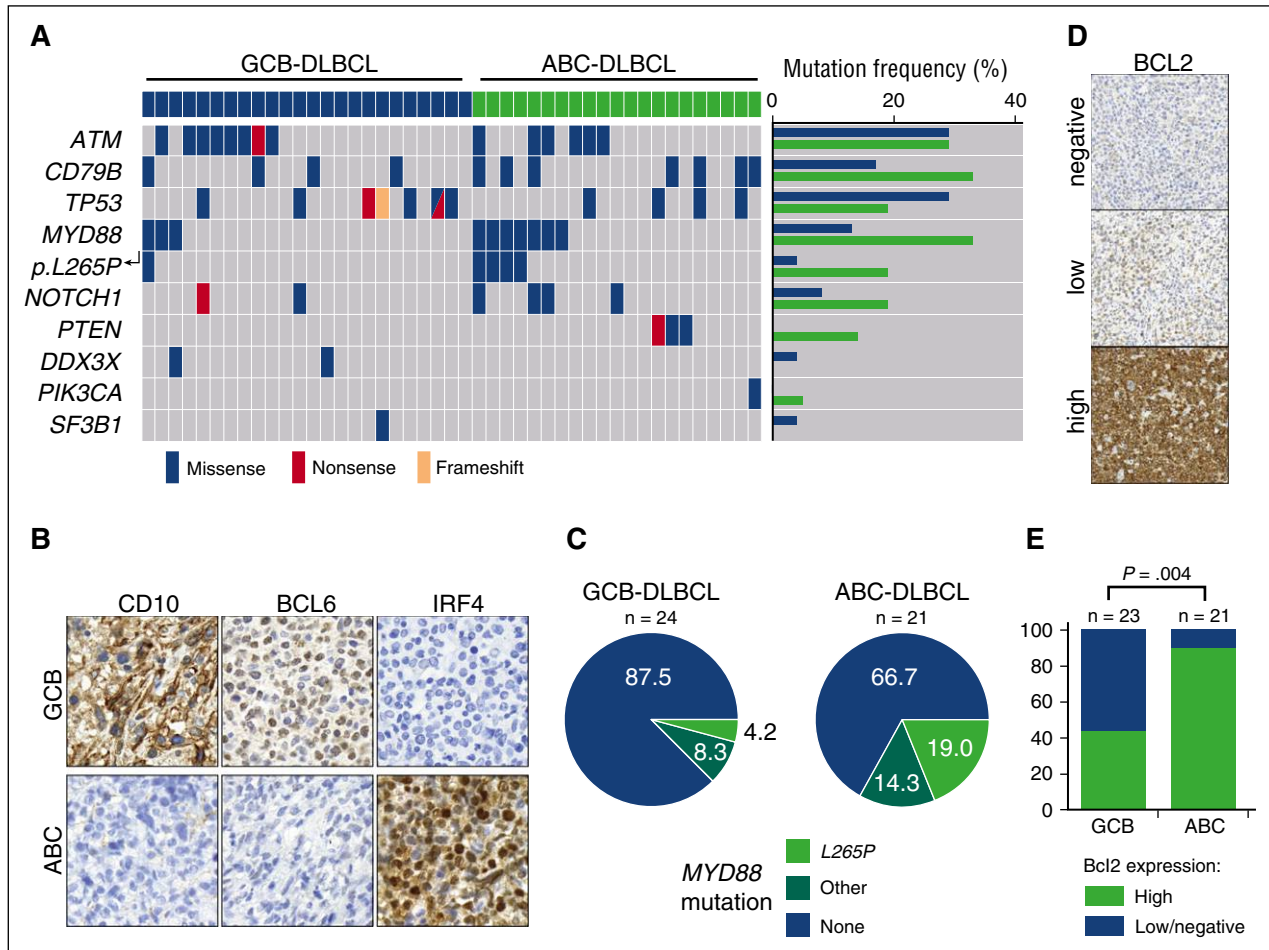


Figure 3. *MYD88* mutations and high expression of *BCL2* are enriched in ABC-DLBCL. (A) *MYD88* mutations are substantially enriched in human ABC-DLBCL. Human DLBCLs were stratified as ABC- or GCB-DLBCL following the Hans algorithm, as depicted in (B). After immunohistochemistry-based stratification, DNA was isolated from tissue sections and subjected to targeted deep sequencing by a multiplex PCR, which covered the *ATM*, *BTK*, *CD79B*, *DDX3X*, *FBXW7*, *MAPK1*, *MYD88*, *NOTCH1*, *PIK3CA*, *PIK3CD*, *PTEN*, *PTPN6*, *SF3B1*, *TP53*, and *XPO1* genes. *MYD88* mutations per se, and particularly the *MYD88*^{L265P} mutation were substantially enriched in ABC-DLBCL. Similarly, *CD79B* and *PTEN* mutations were enriched in ABC-DLBCL. (C) Distribution of *MYD88* mutations detected in human DLBCL samples is shown in pie charts. The samples were classified into high and low or negative expression of *BCL2*, as shown in (D). (E) High protein expression levels of *BCL2* are significantly enriched in ABC-DLBCL (Fisher's exact test).

validated these observations in our cohort of DLBCL samples by using immunohistochemistry to assess *BCL2* expression. As shown in Figure 3D-E, high-level *BCL2* expression was significantly enriched in ABC-DLBCL patients compared with patients who had GCB-DLBCL. Altogether, these genomic, transcriptomic, immunohistochemical, and functional data strongly suggest that enhanced *BCL2* activity plays a central pathomechanistic role in the development of ABC-DLBCL. On the basis of these considerations, we next asked whether lymphoma development in M-Cd19 mice could be accelerated by increased *BCL2* gene dosage in vivo. To mimic *BCL2* amplification in vivo, we generated a novel conditional allele in which *BCL2*.*IRES*.*GFP* was targeted into the *Rosa26* locus. *BCL2* expression was prevented by the insertion of a *LoxP*.*STOP*.*LoxP* cassette upstream of the translation-initiating codon (supplemental Figure 5C). By using this allele, we generated *Cd19*^{Cre/wt};*Rosa26*^{LSL*BCL2*.*IRES*.*GFP*/wt};*Myd88*^{Cre-p.L252P/wt} mice (hereafter M-B-Cd19), in which *Cd19*-driven Cre mediates the excision of the *LoxP*.*STOP*.*LoxP* cassette from the *Rosa26* locus, leading to CAGS-promoter-driven *BCL2* overexpression in addition to *Myd88*^{p.L252P} expression. MRI scans revealed that M-B-Cd19 mice developed splenomegaly substantially earlier than M-Cd19, M-Aid, or M-Cd21 mice (Figures 2B and 4B; supplemental Figure 1A). Furthermore, M-B-Cd19 mice displayed a significantly reduced overall

survival (median survival, 179 days) compared with M-Cd19, M-Aid, and M-Cd21 mice (long-rank $P = .0008$, $P = .0001$, and $P < .0001$, respectively) (Figure 4A). Of note, all of the M-B-Cd19 mice displayed excessive lymphadenopathy and/or splenomegaly at the time of death (splenomegaly in 7 of 8 mice; lymphadenopathy in 6 of 8 mice).

To further characterize the nature of the lymphadenopathy that we observed in M-B-Cd19 mice by MRI, we next performed histologic examination of infiltrated organs (livers, lymph nodes, and spleens). Morphologic assessment of the splenic, hepatic, and lymph node lesions detected in M-B-Cd19 mice revealed that these infiltrates consisted almost entirely of diffuse large lymphoid cells with blastoid to plasmablastoid appearance (Figure 4C). Intriguingly, plasmablasts have been implicated as the cell of origin of ABC-DLBCL.²⁴ In marked contrast to the phenotype observed in M-Cd19 mice, in which the bulk of the lesions consisted of LPD-like infiltrates (Figure 2D), areas of LPD were only rarely detectable in M-B-Cd19 mice. Furthermore, the Ki-67 index of the lymphoid lesions in M-B-Cd19 mice was significantly higher compared with lesions that emerged in M-Cd19 mice (median, $71.57\% \pm 3.65\%$ in M-B-Cd19 mice vs $28.27\% \pm 2.40\%$ in M-Cd19 mice) (Figure 4F). The Ki-67 indices observed in M-B-Cd19 mice

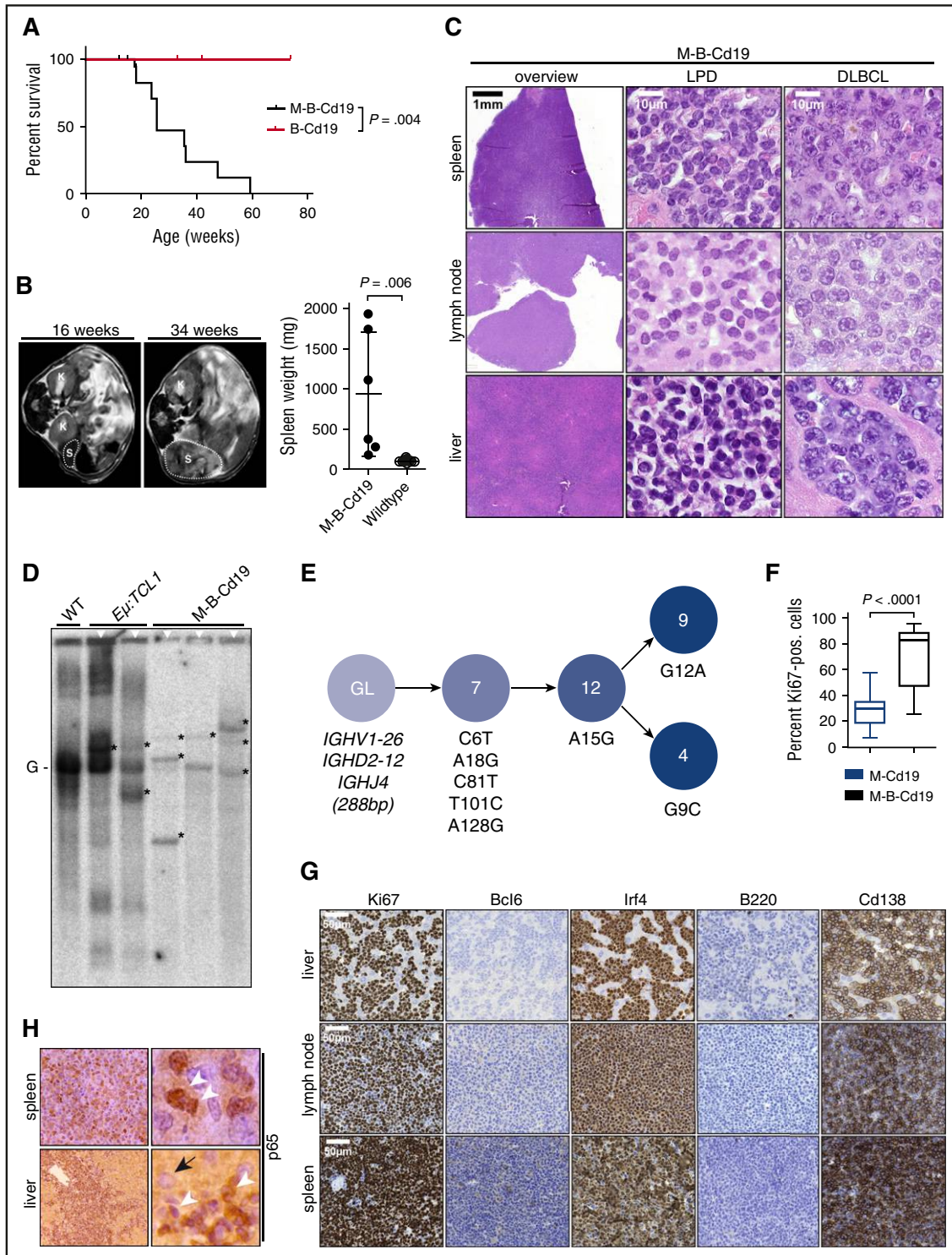


Figure 4. Combination of B-cell-specific *BCL2* overexpression and *Myd88^{p.L252P}* expression drives ABC-DLBCL development in vivo. (A) Kaplan-Meier curve illustrates overall survival of M-B-Cd19 mice. M-B-Cd19 mice display a significantly reduced survival compared with the respective controls (log-rank test). (B) M-B-Cd19 mice develop splenomegaly at 25 to 35 weeks. MRI scans and quantification of spleen weights at autopsy are shown (two-tailed Student *t* test). (C) The architecture of spleens and lymph nodes isolated from M-B-Cd19 mice was completely disrupted by a homogeneous infiltrating population of large lymphoblastoid cells (DLBCL column). Small areas of more heterogeneous lymphoid cell populations consisting of small lymphoid cells were occasionally detectable (LPD column). The livers of M-B-Cd19 mice were diffusely infiltrated by large blastoid cells with DLBCL-like morphology. (D) Variable-diversity-joining-recombination analysis by Southern blot analysis revealed the presence of clonal populations in tumors of M-B-Cd19 mice. Tumors from leukemic Eμ:*TCL1* mice were used as oligoclonal controls. The germ line configuration is also present in the WT controls and is depicted as "G." Asterisks indicate clonal rearrangements. Samples of mice with lymphoma detected by histologic analyses are marked with white triangles (top). (E) Genealogic trees of murine DLBCL isolated from M-B-Cd19 mice. *Igh* rearrangements were cloned and individual clones were sequenced. Analysis focused on the *Ighv* segments starting with FR1. Genealogic tree was derived from 32 sequences. The tree-like structures demonstrate ongoing somatic hypermutation. Numbers in circles show number of sequence reads; numbering of mutations is in regard to the germ line sequence of *Ighv1-26*. (F) The Ki-67 index of DLBCL lesions from M-B-Cd19 mice was significantly higher than that in M-Cd19 mice ($P = 2.27 \times 10^{-12}$, two-tailed Student *t* test; $n = 3$ mice per genotype; 20 fields of view per lesion). (G) Lymphoma infiltrates of M-B-Cd19 mice were immunohistochemically analyzed and stained positive for Irf4 and Cd138 and negative for B220 and Bcl6. The Ki-67 stainings are quantified in (F).

matched those typically detected in aggressive human lymphoma.³² Leukemic effusion was detectable only in M-B-Cd19 mice (supplemental Figure 6).

Consistent with the monomorphic appearance of the blastoid lesions detected in M-B-Cd19 mice, clonality analyses using Southern blot to detect Ig rearrangements revealed that these infiltrates were of clonal or oligoclonal origin (3 of 3 lesions isolated from independent mice) (Figure 4D). Expression of the mutant *Myd88* allele in the lymphomas was verified by reverse transcriptase PCR sequencing analysis (supplemental Figure 7). To ask whether the oligoclonal lymphomas underwent somatic hypermutation, as would be expected in the case of ABC-DLBCL, we analyzed a clonal lesion for *Igh* rearrangements by direct sequencing. As shown in Figure 4E, we could detect a clonal *Ighv1-26*, *Ighd2-12*, *Ighj4* rearrangement. The sequence shows an in-frame rearrangement with mutations but no stop-codons. Thus, this rearrangement is potentially functional. Next, we cloned the amplified *Ighv* gene rearrangements from this sample and sequenced 32 subclones to investigate intraclonal diversity. All analyzed sequences shared 5 mutations compared with the most homologous germ line sequence (*Ighv1-26*). Further mutations could be detected in several subclones (Figure 4E), demonstrating intraclonal diversity for this analyzed case.

Next, we used immunohistochemistry to further characterize the lymphomas isolated from M-B-Cd19 mice. As shown in Figure 4G and supplemental Figure 8, the lesions stained uniformly positive for Irf4 and were entirely negative for Bcl6, consistent with the somatic hypermutation detected in Figure 4E and further corroborating the post-germinal center (GC) nature of the infiltrates.²⁴ Furthermore, the lymphomas lost B220 expression and stained strongly positive for Cd138, indicating ABC origin of these lesions.³³⁻³⁶ This is in line with analyses of human DLBCL, which indicate that GC-experienced Cd138-positive non-GC-DLBCLs are typically of plasmablastoid differentiation.³⁷⁻³⁹ In agreement with constitutive NF- κ B activation, which is typically observed in human ABC-DLBCL, we found almost exclusive nuclear p65 staining in the lymphomas detected in M-B-Cd19 mice (Figure 4H).

Although *MYD88* mutations have not been detected in human multiple myeloma,^{8,40-42} the strong Cd138 positivity of the lymphomas growing in M-B-Cd19 mice suggested the possibility that these lesions might be of plasmacellular origin.^{8,40-42} However, despite considerable effort, we were unable to detect a plasma cell expansion in the bone marrow of M-Cd19, M-Aid, M-Cd21, and M-B-Cd19 mice. More importantly, we did not find lymphoma infiltration of the bone marrow in M-Cd19, M-Aid, and M-Cd21 mice, and only 1 of 4 M-B-Cd19 mice showed an isolated lymphoma infiltrate that did not display signs of plasmacellular differentiation (supplemental Figures 3 and 9). In line with a lack of a consistent bone marrow infiltration, which is an obligatory feature of Waldenström macroglobulinemia and is frequently observed in multiple myeloma, we did not detect mono- or oligoclonal gammopathy in either M-Cd19 or M-B-Cd19 mice (supplemental Figure 10). However, we did detect an increased abundance of the Ig fraction in M-Cd19 mice compared with wild-type controls by using serum electrophoresis (upper and middle panels of supplemental Figure 10). This gammopathy was even more prominent in M-B-Cd19 mice (lower panel of supplemental Figure 10). The overall polyclonality strongly suggests a non-neoplastic origin of the gammopathy observed in M-Cd19 and M-B-Cd19 mice. Thus in summary, the morphologic appearance, as well as the immunophenotype of the lesions observed in M-B-Cd19 mice strongly suggests that these lymphomas represent ABC-DLBCL.

Discussion

Here, we generated a novel conditional *Myd88*^{p.L252P} allele, which is at the orthologous position of the human *Myd88*^{p.L265P} mutation and is expressed from the endogenous locus upon Cre-mediated recombination. We chose to design an allele that allows expression outside the natural genomic context to most faithfully mimic the situation in human malignancies. B cell-specific expression of this mutant leads to an LPD and occasional (~30%) transformation into an aggressive lymphoma that morphologically and immunophenotypically resembles human ABC-DLBCL. Interestingly, development of LPD and lymphoma occurred at a similar rate, regardless of the Cre allele. We specifically used a pan B-cell Cre allele (*Cd19*^{Cre}), a Cre allele that is active in GCB cells (*Aid*^{Cre}), and a Cre allele that is expressed when immature transitional B cells differentiate into mature peripheral B cells (*Cd21*^{Cre}). These data might indicate that *Myd88*^{p.L252P} expression leads to a block in B-cell differentiation after the GC reaction. Consistent with this postgerminal nature of *Myd88*^{p.L252P}-expressing cells, we find that the resulting lymphomas are consistently Bcl6 negative and express Irf4 (Figure 2E; supplemental Figure 1F-G). These observations are in line with data from human lymphoma patients, which suggests that oncogenic *MYD88* mutations are enriched in post-GC malignancies, such as ABC-DLBCL, Waldenström macroglobulinemia, and the *IGHV*-mutant GC-experienced subset of CLL patients.^{7-9,24,43-45} Furthermore, crossing in an additional allele that allows conditional *BCL2* expression from the *Rosa26* locus leads to 100%-penetrant development of ABC-DLBCL.

Both *MYD88*⁴⁶⁻⁴⁸ and *BCL2*^{24,49-51} have been shown to be critical for transition through the GC. Thus, it is conceivable that mice displaying B-cell-specific *Myd88*^{p.L252P} expression (M-Cd19) or *Myd88*^{p.L252P} expression in combination with *BCL2* overexpression (M-B-Cd19) might accumulate post-GCB cells. Consistent with this, we find a polyclonal gammopathy in both M-Cd19 and M-B-Cd19 mice (supplemental Figure 10).

MYD88 is frequently mutated in ~29% of ABC-DLBCL cases.⁹ However, *MYD88* mutations are even more common in Waldenström macroglobulinemia, in which these alterations are detected in ~90% of the cases.^{8,52} Thus, it is somewhat surprising that our *Myd88*^{p.L252P}-expressing mice develop clonal ABC-DLBCL rather than Waldenström macroglobulinemia. One possible explanation might be the different genomic landscapes of these distinct B-cell neoplasias. For instance, mutations in the carboxyl terminal domain of the chemokine receptor CXCR4 have been detected in a substantial portion of patients with Waldenström macroglobulinemia (~30% of the patients), whereas they are absent in ABC-DLBCL.^{2,9,53} Furthermore, somatic mutations affecting *CD79A* and *CD79B* are present in ~20% of human ABC-DLBCLs, whereas these alterations are rare or absent in other lymphoma entities.^{20,24} Thus, it will be interesting to combine our novel *Myd88*^{p.L252P} allele with additional B-cell-specific genetic alterations to determine whether further manipulation might enable a plasmacytoid differentiation.

Our novel models represent a preclinical platform that could be exploited for the in vivo development and validation of novel therapeutic approaches to treat ABC-DLBCL, which remains a difficult-to-treat clinical entity.^{54,55} For instance, it will be interesting to see whether combined targeting of the apoptotic machinery and the myddosome, through *BCL2* inhibition and *IRAK4* repression, will result in synergistic toxicity in *Myd88*-driven ABC-DLBCL.⁵⁵ Our models might also serve to delineate the potential role of NF- κ B and MAPKK6 signaling as actionable pathways downstream of the myddosome

complex in ABC-DLBCL.⁵⁶ For this purpose, it will be interesting to assess the cytotoxic potential of TAK1, I κ B kinase, and MAPKK6 inhibitors in our *Myd88*-driven models of ABC-DLBCL. Altogether, we demonstrated the oncogenic role of a novel *Myd88*^{p.L252P} allele in B-cell neoplastic disease in vivo. The resulting autochthonous mouse models of ABC-DLBCL represent useful preclinical tools for the development, validation, and refinement of novel treatment strategies for ABC-DLBCL in the clinical arena.

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Authorship

Contribution: H.C.R. conceived and supervised the experiments and wrote the manuscript; L.P.F., H.K., M.A.-M., J.M.S., R.B., and F.T.W. provided mouse strains and materials; T.P. and Y.A.-B. performed magnetic resonance imaging scans and volumetric analyses; G.K., S.C.S., K.V., M.P., S.B., P.L., P.M., M. Ortmann, and R.B. performed immunohistochemistry and pathologic analyses; and L.M.S., A.N.R.W., O.-O.W., A.H.S., A.R.K., G.K., P.L., D.K., M. Odenthal, C.F., C.A.S., M.M.-R., M.R., M.A.-M., M.S., C.L., L.T., and R.S. performed experiments.

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