Regulation of B cell homeostasis and activation by the tumor suppressor gene CYLD

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The tumor suppressor gene CYLD was identified in patients with familial cylindromatosis, which is an autosomal dominant predisposition to multiple neoplasms of the skin appendages. The tumors are believed to arise from the eccrine or apocrine cells of the skin (1). Approximately 70% of the cylindromas exhibit loss of heterozygosity on chromosome 16q containing the CYLD gene (2–6).

Sequence analysis of the CYLD gene predicts a protein with several functional domains, including three CAP-Gly domains and a deubiquitination domain at the C terminus (7–9). In vitro studies showed that CYLD contains binding sites for TRAF2 and NEMO (7), and that the deubiquitinating activity of CYLD is directed to lysine 63 (K63)–linked ubiquitin (Ub) (7). Linkage of Ub through K63 assembles a new molecular platform, allowing the recruitment of proteins involved in signal transduction (10–12), whereas Ub chains linked through K48 destined proteins for proteasomal degradation (13, 14). Suppression of CYLD expression by RNA interference causes an increase in the activity of the transcription factor NF-κB, which is activated by TNFα signaling, thus inhibiting apoptosis (7, 8, 15). The increase of NF-κB activation in CYLD knockdown cells was attributed to defective removal of K63 Ub chains.
In a mechanistically analogous manner to its effect on TRAF2, CYLD was found to inhibit signaling from Toll-like receptor 2 (TLR2) by removing K63-linked Ub from TRAF6 and TRAF7 (16). Interestingly, it was shown that after TLR2 activation, CYLD inhibits the phosphorylation of TRAF6 and TRAF7, which, in turn, suppresses MKK3 and MKK6, leading to reduced MAP kinase p38 phosphorylation (17). In addition to p38 signaling, CYLD negatively regulates the stress-activating MAP kinase family, the JNK kinases. CYLD inactivation resulted in hyperactivation of JNK upon TNFα, IL-1β, LPS, and anti-CD40 treatment (17). This hyperactivation is believed to be a result of negatively regulating the activation of MKK7, which is the kinase responsible for JNK activation (17).

To investigate the physiological role of CYLD in vivo, we generated mouse strains with targeted modifications in the CYLD gene. In addition to a complete knockout of CYLD, exon 7 of the CYLD gene was flanked by loxP sites in a second mouse strain to result in conditional, excessive, and solitary expression of a shorter splice variant of the CYLD protein, hereafter termed sCYLD. The sCYLD protein is devoid of both TRAF2 and NEMO binding sites. In this article, we present data describing the profound alterations in the immune system of these mice (CYLD exc7/8), which is characterized by lymphomegaly and splenomegaly and a striking increase in B cell numbers. Overexpression of sCYLD further bestows B lymphocytes with increased IgG1 production and enhances survival capabilities of the B cell.

**RESULTS**

**Generation of CYLD exc7/8 mutant mice**

CYLD mutant mice were generated applying standard gene-targeting techniques in mouse embryonic stem cells (Fig. S1, available at http://www.jem.org/cgi/content/full/jem.20070318/DC1). Using Cre/loxP technology, three different mouse strains were generated. The first strain, CYLD neo, contains the neo resistance gene upstream of exon 7, decreasing transcription of CYLD (Fig. S1). The second strain, CYLD FL, harbors two loxP sites flanking exon 7 of CYLD (Fig. S1). Finally, CYLD exc7/8 mice lacking exon 7 after Cre-mediated from TRAF2 and NF-κB essential modulator (NEMO) by CYLD (7–9).

**Figure 1.** The identification of a new splice variant of CYLD. (A) Schematic representation of exons 6–9 of the CYLD gene, with respective transcripts. (B) RT-PCR of cDNA generated from MEFs of the indicated genotypes using primer pair P1 and P2. WT full-length CYLD transcript corresponds to the 770-bp band, whereas the alternatively spliced CYLD transcript is 244 bp. Bands were isolated, subcloned, and verified by sequencing. The intermediate band could be determined as an unspecific band. (C) Western blot analysis of lysates from WT tissue and cells. (lane 1) Spleen; (lane 2) T cells; (lane 3) LNs; (lane 4) thymus; (lane 5) B cells. Actin was used as loading control. (D) Western blot analysis of lysates from WT and CYLD exc7/8 B cells using anti-CYLD antibody. Actin was used as loading control. (E) Schematic representation of the different CYLD constructs used for transfection experiments in HeLa cells. Highlighted are the important domains of CYLD. FL-CYLD (full-length CYLD), C/S-mut CYLD (full-length CYLD carrying a mutation from C to S to generate a catalytically inactive form), and sCYLD alternatively spliced CYLD lacking exons 7 and 8. (F) HeLa cells were transfected with plasmids encoding either His-FL-CYLD or His-sCYLD together with either plasmid pcDNA or plasmid encoding for TRAF2. Transfected cells were lysed and communoprecipitated with anti-His antibody for CYLD, separated on an 8% SDS-PAGE, and transferred to PVDF membrane. The membranes were incubated with TRAF2 and CYLD antibodies, respectively. (G) MEFs were transfected with TRAF2-FLAG cDNA and TRAF2 was immunoprecipitated with anti-FLAG antibody. Thereafter, TRAF2 was separated on a 4–12% SDS-PAGE and transferred to PVDF membrane. The membrane was incubated with Ub-specific antibody. (lane 1) Mock-transfected WT MEFs; (lane 2) WT MEFs transfected with TRAF2-FLAG; (lane 3) CYLD exc7/8 MEFs transfected with TRAF2-FLAG. Data presented are representative of three different experiments. (H) HeLa cells were transfected with the indicated CYLD construct encoding plasmids together with plasmids coding for Flag-Bcl-3 and His-Ubiquitin. Protein lysates of transfected cells were used for immunoprecipitations with anti-His antibody, and Western blots were incubated with antibodies against the His and the Flag tags. Flag-Bcl-3 was used as a loading control for the Bcl-3 levels in the different cell types.
recombination through transient transfection of the targeted embryonic stem cells with a Cre-expressing plasmid (Fig. S1). CYLD<sup>ex7/8</sup> mice were born at the expected Mendelian frequencies and survived normally when housed under pathogen-free conditions.

Germline deletion of exon 7 should lead to splicing from exon 6 to 8 resulting in an out-of-frame translation of CYLD (Fig. 1 A). RT-PCR was applied to cDNA from mouse embryonic fibroblasts (MEFs) of the indicated genotypes using primers located in exon 6 and 9 of the CYLD transcript to verify the absence of the WT allele in CYLD<sup>ex7/8</sup> MEFs. This analysis revealed an unexpected shorter amplified product besides the expected band from the full-length transcript (FL-CYLD) in WT MEFs. This shorter product represents an alternative splice variant of CYLD lacking exons 7 and 8, termed sCYLD. CYLD<sup>ex7/8</sup> MEFs are devoid of the full-length transcript, but express the sCYLD splice variant excessively (Fig. 1 B). Using CYLD-specific antibodies, sCYLD protein could be detected in different tissues and cells of WT animals (Fig. 1 C). To investigate whether deletion of WT CYLD (FL-CYLD) in CYLD<sup>ex7/8</sup> mice results in higher expression levels of sCYLD, protein extracts were prepared from purified B cells of WT and CYLD<sup>ex7/8</sup> mice, revealing high expression of sCYLD in CYLD<sup>ex7/8</sup> cells compared with WT cells (Fig. 1 D).

Previously, CYLD was identified as a deubiquitinating enzyme, removing K63-conjugated Ub molecules from TRAF2, TRAF6, and NEMO (7–9). However, the putative TRAF2 and NEMO binding sites are absent in the sCYLD protein (Fig. 1 E). To verify whether sCYLD is, indeed, unable to bind TRAF2, HeLa cells were transfected with expression plasmids encoding either FL-CYLD or sCYLD together with TRAF2-encoding or control plasmids. As shown in Fig. 1 F, FL-CYLD protein, but not sCYLD, coimmunoprecipitated with TRAF2, indicating that in contrast to FL-CYLD, sCYLD does not bind TRAF2. However, in an 8% SDS-PAGE, the difference in size between FL-CYLD and sCYLD could not be unambiguously demonstrated (Fig. 1 F). In agreement, immunoprecipitated TRAF2 from extracts of CYLD<sup>ex7/8</sup> MEFs was highly ubiquitinated compared with TRAF2 immunoprecipitated from WT cells (Fig. 1 G).

To investigate whether sCYLD is still able to function as a deubiquitinating enzyme, similar transfection experiments were applied to determine whether both CYLD proteins are capable of removing K63-conjugated Ub from Bcl-3, which is a novel substrate for CYLD (18). The Bcl-3 binding site in CYLD is predicted to reside outside the area absent in sCYLD (Fig. 1 E). We cotransfected HeLa cells with expression vectors encoding His-tagged Ub and Flag-tagged Bcl-3, together with plasmids coding either for FL-CYLD, or for a catalytically inactive form of CYLD (C/S-CYLD) or for sCYLD. Fig. 1 H shows that both sCYLD and FL-CYLD are able to remove Ub from Bcl-3. Collectively, our results demonstrate that sCYLD is not able to deubiquitinate TRAF2, but still removes K63-linked Ub from Bcl-3.

Lymphoid system defects in CYLD<sup>ex7/8</sup> mice

CYLD<sup>ex7/8</sup> mice exhibited considerably larger spleens, LNs, and Peyer’s patches (Fig. 2 A and not depicted). Careful microscopic examination revealed a disorganized structure of the spleen, where the distinct B cell and T cell zones of the follicle were not uniform as in WT spleens (Fig. 2 B). Importantly, such abnormalities of the secondary immune organs were not reported in mice with complete inactivation of the CYLD gene (18) (unpublished data).

The CYLD<sup>ex7/8</sup> mice displayed massively enlarged secondary lymphoid organs as early as 4 wk of age, whereas other tissues appeared normal (unpublished data). To investigate the mechanisms underlying these enlarged lymphoid organs, we examined lymphocyte populations of BM, spleen, LNs, and peritoneal cavity (PC). Analysis of B220<sup>+</sup> cells in the BM of CYLD<sup>ex7/8</sup> mutant mice shows a small decrease in the percentage of the immature B lymphocyte compartments (Fig. 3 A, top). To analyze early B cell development, B220<sup>+</sup>IgM<sup>−</sup> cells were analyzed for the expression of the pro-B cell marker, c-Kit. We observed an increase in the pro-B cell fraction, indicating that deletion of the FL-CYLD leads to a partial block in early B cell differentiation (Fig. 3 A, bottom).

Strikingly, CYLD<sup>ex7/8</sup> mice, but not heterozygous and control mice, demonstrate a dramatic accumulation of B cells in all secondary lymphoid organs tested (Fig. 3 B and Table I; not depicted for Peyer’s patches). The proportion and absolute number of immature B cells was reduced in CYLD<sup>ex7/8</sup> mice (Fig. 3 C and Table I), although they displayed significantly higher total B cell numbers (Table I).

As seen in Fig. 3 D and Table I, the proportion, as well as the absolute number, of marginal zone (MZ; CD21<sup>high</sup>CD23<sup>low</sup>)}
and follicular (CD21^hiCD23^lo) B cells were increased by two- to threefold in CYLD^ex7/8^ mice compared with WT mice. Interestingly, expression of CD21 and CD23 by B cells isolated from CYLD^ex7/8^ mice was elevated compared with control B cells (Fig. 3 D). The latter was accompanied by an increase in cell size of the CYLD mutant B cells (unpublished data). To analyze the structure of MZ B cells, spleen sections from CYLD^ex7/8^ and littermate control mice were stained with anti-MOMA1 antibody, which binds to metallophilic macrophages, marking the border of the MZ. As shown in Fig. 3 E, MOMA1^+^ cells can be detected in the spleens of CYLD^ex7/8^; however, the MZ appeared to be disordered and discontinuous compared with WT spleens. Furthermore, the amount of MOMA1^+^ cells was increased without forming a closed ring around the follicle. Notably, the B cell region located outside of the MOMA1^+^ cells corresponding to the MZ B cells was larger in the mutant spleens compared with controls (Fig. 3 E). Interestingly, the expansion of MZ and follicular B cells was accompanied by a decrease in the proportion and absolute number of B-1a (CD19^hiCD5^-CD23^-) and B-1b (CD19^hiCD5^-CD23^-) B cells in the PC (Fig. 3 F and Table I). These results suggest that either overexpression of sCYLD or disruption of FL-CYLD is critical for regulating the maturation of BM B cells and the distribution of peripheral B cells.

To address whether these differences in the B cell compartment resulted from the overexpression of sCYLD, we analyzed another CYLD mutant strain, CYLD^ex8^, in which the neomycin resistance gene was introduced into the first coding exon of the CYLD gene, thus preventing its expression (18).
Fig. S2 A (available at http://www.jem.org/cgi/content/full/jem.20070318/DC1) shows that the B cell development in the BM was unchanged in CYLDko mice. Furthermore, in contrast to CYLDex7/8 mice, CYLDko mice showed neither differences in B/T cell ratio nor an elevation of MZ and follicular B cells in the spleen nor a decrease of B-1 B cells in the PC (Fig. S2, B–D). Moreover, B cell numbers in these mice were comparable to WT B cell numbers (Table I). Thus, we conclude that the absence of the FL-CYLD, in tandem with the overexpression of sCYLD, is responsible for the increased proportion of B cells in the lymphoid organs of CYLDex7/8 mice.

Skewed humoral immune response of CYLDko7/8 mice

To determine whether the expansion of B cells in CYLDex7/8 mice affects the immune response, we evaluated serum antibody levels in CYLDex7/8 and WT mice. The sera of CYLDex7/8 mice contained significantly lower IgM and IgG3 antibody titers compared with WT littermates, whereas all other tested antibody classes, including IgG1, IgG2a, IgG2b, and IgA, were elevated in CYLDex7/8 mice (Fig. 4 A). The latter antibody isotypes are more frequently secreted by class-switched B cells after encountering T cell–dependent (TD) antigens. Therefore, we next investigated TD antigen responses of CYLDex7/8 B cells and control B cells. To this end, mice were challenged with nitrophenol-conjugated chicken y-globulin (NP-CG) and immune responses were measured by ELISA. Upon in vivo challenge with NP-CG, CYLDex7/8 mice developed significantly lower antibody titers of both IgM and IgG3 isotypes, whereas secretion of NP-specific IgG1 and IgG2a antibodies was enhanced compared with WT mice (Fig. 4 B). In addition, CYLDex7/8 mice developed higher titers of NP-specific IgA antibodies, whereas the NP-specific IgE antibody levels were similar to the levels in WT mice (Fig. 4 B, top). Previously, it was shown that WT mice harbor a relatively high number of NP-specific precursor cells (~1%) (19). To test whether these findings hold true for another TD antigen with lower precursor frequency, we immunized mice with KLH. CYLDex7/8 mice developed significantly higher levels of KLH-specific IgG1 antibodies after immunization compared with WT mice (Fig. S3 A, available at http://www.jem.org/cgi/content/full/jem.20070318/DC1). CYLDex7/8 mice and littermate controls were also immunized with the T cell–independent (TI) antigen NP-Ficoll, resulting in similar levels of NP-specific IgM and IgG3 antibodies in CYLDex7/8 and WT mice (Fig. 4 C).

To examine whether the high IgG1 antibody levels observed in nonimmunized mice, as well as in the immunized mutant mice, were a consequence of increased antibody production by plasma cells or whether CYLDex7/8 B cells switch more efficiently to other antibody classes, B cells from WT and CYLDex7/8 mice were isolated and, subsequently, incubated for 5 d ex vivo with optimal concentrations of LPS and IL-4. A higher proportion of CYLD mutant B cells switched to the IgG1+ isotype compared with WT B cells (Fig. S3 B). B cells in the Peyer’s patches are constantly encountered by the gut flora, and therefore yield higher proportions of class-switched B cells. Peyer’s patches from CYLDex7/8 mice contain a threefold increased B cell population rich in IgG1-expressing B cells compared with WT B cells (Fig. 4 A). The B cell phenotype in CYLDex7/8 mice is B cell intrinsic

As all cells in CYLDex7/8 mice lacked exon 7, it was unclear whether the defect in B cells was intrinsic to B cells or secondary

### Table I. Lymphocyte population sizes

| Spleen          | Genotype                  | Total  | B cells | Immature | MZ |
|-----------------|---------------------------|--------|---------|----------|----|
| WT (14)         |                           | 79.4 (8.2) | 46.7 (4.8) | 7.7 (0.8) | 2.4 (0.3) |
| CYLDex7/8 (9)   |                           | 96.9 (13.0) | 49.3 (6.6) | 6.2 (0.9) | 2.9 (0.4) |
| CYLDex7/8 (14) |                           | 138.7 (21.6) | 111.1 (17.3) | 4.8 (0.7) | 10.3 (1.6) |
| CYLDex7/8 (5)   |                           | 84.7 (8.4) | 42.4 (4.1) | 14.4 (1.4) | 4.6 (0.4) |
| CD19-Cre/CYLDex7/8 (5) |            | 134.5 (12.6) | 100.8 (9.5) | 11.1 (1.0) | 10.1 (0.9) |

| Peritoneal cavity | Genotype                  | Total  | B cells | Immature | MZ |
|-------------------|---------------------------|--------|---------|----------|----|
| WT (4)            |                           | 1.4 (0.3) | 0.9 (0.2) |         |     |
| CYLDex7/8 (4)     |                           | 1.1 (0.1) | 1.6 (0.1) |         |     |
| CYLDex7/8 (4)     |                           | 0.3 (0.2) | 2.1 (0.2) |         |     |
| CYLDex7/8 (5)     |                           | 1.5 (0.3) | 0.3 (0.1) |         |     |
| CD19-Cre/CYLDex7/8 (5) |             | 0.4 (0.1) | 0.8 (0.2) |         |     |

The mean size of lymphocyte populations was calculated based on their frequency as determined by flow cytometry (Fig. 2). Total numbers (× 10⁶) are indicated (the SEM is shown in parentheses). Data are the mean of the number of mice indicated in parentheses after the mice genotypes. Values are the mean ± the SEM.

P ≤ 0.05.

P ≤ 0.001 versus control.
because of defects in other cell types. Therefore, we crossed CYLD<sup>FL/FL</sup> mice to CD19-Cre mice (20, 21), leading to Cre-mediated excision of exon 7 of CYLD solely in B cells. The resulting mice, CD19-Cre/ CYLD<sup>FL/FL</sup> exhibited enlarged secondary immune organs, including spleen, LNs, and Peyer’s patches similar to the CYLD<sup>ex7/8</sup> mice that harbor the mutation in the germline (unpublished data). To analyze B cell development in these mice, BM cells were stained for B220 and IgM. As seen in Fig. 5 A (top), BM cells of CD19-Cre/CYLD<sup>FL/FL</sup> mice show a small decrease in the percentage of immature cells, similar to BM of CYLD<sup>ex7/8</sup> mice. B220<sup>+</sup>IgM<sup>−</sup> cells were gated and analyzed for the expression of the pro-B cell marker, c-Kit. As for the CYLD<sup>ex7/8</sup> mice, an accumulation of c-Kit−positive cells in the BM of CD19-Cre/CYLD<sup>FL/FL</sup> was detected (Fig. 5 A, bottom). Importantly, also in CD19-Cre/CYLD<sup>FL/FL</sup> mice, the total number of B cells was increased by two- to threefold (Fig. 5 B and Table I) compared with WT mice. All other characteristics of the B lymphocytes detected in CYLD<sup>ex7/8</sup> mice were also manifested in CD19-Cre/CYLD<sup>FL/FL</sup> mice, including a reduction in the percentage of transitional B cells, a reduction of immature B cells (Fig. 5 C), and an increase in the total number of MZ B cells (Fig. 5 D and Table I). Finally, in both CD19-Cre/CYLD<sup>FL/FL</sup> and CYLD<sup>ex7/8</sup> mice, the ratio of B-1 to B-2 B cells in the PC was inverted (Fig. 5 E). Together, these data indicate that the B cell defects seen in the CYLD<sup>ex7/8</sup> mice are cell autonomous.

Overexpression of sCYLD results in prolonged B cell survival
Bcl-2 transgenic mice exhibit enhanced B cell survival in vitro; this is associated with an increased size of secondary immune organs and B cell compartments in vivo (22), which is similar to CYLD<sup>ex7/8</sup> mice. To examine the survival capacities, B cells from CYLD<sup>ex7/8</sup> and WT mice were cultured in enriched medium and monitored daily for survival by cell counting and FACS analysis. The CYLD mutant B cells exhibited prolonged survival compared with WT B cells (Fig. 6 A, top).
assessed by quantitative real-time PCR. CYLD ex7/8 B cells express fivefold more Bcl-2 mRNA compared with WT B cells (Fig. 6D). These results were further verified at protein level by intracellular staining with Bcl-2–specific antibody (Fig. 6D). In addition, we found that primary fibroblasts isolated from CYLD ex7/8 mutant mice are more resistant to TNFα/H9251-induced apoptosis in the absence of protein synthesis (Fig. 6E). Therefore, it is tempting to speculate that the deletion of the FL-CYLD, concomitant with the overexpression of sCYLD, conveys increased resistance to apoptosis, leading to the enlarged B cell compartment in CYLD ex7/8 mice.

Next, we determined the responses of CYLD ex7/8 and WT B cells to activating stimuli by incubating CFSE-labeled B cell cultures with optimal concentrations of anti-IgM F(ab)2 and anti-CD40 antibodies, or with anti-RP105, LPS, or BAFF. As shown in Fig. 6F, B cells from CYLD ex7/8 mice responded similarly to the different stimuli compared with WT B cells. Although the addition of BAFF to culture medium resulted in a more activated status of the mutant B cells compared with the WT B cells (Fig. 6D), supplementing the culture with BAFF together with LPS failed to increase proliferation (Fig. 6F). Therefore, we conclude that sCYLD contributes to the survival capacity of B cells, but does not affect B cell proliferation.

Elevated expression of NF-κB and TRAF proteins in resting CYLD ex7/8 B cells

Previously, CYLD was shown to act on different proteins involved in the NF-κB (7–9) and mitogen-activated protein kinase (MAPK) signaling pathways (17, 24). To study whether the enlargement of the B cell compartment in CYLD ex7/8 mice was assessed by quantitative real-time PCR. CYLD ex7/8 B cells express fivefold more Bcl-2 mRNA compared with WT B cells (Fig. 6D). These results were further verified at protein level by intracellular staining with Bcl-2–specific antibody (Fig. 6D). In addition, we found that primary fibroblasts isolated from CYLD ex7/8 mutant mice are more resistant to TNFα-induced apoptosis in the absence of protein synthesis (Fig. 6E). Therefore, it is tempting to speculate that the deletion of the FL-CYLD, concomitant with the overexpression of sCYLD, conveys increased resistance to apoptosis, leading to the enlarged B cell compartment in CYLD ex7/8 mice. After 4 d of culture, the number of living CYLD ex7/8 B cells was increased by twofold compared with WT B cells. Furthermore, the survival properties of CYLD ex7/8 B cells were compared with those of WT B cells cultured in medium supplemented with the B cell survival factor BAFF. BAFF treatment led to improved survival of both WT and mutant B cells (Fig. 6A, bottom). However, at all time points, CYLD ex7/8 B cells responded better to BAFF treatment compared with control B cells. Additionally, BAFF triggering of CYLD ex7/8 B cells resulted in the up-regulation of the activation markers CD40 and CD86, respectively, in comparison to WT B cells, whereas CpG treatment had similar effects on both B cell cultures (Fig. 6B). To investigate whether the increased numbers of mature B cells in CYLD ex7/8 mice was related to response to BAFF in vivo, CYLD mutant and control mice were treated with either TACI- or human-Ig as control. It was shown that TACI-Ig blocks BAFF signaling, and that this treatment resulted in a dramatic decrease in B cell numbers (23). In treated WT mice, we detected a significant decrease in the percentage of mature B cells in the mesenteric LN (Fig. 6C and Fig. S4A, available at http://www.jem.org/cgi/content/full/jem.20070318/DC1) and spleen (Fig. S4, B and C). In contrast, the percentage of B cells in CYLD ex7/8 mice was hardly affected by the TACI-Ig treatment (Fig. 6C and Fig. S4). No differences in the total numbers of T cells in CYLD ex7/8 and WT mice were observed upon TACI treatment, showing that, indeed, TACI-Ig acts specifically on B lymphocytes (unpublished data).

To investigate the increased viability of CYLD ex7/8 B cells in vitro, the expression of the antiapoptotic gene Bcl-2 was assessed by quantitative real-time PCR. CYLD ex7/8 B cells express fivefold more Bcl-2 mRNA compared with WT B cells (Fig. 6D). These results were further verified at protein level by intracellular staining with Bcl-2–specific antibody (Fig. 6D). In addition, we found that primary fibroblasts isolated from CYLD ex7/8 mutant mice are more resistant to TNFα-induced apoptosis in the absence of protein synthesis (Fig. 6E). Therefore, it is tempting to speculate that the deletion of the FL-CYLD, concomitant with the overexpression of sCYLD, conveys increased resistance to apoptosis, leading to the enlarged B cell compartment in CYLD ex7/8 mice.
mice results from alterations in the expression levels of proteins involved in the NF-κB signaling pathway, whole-cell lysates of purified B cells from three CYLDex7/8 mutant mice and three control mice were subjected to Western blot analysis. This analysis revealed increased protein levels of TRAF2 in the CYLD mutant B cells compared with WT B cells (Fig. 7 A). Additionally, the protein levels of TRAF3 and NEMO were determined, showing that TRAF3 levels were increased, whereas NEMO levels were unchanged (Fig. 7 A). The levels of 1κBα protein were markedly elevated in CYLDex7/8 B cells, although its phosphorylation on conserved serine residues was unaltered. Most interestingly, NfκB2/p100 protein levels primarily involved in the noncanonical NF-κB pathway were dramatically increased, whereas the levels of its processed form, p52, were only slightly increased (Fig. 7 A).

Furthermore, we could show that the RelB protein levels were also highly increased in the cytoplasm and nucleus of CYLDex7/8 B cells (Fig. 7 B). The elevated protein levels of TRAF2, TRAF3, p100, RelB, and 1κBα in CYLDex7/8 B cells likely resulted from increased protein stability rather than from increased transcriptional activity, as quantitative real-time PCR revealed no significant differences in relative expression of these genes (Fig. S5, available at http://www.jem.org/cgi/content/full/jem.20070318/DC1).

To investigate whether the noncanonical NF-κB pathway is affected in CYLDex7/8 B cells, purified B lymphocytes of control and CYLDex7/8 mice were cultivated with optimal concentrations of BAFF for the indicated time points (Fig. 7 C).
Nonstimulated CYLD<sup>ex7/8</sup> B cells exhibited detectable NF-κB activation, which was not observed in WT B cells (Fig. 7 E). However, the NF-κB–activating response to anti-IgM was unaltered at the different time points of stimulation in CYLD<sup>ex7/8</sup> B cells compared with WT B cells. This result suggests that the basal activity of NF-κB may reflect an increased response to a yet unknown endogenous activator.

To measure canonical NF-κB activity in CYLD<sup>ex7/8</sup> B cells, splenic B cells were stimulated with anti-CD40, LPS, and anti-BCR (Fig. 7 D). We did not observe significant differences in IkBα degradation upon triggering of CYLD<sup>ex7/8</sup> and WT B cells with the aforementioned stimuli (Fig. 7 D). Similarly, we did not detect changes in IkBα degradation and phosphorylation after anti-BCR treatment (Fig. 7 D). As a direct measure of NF-κB activity, we performed electrophoretic mobility shift assay (EMSA) using NF-κB consensus-binding oligonucleotides with nuclear extracts from resting B cells stimulated with optimal concentrations of anti-BCR (Fig. 7 E). Nonstimulated CYLD<sup>ex7/8</sup> B cells exhibited detectable NF-κB activation, which was not observed in WT B cells (Fig. 7 E). However, the NF-κB–activating response to anti-IgM was unaltered at the different time points of stimulation in CYLD<sup>ex7/8</sup> B cells compared with WT B cells. This result suggests that the basal activity of NF-κB may reflect an increased response to a yet unknown endogenous activator.

To estimate the effect of the sCYLD in B cells on MAPK signaling, we performed cytometric bead assay (CBA) to determine quantitatively phosphorylated MAPKs, which can be visualized using FACS analysis. In this assay, LPS stimulation resulted in increased phosphorylation of MAPK p38 in CYLD<sup>ex7/8</sup> B cells compared with WT B cells (Fig. 7 F). Similar results were observed by Western blotting of anti-BCR–stimulated B cells (Fig. 7 G). In contrast, we observed
reduced phosphorylation of ERK1 and ERK2 upon LPS stimulation (Fig. 7 F). This could be further confirmed with Western blot analysis (Fig. 7 G).

In keratinocytes isolated from CYLD<sup>ko</sup> mice, Bcl-3 associates with the NF-κB proteins p52 and p50 and promotes increased transactivation of cyclin D1 compared with WT cells (18). Although we have shown that sCYLD is able to interact with Bcl-3 and to remove K63-conjugated Ub, we were interested to see whether Bcl-3 is present in the nuclei of B cells from CYLD<sup>ex7/8</sup> mice without activation. To this end, we prepared cytospins of CYLD<sup>ex7/8</sup>, CYLD<sup>ex7/8;wt</sup>, CYLD<sup>ko</sup>, and WT B cells. Fig. 8 A and B demonstrates that Bcl-3 was predominantly located in the nucleus, or in close proximity to the nuclei, of untreated CYLD<sup>ex7/8</sup> B cells, whereas in WT B cells, Bcl-3 was primarily found in the cytoplasm. Calculating the percentage of B cells of the various genotypes containing nuclear Bcl-3 revealed that 60% of the CYLD<sup>ex7/8</sup> B cells contain nuclear Bcl-3, whereas B cells from WT, CYLD<sup>ex7/8;wt</sup>, and CYLD<sup>ko</sup> exhibit 8, 3, and 2.5% of Bcl-3 in the nucleus, respectively (Fig. 8 C). Furthermore, we could corroborate these results using Western blot analysis of extracts from CYLD<sup>ex7/8</sup> mice (Fig. 8 D). By quantitative real-time PCR, we could show that the increased protein levels of Bcl-3 in the B cells of the CYLD<sup>ex7/8</sup> mice are not a result of higher Bcl-3 gene expression (Fig. S5). Interestingly, mice that are deficient for CYLD do not show spontaneous translocation of Bcl-3 to the nucleus. These results are reminiscent of the previously published Bcl-3 transgenic mice (25), in which overexpression of Bcl-3 in B cells resulted in lymphadenopathy, splenomegaly, and altered immunoglobulin production, similar to what was observed in CYLD<sup>ex7/8</sup> mice.

**DISCUSSION**

In this article, we report the function of a splice variant of CYLD devoid of the TRAF2 and NEMO binding sites. Expression of this shorter form of CYLD (sCYLD) with a simultaneous deficiency of the full-length (FL) protein gave rise to a dramatic accumulation of mature B lymphocytes in all secondary lymphoid organs in vivo. This B cell accumulation is a consequence of increased survival, rather than increased proliferation, putatively resulting from aggravated Bcl-2 expression caused by various alterations in signal transduction pathways, such as NF-κB and MAPK signaling. In contrast, mice that are entirely devoid of CYLD (CYLD<sup>ko</sup> mice) do not show any abnormalities in the B cell compartment. Therefore, we reason that sCYLD has other functions in B cell homeostasis than FL-CYLD. Under physiological conditions FL-CYLD expression might be diminished, whereas the expression of sCYLD is elevated. Indeed, we detected various expression levels of both CYLD forms in different tissues and cell types. Therefore, we suggest that also

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**Figure 8.** Bcl-3 accumulation in CYLD<sup>ex7/8</sup> B cells. (A) Cytospins were prepared from B cells isolated from CYLD<sup>ex7/8</sup>, CYLD<sup>ex7/8;wt</sup>, CYLD<sup>ko</sup>, and WT mice and immunostained with anti-Bcl-3 and counterstained with Hoechst 33258. Shown are unstimulated B cells (top) and B cells treated with anti-BCR (bottom). Bar, 5 μm. (B) Same as in A, but showing more cells. Bar, 5 μm. (C) Quantification of nuclear Bcl-3 from unstimulated B cell cytospins from the indicated genotypes. For each individual genotype, 400 cells were counted and quantified for nuclear Bcl-3 localization. The frequency of nuclear Bcl-3 localization in CYLD<sup>ex7/8</sup> B cells were statistically significant (P < 0.001) compared with B cells isolated from WT, CYLD<sup>ex7/8;wt</sup>, and CYLD<sup>ko</sup> mice. (D) Lysates of MACS-purified B cells from the indicated genotypes subjected to Western blot analysis using Bcl-3 antibody. Duplicates for each mouse strain are shown. Actin was used as a loading control.
under normal conditions, the unique function of sCYLD can be manifested.

The CYLD-deficient mice generated by Reiley et al. (26) demonstrate an elevated number of B lymphocyte, which is explained by a compensatory effect caused by the reduction in the T-lymphocyte number (26). In CYLD<sup>−/−</sup> mutant mice, the generalized accumulation of mature B cells seems to be B cell intrinsic because CD19-Cre/CYLD<sup>+/+L/FL</sup> mice, in which only the B cells express sCYLD and concomitantly lack the FL-CYLD protein, show a dramatically enlarged B cell compartment. We further observed that CYLD<sup>−/−</sup> mice have an increased number of Igκ-positive B cells compared with WT mice. This might be caused by the partial block in B cell development, which is demonstrated by an increase in the number of c-Kit–positive B cells in the BM. A block during B cell development may allow the developing B cells to edit their light chain locus, as seen in the process of B cell editing (27), which, indeed, results in higher numbers of λ-positive B cells.

We observed increased protein levels of several molecules involved in the NF-κB signaling pathway in CYLD<sup>−/−</sup> B cells, including IkBα. However, the degradation and phosphorylation of IkBα is as efficient in the CYLD<sup>−/−</sup> B cells as in the WT B cells upon stimulation, showing that the canonical NF-κB activation is unaltered. Consistently, the relative expression of IkBα, p100, and RelB, which are all under the regulation of the NF-κB pathway, was unaltered as determined by quantitative real-time PCR, thus pointing to increased protein stability resulting from differences in post-translational modifications. Nevertheless, EMSA revealed elevated binding of nuclear transcription factors to the NF-κB consensus probe in immunostimulated B cells.

We could demonstrate elevated protein levels of TRAF2 and TRAF3 in CYLD<sup>−/−</sup> B cells, which is a consequence of impaired posttranslational modification because relative expression revealed by real-time PCR was unaltered. Consistently, TRAF2 was highly ubiquitinated in CYLD<sup>−/−</sup> B cells, presumably by K63-linked Ub chains, as shown by Western blot analysis. Therefore, it is tempting to speculate that K63-ubiquitinated TRAF2 exhibit increased protein stability and possibly have impaired functionality. This is supported by the aggravated TRAF3 protein levels in CYLD<sup>−/−</sup> B cells because TRAF2 was shown to be a negative regulator of TRAF3 upon stimulation with CD40 (28). B lymphocytes deficient for TRAF2 demonstrate a phenotype similar to that of CYLD<sup>−/−</sup> B cells with increased TRAF3 protein levels, but also show constitutive processing of p100 to p52, which is contrary to CYLD<sup>−/−</sup> B cells (29). In contrast, however, TRAF3-deficient cells also display constitutive processing of p100 to p52, resulting in embryonic lethality. This phenotype can be rescued in vivo by simultaneous deficiency of p100 (30). Therefore, elevated TRAF3 levels should lead to increased p100 levels. Consistently, CYLD<sup>−/−</sup> B cells exhibit increased p100 protein levels, but not elevated processing to p52. In addition, elevated TRAF2 and TRAF3 levels might promote MAPK p38-mediated isotype switching to IgG1, as essential functions for these adaptor molecules in CD40-induced class switch recombination and Ig production were reported (31, 32). Indeed, CYLD<sup>−/−</sup> B cells switch more efficiently to IgG1, but whether this is a result of the p38 hyperactivation observed in CYLD<sup>−/−</sup> B cells or a result of impaired ERK activation still needs further investigation.

Furthermore, CYLD<sup>−/−</sup> B cells exhibit a significant accumulation of Bcl-3 in the nucleus that cannot be explained by an inability of sCYLD to associate with Bcl-3, as we could show that sCYLD binds and removes K63-linked ubiquitin chains from Bcl-3. Western blot analysis of B lymphocytes using Bcl-3 antibody revealed aggregated Bcl-3 protein levels, whereas on the RNA level, expression of Bcl-3 was unaltered in CYLD<sup>−/−</sup> B cells. Ultimately, CYLD<sup>−/−</sup> mice show a very similar phenotype to mice overexpressing Bcl-3 in B lymphocytes (25). Both transgenic mice show lymphadenopathy and splenomegaly and exhibit distorted follicles in the spleen, resulting from an enlarged B cell compartment. Further, both mouse strains showed an increase in IgG1 and decreased levels of IgM and IgG3 in the sera compared with WT mice. In addition, enhanced B cell survival, accumulation of MZ B cells, as well as increased B cell size, and the increased expression of CD21/CD23 observed in CYLD<sup>−/−</sup> mice are also found in Bcl-3 transgenic mice.

The elevated expression of Bcl-2 in CYLD<sup>−/−</sup> B cells may be a consequence of the dramatic overexpression of the NF-κB proteins p100 and RelB, which might activate transcription of Bcl-2 in a complex with Bcl-3 upon signaling through the noncanonical NF-κB pathway (33). Thus, nuclear Bcl-3 is in combination with p100 can induce the expression of Bcl-2 (33) and might account for the elevated B cell survival observed in the CYLD<sup>−/−</sup> mice, as shown previously for Bcl-2 transgenic mice (22, 34). Concomitantly, CYLD<sup>−/−</sup> B cells are less dependent on BAFF signaling by a yet unsolved mechanism. This was demonstrated by ineffective BAFF blocking by TACI-Ig. Collectively, our data suggest that the newly identified splice variant sCYLD regulates B cell survival mediated through a variety of signaling disarrangements.

**Materials and Methods**

**Generation of CYLD mutant mice.** The generation of CYLD mutant mice is described in the Supplemental materials and methods. All animal experiments were in accordance with the guidelines of the Central Animal Facility Institution of the University of Mainz.

**RNA analysis.** For RNA analysis, total RNA was isolated using Trizol reagent (Invitrogen) according to the manufacturer’s instruction. The following primers were used for RT-PCR amplification of mouse CYLD: P1, 5′-CTCAAGGGCTGGAATGAA-3′; P2, 5′-TGGTATGCCCCT-GGATGCC-3′. Quantitative real-time PCR of Bcl-2, p100, RelB, IkBα, TRAF2, TRAF3, and Bcl-3 was performed using primers from QIAGEN, as described on their homepage (https://www1.qiagen.com/GeneGlobe/Default.aspx).

**Immunostaining of purified B cells.** Cytospins of purified B cells were fixed in acetone for 10 min at −20°C and stained for Bcl-3 according to standard methods, using polyclonal rabbit IgG Bcl-3 antibody. All incubation steps of the staining procedure were performed at room temperature.
The cytoplasmic tails of HMGB-1 were counterstained with Hoechst 33258. Cytoplasmic tails were analyzed by confocal microscopy.

**Histological analysis.** For histological staining of B and T cells and germinal centers, frozen 6-μm spleen sections were thawed, air dried, fixed in acetone, and stained for 1 h at room temperature in a humidified chamber with biotinylated rat anti-CD19 (BD Biosciences), rat anti-B220, anti-CD3ε, and anti-MOMA1, respectively (eBioscience), followed by horseradish peroxidase–conjugated secondary antibody and alkaline phosphatase–conjugated streptavidin.

**Flow cytometry.** Single-cell suspensions were prepared from the different organs. Red blood cells of LN and spleen were lysed in cell suspension with tris-ammonium chloride, pH 7.2. Cells were incubated with combinations of antibodies to cell surface determinants, conjugated to PE, FITC, Cy-Cyochrome, or biotin. Antibodies specific to the following surface markers were purchased from BD Biosciences: CD3ε, CD19, CD21 (766), CD23, HSA (CD24), CD25, CD40,2, CD43, and IgD. Anti-CD45R (B220; clone RA3-6B2) and anti-IgM (R33-24-12) were prepared in our laboratory. Biotinylated cells were visualized with streptavidin conjugated to PE or Cy-Cyochrome (BD Biosciences). Bcl-2 was visualized using anti-Bcl-2 antibody (Santa Cruz Biotechnology) and secondary donkey anti-rabbit sera conjugated to FITC (Jackson ImmunoResearch Laboratories). All samples were acquired on a FACSCalibur (BD Biosciences), and results were analyzed with CellQuest software (BD Biosciences). Absolute numbers of splenocyte subpopulations were calculated based on their percentage and the total number of splenocytes.

**Immunostaining and transfection.** The coding regions of murine Bcl-3, TRAF2, FL-CYLD, or CYLD were cloned after RT-PCR using specific primers from the total RNA of activated B cells. PCR products were cloned in pcDNA3.1 vector (Invitrogen). HeLa cells and MEFs were transfected with the different constructs or histidine-ub (His-ub), as described in Fig. 1, using Lipofectamine2000 (Invitrogen). For anti-Bcl-3 or anti-TRAF2 immunoprecipitation, lysates were preclarified for 30 min at 4°C. The protein content was determined and compensated for equal content in all supernatants. For testing protein input, a small part of the resulting lysates was gel separated and immunoblotted (protein input), and the remaining part was used for immunoprecipitation. Immunoblots were developed with the ECL Plus reagent (GE Healthcare) according to the manufacturer’s guidelines.

**Immunization.** 8-wk-old WT and CYLD<sup>−/−</sup> mice were immunized intraperitoneally with 20 μg of NP<sub>28</sub>-CCG or 100 μg of KLH for TD response, or 20 μg of NP-Fcoll for TI responses. 3 days after the last injection of 50 μg of NCP28-CCG and 50 μg of KLH, the mice were sacrificed and the spleens were harvested. Spleens were cut into small pieces and placed into 50 μl of digestion buffer (1% Trypsin, protease inhibitors). The resulting Ficoll fractions were lysed in RIPA buffer and subjected to Western blotting.

**Cell preparation and B cell proliferation assay.** Total mouse splenic B cells were purified through negative depletion with CD43 beads (Miltenyi Biotech). The purity (>98%) was determined by flow cytometry. For in vitro proliferation studies, B cells were labeled with CFSE, and 2×10<sup>5</sup> cells/well were incubated in RPMI medium containing 10% FCS, 2×10<sup>5</sup> cells/well were incubated in RPMI medium plus 10% FCS or untreated or treated with 5 μg/ml anti-CD40, 5 μg/ml anti-IgM, 2 μg/ml LPS, and 5 μg/ml LPS plus IL-4 with and without 100 ng/ml BAFF for 4 d.

**Measurement of TNFα-induced cell death.** MEFs (10<sup>5</sup> per well) were plated onto 6-well plates and cultured for 24 h in DMEM containing 10% FCS. Subsequently, the cells were stimulated with either 10 ng/ml TNFα with and without 300 ng/ml cycloheximide. Triplicate samples of TNFα-induced cell death were measured by FACS analysis using Toto3 and Annexin V.
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