CTP:phosphocholine Cytidyltransferase \( \alpha \) Is Required for B-cell Proliferation and Class Switch Recombination*§

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CTP:phosphocholine cytidyltransferase (CCT) is a key rate-controlling enzyme in the biosynthetic pathway leading to the principle membrane phospholipid, phosphatidylcholine. CCT\( \alpha \) is the predominant isofom expressed in mammalian cells. To investigate the role of CCT\( \alpha \) in the development and function of B-lymphocytes, mice with B-lymphocytes that selectively lacked CCT\( \alpha \) were derived using the CD19-driven Cre/loxP system. When challenged with a T-cell-dependent antigen, the animals harboring CCT\( \alpha \)-deficient B-cells exhibited a hyper-IgM secretion phenotype coupled with a lack of IgG production. The inability of CCT\( \alpha \)/–/– B-cells to undergo class switch recombination correlated with a proliferation defect \textit{in vivo} and \textit{in vitro} in response to antigenic and mitogenic stimuli. Lipopolysaccharide stimulation of CCT\( \alpha \)/–/– B-cells resulted in an early trigger of the unfolded protein response-mediated splicing of \textit{Xbp-1} mRNA, and this was accompanied by accelerated kinetics of IgM secretion and higher incidence of IgM-secreting cells. Thus, the inability of stimulated B-cells to produce enough phosphatidylcholine prevents proliferation and class switch recombination but leads to unfolded protein response activation and a hyper-IgM secretion phenotype.

Stimulated B-lymphocytes proliferate and/or differentiate into plasma cells, which synthesize and secrete large amounts of Ig. Both events require a significant increase in cellular membrane phospholipid biosynthesis. Proliferation results in a doubling of the cellular membrane content prior to each cell division (1), whereas plasma cell differentiation is accompanied by a substantial increase in membrane phospholipid to support the expansion of the endoplasmic reticulum (ER) synthetic and secretory apparatus (2, 3). The most abundant membrane phospholipid component is phosphatidylcholine (PtdCho), whose synthesis is regulated by the CTP:phosphocholine cytidyltransferase (CCT). CCT is dynamically regulated during cell cycle progression (1, 4, 5), and increased PtdCho biosynthesis during plasma cell formation is accomplished by a program of genetic and biochemical events that up-regulate the flux through CCT (3). An alternative pathway to PtdCho mediated by the phosphatidylethanolamine \( N \)-methyltransferase contributes only about 5% to the total PtdCho following B-cell stimulation (3). Thus, CCT plays a central role in multiple aspects of membrane biogenesis during B-cell development.

Plasma cell differentiation requires \textit{Xbp-1} (X-box-binding protein 1), a transcription factor regulated by the unfolded protein response (UPR) (6). The \textit{Xbp-1} mRNA is processed by a novel, UPR-mediated splicing mechanism, yielding the transcriptional activator \textit{XBP-1(S)} (7). Forced expression of \textit{XBP-1(S)} activates the cytidine diphosphocholine pathway for PtdCho synthesis and triggers an expansion of the ER compartment (8). Consistent with its key regulatory role, enforced expression of CCT\( \alpha \) is sufficient to increase membrane PtdCho, in contrast to other enzymes in the cytidine diphosphocholine pathway (9). As a cellular response to ER stress, the UPR has been most extensively studied as it relates to protein quality control in the ER, but phospholipid metabolism also appears to be vitally important. CCT inactivation leads to PtdCho depletion, activation of some components of the ER stress response, and cell death in cultured fibroblasts (10). Alteration of the ER lipid composition by accumulation of free cholesterol initiates ER stress in macrophages (11, 12), and CCT inactivation renders macrophages more sensitive to the lethal effects of free cholesterol loading (13). These results suggest that ER phospholipid quality control may impact plasma cell differentiation as well as the UPR.

Three CCT isofoms are known in mammals, CCT\( \alpha \), CCT\( \beta \), and CCT\( \beta \)3 (5, 14), and the different CCT isofoms are biochemically similar in enzymatic activity and regulation (15). CCT\( \alpha \) is the dominant isofom expressed in most tissues, including the CH12 B-cell line (3). Accordingly, deletion of the \textit{Pcyt1a} (CCT\( \alpha \)) gene is lethal prior to embryonic day 3.5 (16), whereas the deletion of CCT\( \beta \) only results in premature reproductive senescence (17). The functions of CCT\( \alpha \) in adult animals have been studied by tissue-specific deletion of CCT\( \alpha \) cytidyltransferase; LPS, lipopolysaccharide; NP-KLH, 4-hydroxy-3-nitrophenylacetyl-conjugated keyhole limpet hemocyanin; KO, knock-out; WT, wild-type.
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gene expression using the Cre-loxP system. These studies have revealed distinct roles for CCTα in professional secretory cells, including the formation and secretion of surfactant by alveolar epithelial cells (18), the assembly and secretion of lipoproteins by hepatocytes (19), and cytokine secretion by stimulated macrophages (20). In this study, the role of CCTα in B lymphocyte function was examined by selectively deleting the Pcyt1α gene encoding CCTα in murine B-cells. We found that XBP-1(S) expression and IgM secretion were activated with accelerated kinetics and more robustly in stimulated CCTα-deficient B-cells. However, compromised PtdCho synthesis in CCTα-deficient B-cells correlated with severely reduced proliferation and only minimal Ig class switching. Thus, by regulating the supply of PtdCho, CCTα plays a pivotal role in determining the function and the fate of activated B-cells.

EXPERIMENTAL PROCEDURES

Generation of B-cell-specific CCTα Knock-out (KO) Mice and B-cell Isolation—Pcyta1flo/llo mice (13) were bred with Pcyta1flo/llo/CD19cre/cre mice to obtain Pcyta1flo/llo/CD19cre/cre (KO) mice. Mouse tails were genotyped as previously described (13) and following the protocol for Cre recombinase detection from The Jackson Laboratory. All procedures involving mice were performed according to protocols approved by the Institutional Animal Care and Use Committee of St. Jude Children’s Research Hospital. Splenic B-cells were isolated from wild type (WT) (Pcyta1flo/llo) or KO mice by either a depletion strategy using the B-cell isolation kit or a positive strategy using the Ficoll-PaqueTM PLUS (GE Healthcare) following a one-step gradient sedimentation using Ficoll-PaqueTM PLUS (GE Healthcare) following a published protocol (21). Pcyt1aα−/− and Pcyt1aα−/− genotypes were determined by using the same primers used previously (13), and samples were run on a 2% agarose gel. PCR products were stained with ethidium bromide and quantified using a Typhoon 9200 scanner (GE Healthcare) controlled by Typhoon Scanner Control software, version 2.0 (GE Healthcare) together with ImageQuant TL software, version 2003.02 (Amersham Biosciences). Values were normalized to the number of bp for each PCR product.

Cell Genotyping—DNA was extracted from WT and KO B-cells resuspended in TRizol® (Invitrogen), following the manufacturer’s directions. Because of the low viability 72 h after LPS stimulation, WT and KO B-cells were enriched in viable cells (more than 90%) by one-step gradient sedimentation using Ficoll-Paque™ PLUS (GE Healthcare) following a published protocol (21). Pcyt1aα−/− and Pcyt1aα−/− genotypes were determined by using the same primers used previously (13), and samples were run on a 2% agarose gel. PCR products were stained with ethidium bromide and quantified using a Typhoon 9200 scanner (GE Healthcare) controlled by Typhoon Scanner Control software, version 2.0 (GE Healthcare) together with ImageQuant TL software, version 2003.02 (Amersham Biosciences). Values were normalized to the number of bp for each PCR product.

Immunocytochemistry—Wild-type and KO mice were sacrificed, and the spleens were collected, embedded in OCT™ cryoprotective medium, and frozen on dry ice. Spleens were sliced, 4-μm thickness, using a microtome cryostat HM 505 (MICROM International Gmbh), and fixed in acetone at −20 °C. Tissue slides were incubated first with biotinylated rat anti-mouse IgD (eBiosciences) and rabbit anti-mouse CCTα, previously characterized (22), followed by incubation with AlexaFluor®488 mouse anti-biotin and AlexaFluor®594 goat anti-rabbit IgG (Molecular Probes). Splenic germinal centers were detected by staining with fluorescein isothiocyanate-conjugated peanut agglutinin (Sigma) and R-phycocerythrin-labeled anti-mouse B220 (BD Biosciences). Apoptotic cells were detected in spleen sections from WT and KO mice using the ApopTag® fluorescein in situ apoptosis detection kit from Chemicon International, following the manufacturer’s protocol. Images were acquired using an Olympus BX41 microscope equipped with an UPlanFl ×20/0.50 objective and a 7.3 Three Shot color camera from Diagnostic Instruments, Inc., controlled by SPOT software, version 4.0.4PC, from Diagnostic Instruments, Inc.

CCT Activity Measurement and Metabolic Labeling—CCT activities in B-cell lysates were measured as previously described (3). [methyl-3H]Choline incorporation into PtdCho was performed as described previously (3) using medium containing 6 μM choline and supplemented with 16.4 μCi/ml [3H]choline.

RNA and Lipid Measurements—Total RNA was isolated and analyzed as previously reported (3). The amount of target RNA was normalized to either the endogenous glyceraldehyde-3-phosphate dehydrogenase reference or to the total amount of RNA used to prepare the cDNA. The relative expression of XBP-1 and XBP-1(S) was determined using primers and procedures described elsewhere (23). Lipids from B-cells were quantified as described previously (3).

Flow Cytometry Analysis of B-cells—Splenocytes and cells from the peritoneal cavity were depleted of red blood cells and labeled with anti-mouse B220, anti-mouse IgM, and anti-mouse CD3 (BD Biosciences) in the presence of anti-mouse CD16/32 (eBiosciences). Samples were analyzed using a FACSCalibur cytometer (BD Biosciences), and data were acquired using CellQuest™ Pro (version 5.2.1) software (BD Biosciences) and analyzed using WinMDI (version 2.8) software (Joseph Trotter).

B-cell Proliferation in Vitro—Purified B-cells were suspended in RPMI 1640 culture medium supplemented with 10% fetal bovine serum, seeded at 5 × 10⁶ cells/ml in a 96-well plate, and cultured for 72 h with 10 μg/ml Escherichia coli LPS (E. coli 055:B5; Sigma). Proliferation was measured by adding [3H]thymidine (63 Ci/mmol; American Radiolabeled Chemicals) to the medium (final 50 μCi/ml) and incubating for 6 h. Cells were harvested using MF™-membrane filters (0.45-μm pores; Millipore) and washed, and the radioactivity was measured by scintillation spectroscopy.

B-cell Proliferation in Vivo—Wild-type and KO mice were injected subcutaneously with 30 μg of NP-KLH dispersed in TiterMax Gold adjuvant (Sigma). After 7 days, mice were sacrificed, spleens were isolated, and the total splenic B-cell numbers were determined by flow cytometry of B220-positive cells.

Immunization and Immunoglobulin Measurements—Wild-type and KO mice were injected subcutaneously with 30 μg of NP-KLH dispersed in TiterMax Gold adjuvant (Sigma). Prior to injection and at the indicated times following immunization, blood was collected, and the serum IgM and IgG content (both λ- and κ-chain) was measured by enzyme-linked immunosorbent assay. IgM and IgG were quantified using goat anti-mouse IgM (μ-chain-specific), goat anti-mouse IgG (γ-chain-specific), goat anti-mouse κ- or λ-alkaline phosphatase (Southern-
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Biotech), and 4-methylumbelliferyl phosphate (Sigma) as substrate.

Antibody Secretion in Vitro—Purified B-cells were plated and stimulated with LPS. At times before and after stimulation, cells were collected, washed, counted, and cultured for 4 h in fresh medium. The amount of IgM in the medium was measured by enzyme-linked immunosorbsorbent assay. The number of IgM-secreting cells was assessed by an enzyme-linked immunosorbsorbent spot assay. In practice, after washing the cells before and after stimulation, the cells were cultured for 4 h on MultiScreen™-1P plates (Millipore) coated with goat anti-mouse IgM. After washing the plate to remove the cells, the plate was treated with a mixture of goat anti-mouse κ- and λ-alkaline phosphatases, and the IgM spots were revealed by incubation with 5-bromo-4-chloro-3-indolyl phosphate and p-nitro blue tetrazolium salt (Sigma).

Cell Cycle Arrest with Aphidicolin—Purified B-cells were stimulated with LPS with or without 2 μg/ml aphidicolin (Sigma). After 24 h, cells were harvested, and the number of secreting cells was estimated by enzyme-linked immunosorbsorbent spot assays.

RESULTS

Selective Deletion of CCTα in B Lymphocytes—Mice were derived whereby exons 5 and 6 of the Pcyt1a gene, encoding CCTα, were deleted in B lymphocytes. This was accomplished using Cre/loxP-mediated gene deletion under the transcriptional control of the B-cell lineage-restricted CD19 gene (24). Mice homozygous for the “floxed” CCTα allele (Pcyt1a<sup>fl/fl</sup>/CD19<sup>cre+/−</sup>) were crossed with mice with a Cre-expressing construct (Pcyt1a<sup>fl/fl</sup>) to delete the CCTα allele. The resulting KO mice were crossed with mice expressing Cre under the control of the B220<sup>+/−</sup> allele (Pcyt1a<sup>fl/fl</sup>/CD19<sup>cre</sup>) to delete the CCTα allele in B cells. The resulting KO mice were homozygous for the “floxed” CCTα allele (Pcyt1a<sup>fl/fl</sup>/CD19<sup>cre−/−</sup>), and they were crossed with mice expressing Cre under the control of the B220<sup>+/−</sup> allele (Pcyt1a<sup>fl/fl</sup>/CD19<sup>cre</sup>) to delete the CCTα allele in B cells. The resulting KO mice were homozygous for the “floxed” CCTα allele (Pcyt1a<sup>fl/fl</sup>/CD19<sup>cre−/−</sup>). B-cells were isolated from the spleens of KO and WT mice and quantified (Fig. 3A). Splenic B-cells, which are primarily B2-cells originating from adult bone marrow (26), were reduced by about 25%. In contrast, the B-cell abundance in WT control mice was reduced by about 60% compared with B-cells from WT animals (Fig. 2A, inset). The contribution of CCTβ2 to the total level of CCT activity in the KO cells accounted for the slightly higher levels of CCT activity in the KO cells than would be predicted from the reduction in the CCTα mRNA abundance. However, the reduction in CCTα transcript and total CCT activity did not affect the amounts of the major membrane lipid components in the B-cells (Fig. 2B). These data are consistent with earlier analytical results with macrophages (20) or hepatocytes (19), which were selectively deleted in CCTα expression and possessed normal amounts of PtdCho in vivo. The results indicated that the cellular PtdCho level in the KO B-cells was sustained by CCTβ2 and/or the acquisition of PtdCho from extracellular sources, such as serum high density lipoprotein (25). A reduction in the rate of PtdCho synthesis via the CCTα-dependent cytidine diphosphocholine pathway was confirmed by a significant decrease in the rate of PtdCho synthesis measured by the metabolic labeling of intact B-cells (Fig. 2B, inset). Thus, CCTα expression and de novo PtdCho synthesis were reduced in a significant proportion of B lymphocytes in the KO animals. However, between 20 and 30% of the splenic B-cell population in the KO animals expressed wild-type CCTα, due to incomplete penetrance of Cre recombination expression, as has been reported before (24).

B-cell Population and Serum Ig Levels in B-cell-specific CCTα KO Mice—B-cells were isolated from several locations in WT and KO mice and quantified (Fig. 3A). Splenic B-cells, which are primarily B2-cells originating from adult bone marrow (26), were reduced by about 25%. In contrast, the B-cell abundance in
adult bone marrow was equivalent in WT and KO animals (Fig. S7). CCTα deletion in the KO bone marrow B-cells was confirmed by PCR (data not shown). There was no reproducible change in the distribution of marginal zone cells or follicular cells isolated from KO spleens (data not shown). On the other hand, peritoneal B-cells, primarily B1-cells that are a self-renewing population produced by fetal liver, were reduced by 75% in the CCTα-deficient animals. These results suggested that proliferation and/or survival of the peripheral B-cell populations were compromised by the loss of CCTα expression. Terminal deoxynucleotidyltransferase-mediated dUTP nick-end-labeling staining of splenic sections from KO animals did not show increased apoptosis compared with spleens from WT animals (Fig. S1A), and cell viability in both KO and WT B-cell populations was ≥80% following stimulation for 48 h in vitro (Fig. S1B). Thus, these data indicated that the CCTα deficiency primarily affected B-cell proliferation in the periphery.

Basal serum immunoglobulin levels in the KO mice were also affected (Fig. 3B). Although the levels of circulating IgM were not statistically different between the WT and KO animals, there was a 75% drop in the serum levels of IgG in the KO mice. These data indicated that the KO mice were selectively deficient in the production of IgG.

**Proliferative Response of CCTα-deficient B-cells**—B lymphocytes were isolated from WT and KO spleens and stimulated in vitro to evaluate the proliferation of CCTα-deficient cells. Proliferation, as measured by thymidine incorporation into DNA, was significantly lower in the KO population in response to either anti-IgM, which stimulates the B-cell antigen receptor (Fig. 4A), or bacterial LPS, an activator of the Toll-like receptor 4 (Fig. 4B). The low level of [3H]thymidine incorporation that was observed in B-cells from the KO animals was explained by the fraction that still expressed CCTα. This result was not unexpected, because it was clear from previous work that CCTα was essential for the proliferation of immortalized cells, although inhibition of CCT activity in those systems resulted in apoptosis (27). Although the KO B-cells were unable to proliferate, receptor signaling mechanisms appeared to be intact, as demonstrated by tyrosine phosphorylation after stimulation with anti-IgM (Fig. S2). Since the WT cells would eventually become dominant in the KO population due to their proliferative advantage, receptor signaling mechanisms appeared to be intact, as demonstrated by tyrosine phosphorylation after stimulation with anti-IgM (Fig. S2).
data indicated that the CCTα-deficient B-cells had a defect in proliferation in response to immunization in vivo.

Antibody Production in B-cell-specific CCTα KO Mice—Antibody production was evaluated in wild-type and KO mice in response to immunization with NP-KLH. The response to immunization with NP-KLH is dependent on the interaction between T-cells and B-cells and is accompanied by B-cell proliferation in the spleen and antibody production by differentiated plasma cells. The proliferation and differentiation process results in a shift from synthesis and secretion of IgM by the naive B-cells to the production of higher affinity IgG by the differentiated plasma cells. The normal production of IgG in response to NP-KLH was significantly reduced in the KO animals (Fig. 5B). In contrast, the KO animals produced significant amounts of IgM rather than IgG (Fig. 5A). These results indicated that despite the reduced CCTα activity (Fig. 6A), the KO B-cells were capable of secreting antibodies but were unable to proliferate and undergo isotype switching. This was a general feature of the KO B-cells, since the IgG response to immunization with a T-cell-independent antigen, NP-Ficoll, was blunted as well (Fig. S5). The ability of CCTα-deficient B-cells to secrete large amounts of IgM was corroborated in vitro (Fig. 6B). Both before and after stimulation with LPS, the CCTα−/− B-cell population secreted IgM at a significantly higher rate compared with the WT B-cells. B-cells can secrete small amounts of IgG in vitro, in addition to IgM, and we measured this feature in both WT and KO cells. The KO B-cells secreted more IgG with earlier kinetics compared with cultured WT B-cells, similar to the results for IgM secretion (Fig. S6). These results suggested an early activation of plasma cell differentiation in the KO animals.

Activation of the UPR in CCTα-deficient B-cells—The primary biochemical defect in CCTα KO B-cells was demonstrated to be a reduction in the rate of PtdCho synthesis (Fig. 2B), and stimulation imposed an even greater demand for PtdCho (Fig. 6A). The correlation between the reduced PtdCho synthesis and elevated IgM secretion in the KO cells (Fig. 6B) was counterintuitive, based on the implied need for membrane PtdCho expansion during plasma cell differentiation (3). Our observations suggested that PtdCho deficiency triggered the secretion program in B-cells. We hypothesized that the mechanism involved in B-cell differentiation and in the response to PtdCho deficiency was activation of the UPR. The splicing of XBP-1 to form XBP-1(S) is a hallmark of UPR activation, and the XBP-1(S) is a key transcription factor in the control of membrane phospholipid biogenesis (9). We found that the CCTα-deficient B-cell population had significantly higher levels of XBP-1(S) mRNA after LPS stimulation, and the amount of this UPR-generated transcript increased substantially faster compared with the kinetics of activation in WT cells (Fig. 7A). This elevation in XBP-1(S) correlated with a higher number of
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FIGURE 6. PtdCho biosynthesis and IgM secretion in vitro by WT and KO B-cells in response to LPS. A, the rate of incorporation of [3H]choline into PtdCho was determined before and at 24 or 48 h after the addition of LPS to KO (dark gray) or WT (light gray) purified splenic B-cells. B, the amount of IgM secreted in vitro during a 1-h period from KO (dark gray) or WT (light gray) B-cells was determined before and 24 or 48 h after stimulation with LPS. The data are the mean ± S.E. from at least three independent determinations in two independent experiments.

FIGURE 7. IgM secreting cells and Xbp-1 mRNA splicing following B-cell stimulation with LPS. WT (light gray) and KO (dark gray) B-cells were purified by negative selection and cultured in the presence or absence of LPS. A, real-time quantitative reverse transcription-PCR detected the expression of unspliced and spliced Xbp-1 mRNA before and after stimulation with LPS for 16 or 24 h. B, the number of IgM-secreting B-cells was determined using an enzyme-linked immunosorbent spot assay, and the WT value was set at 1. The data are the mean ± S.E. from at least three independent determinations in two independent experiments.

FIGURE 8. Effect of aphidicolin on proliferation and secretion of WT B-cells. Purified splenic WT B-cells were stimulated with LPS for 24 h with (dark gray) or without (gray) the addition of aphidicolin (2 μg/ml). A, proliferation was measured by [3H]thymidine incorporation into cells. B, IgM secretion was measured by enzyme-linked immunosorbent spot assays. The data are the mean ± S.E. from three independent determinations in two independent experiments.

Ig-secreting cells in the CCTα−/− population (Fig. 7B) corresponded to the increased rate of secretion observed in the stimulated KO B-cells (Fig. 6 and Fig. S6). These data indicated an accelerated activation of the UPR in the stimulated KO cells that, in turn, promoted antibody secretion. Since both CCT deficiency (28, 29) and pharmacologic induction of the UPR (30) block proliferation in immortalized cell lines, we tested whether simply blocking cell proliferation in LPS-stimulated B-cells by another means would trigger IgM secretion as well as halt proliferation. However, aphidicolin, an inhibitor of DNA synthesis that blocks cells in the G1-S phase transition, did not change the kinetics of secretion in WT cells, despite an inhibition of thymidine incorporation (Fig. 8). These data indicated that an inability to undergo cell division does not necessarily commit B-cells toward terminal differentiation into antibody-secreting cells. Rather, we propose that the early and potent induction of XBP-1(S) in CCTα-deficient B-cells accelerates and augments the transition into antibody secretion (Fig. 9).

DISCUSSION

The interruption of membrane phospholipid synthesis in CCTα-deficient cells elicited by a block in PtdCho formation at the CCTα step results in early activation of the UPR, as demonstrated by Xbp-1 mRNA splicing (Fig. 7). Induction of XBP-1(S) mRNA correlates with the inhibition of proliferation (Fig. 4) and an increased rate of IgM secretion from a greater number of stimulated B-cells (Figs. 6 and 7). XBP-1(S) is the active transcription factor that is derived from the splicing of the Xbp-1 mRNA by IRE1α (inositol-requiring 1α) in response to ER stress (7). XBP-1 is an essential prerequisite for immunoglobulin secretion via the ER (31). XBP-1−/− B-cells are found in normal numbers in vivo, but in response to stimulation they secrete neither IgM nor IgG at normal levels (6). IRE1α, an integral ER membrane protein and proximal transducer of the UPR, oligomerizes in response to the accumulation of misfolded proteins in the ER lumen to activate its cytosolic ribonuclease domain. Activated IRE1α then executes site-specific cleavage of Xbp-1 transcript to initiate the UPR-mediated splicing mechanism that yields XBP-1(S) mRNA (23, 32). To date, the only known way to initiate the novel splicing of XBP-1 transcripts requires that the IRE1 ribonuclease site-specifically cleave the XBP-1 mRNA, and to date, the only known ways to activate the IRE1 protein involve some type of event that “stresses” the ER, such as disruption of protein folding in the ER, augmentation of protein traffic through the ER, or disruption of lipid biosynthesis. One function of XBP-1(S) appears to involve directing increased production of phospholipids necessary for intracellular membrane biogenesis (9), and CCTα plays a key role providing PtdCho to support ER expansion in differ-

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entitatively B-cells (8). In addition, XBP-1(S) is required for increased expression of a large cohort of genes that enhance both the protein biosynthetic and secretory capacity of differentiating B-cells (33). Thus, the enhanced IgM secretion in stimulated CCTα−/− B-cells reflects an acceleration of the normal processes that occur in B-cells following the activation of Xbp-1 splicing.

Previous work has established that synthesis of Ig μ-heavy chain is required for maximal induction of XBP-1(S) in LPS-stimulated B-cells (34). This fits well with the observation that the level of XBP-1(S) peaks after maximal synthesis of μ-chains has been achieved (35, 36). However, it has also been observed that XBP-1(S) synthesis initiates prior to the strongest increase in μ-chain synthesis (35, 37), suggesting that another signal(s) might contribute to the activation of UPR-mediated Xbp-1 mRNA splicing in the early stages of LPS stimulation. Indeed, our studies of CCTα-deficient B-cells reveal that the IRE1α-XBP-1 branch of the UPR responds to increased demand for phospholipids as well as increased demand on the protein folding capacity of the ER. In agreement with our findings, the inhibition of PtdCho syntheses (10) and fatty acid synthesis (38) have both been shown to elicit activation of UPR components. Depletion of ER phospholipids or alteration of the ratio between PtdCho and other membrane lipids may lead to ER membrane protein aggregation and activation of IRE1α to trigger the splicing of Xbp-1 mRNA and, perhaps, activation of the UPR in general. Alternatively, the defect in producing membrane phospholipids for the ER may adversely affect the folding and insertion of newly synthesized ER membrane protein or interfere with vesicular trafficking in the secretory apparatus. These types of perturbations could certainly lead to an inappropriate build-up of proteins in the ER and, in turn, activate IRE1α by established mechanisms. It will be of interest to assess the activation status of other proximal transducers of the UPR, PERK (PKR-like ER kinase) and ATF6 (activating transcription factor 6), in the CCTα-deficient B-cells. We propose that the signals and mechanisms underlying activation of the UPR warrant further investigation.

Inhibition of PtdCho synthesis by pharmacological or genetic reduction in CCT activity is known to block cell proliferation and trigger cell death in cultured immortalized cell lines (10, 39, 40). However, CCTα−/− primary animal cells survive and differentiate, as exemplified by macrophages (13), hepatocytes (19), lung alveolar cells (18), and B-cells (this study). These results strongly suggest that another source of PtdCho is available that circumvents a requirement for CCTα. It is likely that the uptake of serum lipoproteins (25), coupled with a contribution from CCTβ expression, is sufficient to support development. On the other hand, B-cell numbers do not increase following immunization of KO animals, and the challenge of rapid proliferation in stimulated B-cells reveals a requirement for de novo PtdCho synthesis. The inability to proliferate accounts for the lack of IgG production by CCTα-deficient B-cells, since class switch recombination is dependent on the DNA synthesis that accompanies proliferation. Interestingly, the low IgG levels and the hyper-IgM secretion phenotype exhibited by the CCTα-deficient animals in response to immunization are similar to the phenotypes of murine class switch recombination deficiencies. However, it is difficult to envision how a reduction in PtdCho synthesis would lead to a block in class-switch recombination, particularly since phospholipid synthesis and DNA synthesis are not co-dependent (4). In light of our analysis of CCTα-deficient B-cells and previous data showing that UPR-mediated splicing of Xbp-1 mRNA is a prerequisite for terminal differentiation of B-cells into antibody-secreting plasma cells (34), we speculate that XBP-1(S) might play a role in the cessation of B-cell proliferation. Further investigation of these questions should provide additional insight into the complex relationship of increased need for membrane lipids and protein production capacity, the UPR, and the function and fate of activated B-cells.

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