

INNATE IMMUNITY

The interaction between IKK α and LC3 promotes type I interferon production through the TLR9-containing LAPosome

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Toll-like receptor 9 (TLR9) recognizes DNA in endosomes and activates distinct signaling pathways to stimulate the production of proinflammatory cytokines and type I interferons (IFNs). The assembly of signaling platforms on microtubule-associated proteins 1A/1B–light chain 3 (LC3)–decorated endosomal vesicles is required to transduce TLR9 signals that stimulate the production of IFN but not interleukin-12 p40 (IL-12p40). LC3-associated phagocytosis (LAP), a form of noncanonical autophagy, is critical for the activation of interferon regulatory factor 7 (IRF7) and for IFN synthesis. We showed that after the stimulation of TLR9 by CpG oligonucleotides, the autophagy protein LC3 and the kinase IKK α were recruited to endosomes that contained TLR9. The recruitment of IKK α and LC3 to such signaling endosomes was not stimulated by catalysts of classical autophagosome formation but involved LAP formation, which required ATG5 but not FIP200. In addition, we found that the LC3–IKK α complex further associated with both TRAF3 and IRF7. We identified three putative LC3-interacting regions (LIRs) in IKK α , and mutagenesis suggested that two of these were critical for direct binding to LC3. Moreover, mutation of the same LIR sequences failed to rescue type I IFN production in IKK α -deficient dendritic cells upon reconstitution. Together, these data suggest a direct link between LAP formation and IKK α recruitment downstream of TLR9 activation that is necessary to facilitate type I IFN production.

INTRODUCTION

Innate immune cells are equipped with an array of pattern recognition receptors (PRRs) that recognize evolutionarily conserved pathogen-associated molecular patterns and activate the production of anti-microbial defense effector molecules. Toll-like receptor 9 (TLR9) is a PRR expressed in the endosomes of immune cells that recognizes DNA (1). Upon ligand engagement, TLR9 activates two distinct downstream signaling pathways to stimulate the production of proinflammatory cytokines and type I interferons (IFNs), including IFN- α and IFN- β (2). DNA-dependent dimerization of TLR9 stimulates the recruitment of MyD88 (or myeloid differentiation primary response 88) and the following signaling molecules: interleukin-1 (IL-1) receptor–associated kinase 4 (IRAK4), IRAK2, tumor necrosis factor (TNF) receptor–associated factor 6 (TRAF6), transforming growth factor β -activated kinase 1 (TAK1), and its regulatory components TAK1-binding protein 2/3 (TAB2/3). This signaling complex ultimately triggers the transcription of proinflammatory cytokines through activation of nuclear factor κ B (NF- κ B), IFN regulatory factor 5 (IRF5), and mitogen-activated protein kinases (2). Activation of TLR9 also stimulates a second MyD88-dependent pathway that requires IRAK1 (3), TRAF3 (4, 5), inhibitor of NF- κ B kinase α (IKK α) (6), osteopontin (7), dedicator of cytokinesis 2 (8), a plasmacytoid dendritic cell (pDC)–specific receptor (pDC-TREM) (9), viperin (10), and adaptor protein 3 (11). This pathway results in IRF7 phosphorylation and subsequent type I IFN production.

In macrophages stimulated with the TLR9 ligand–like DNA-containing immune complexes or CpG, microtubule-associated protein 1A/1B–light chain 3 (LC3), an autophagy-associated protein, is recruited to the phagosome containing TLR9 independently of canonical autophagy through a process known as LC3-associated

phagocytosis (LAP) (12). LAP facilitates the fusion between zymosan-containing phagosomes and lysosomes to mediate rapid acidification of the cargo-containing phagosomes in a TLR2-dependent manner (13). This process is associated with the recruitment of LC3, Beclin-1, and phosphatidylinositol 3-kinase (PI 3-kinase or PI3K), as well as the components of the ubiquitin-like conjugation system, autophagy-related 5 and 7 (ATG5 and ATG7). Similarly, autophagy protein–dependent vacuole formation is also observed in macrophages treated with other TLR ligands such as lipopolysaccharide (LPS; TLR4 ligand), *Porphyromonas gingivalis* LPS and lipoteichoic acid (PAM3CSK4; a TLR1/2 ligand), and DNA-immune complex and CpG oligodeoxynucleotides (TLR9 ligands) (12, 13). Notably, LAP is required for TLR9-mediated IRF7 activation and production of type I IFN but not for IL-12p40 secretion (12).

Three distinct features of LAP distinguish it from canonical autophagosomes. First, unlike the double-membrane structures characteristic of canonical autophagosomes, LAP generates single membrane–enveloped vesicles coated with LC3 on the cytosolic side of the phagosomal membrane. Second, LAP initiation occurs independently of proteins associated with the autophagy preinitiation complex, namely, Unc-51–like autophagy activating kinase 1/2 (ULK1/2), retinoblastoma-associated protein–inducible coiled-coil protein 1 (RB1CC1 or FIP200), or ATG13, which are required for initiation of canonical autophagy. Finally, the class III PI3K-associated protein, rubicon, is required for LAP (14). It stabilizes PI3K catalytic subunit type 3 (PI3KC3 or VPS34) activity to sustain PtdIns(3)P presence on LAP-engaged phagosome (LAPosome) and stabilize the cytochrome b-245 heavy chain complex for reactive oxygen species production, both of which are critical for progression of LAP. Other autophagy-related genes identified to be involved in LAP formation are other components of class III PI3K-associated proteins, Beclin-1, VPS34, ATG14, UVRAG (ultraviolet radiation resistance associated or VPS38) (14), and components of ubiquitin-like conjugation system (ATG5 and ATG7) (14).

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Here, we examined the relationship between IKK α and LAP formation after TLR9 stimulation. We found that LC3 was recruited to TLR9-containing LAPosomes and provided an anchor for IKK α recruitment. We identified two LC3-binding motifs within IKK α that were critical for both its interaction with LC3 and the stimulation of type I IFN production downstream of TLR9. Our findings provide a missing mechanistic link between IKK α - and LAP-dependent signaling downstream of TLR9.

RESULTS

IKK α is recruited to TLR9-containing endosomes and associates with LC3 after stimulation with CpG-A

Because LAP and IKK α are required for TLR9 signaling and IFN- α production, we examined the localization of IKK α and LC3 after transfection of macrophages with cationic lipid vesicles [DOTAP or 1,2-di-(9Z-octadecenoyl)-3-trimethylammonium-propane (chloride salt)] containing the TLR9 ligand CpG-A. In macrophages, this approach delivers TLR9 ligands to the appropriate cellular compartment to stimulate robust IFN- α production downstream of TLR9 (15). We found that IKK α and LC3 colocalized only after DOTAP-CpG-A stimulation in mouse fetal liver macrophages and RAW264.7 cells (Fig. 1A and fig. S1A). Throughout this study, LC3 refers to the LC3b isoform. After the activation of TLR9 in macrophages, both early [vesicle-associated membrane protein 3 (VAMP3⁺)] and late [lysosome-associated membrane protein 1 (LAMP1⁺)] endosomes recruited IKK α and LC3 (Fig. 1A, top and bottom, respectively). IKK α and LC3 also colocalized with TLR9-containing endosomes after stimulation of fetal liver macrophages or RAW264.7 cells stably expressing hemagglutinin (HA)-tagged TLR9 (TLR9-HA) and mCherry-tagged LC3 (Cherry-LC3) (Fig. 1B and fig. S1B). A time course analysis revealed that IKK α and LC3 began to colocalize as early as 30 min after CpG-A stimulation and decorated large LAMP1⁺ endosomes by 3 to 6 hours after stimulation (Fig. 1C). Immunoprecipitation analysis of lysates of human embryonic kidney-293T (HEK293T) cells transfected with IKK α -HA and green fluorescent protein (GFP)-LC3 showed that IKK α and LC3 physically interacted with each other (Fig. 1D). This interaction was

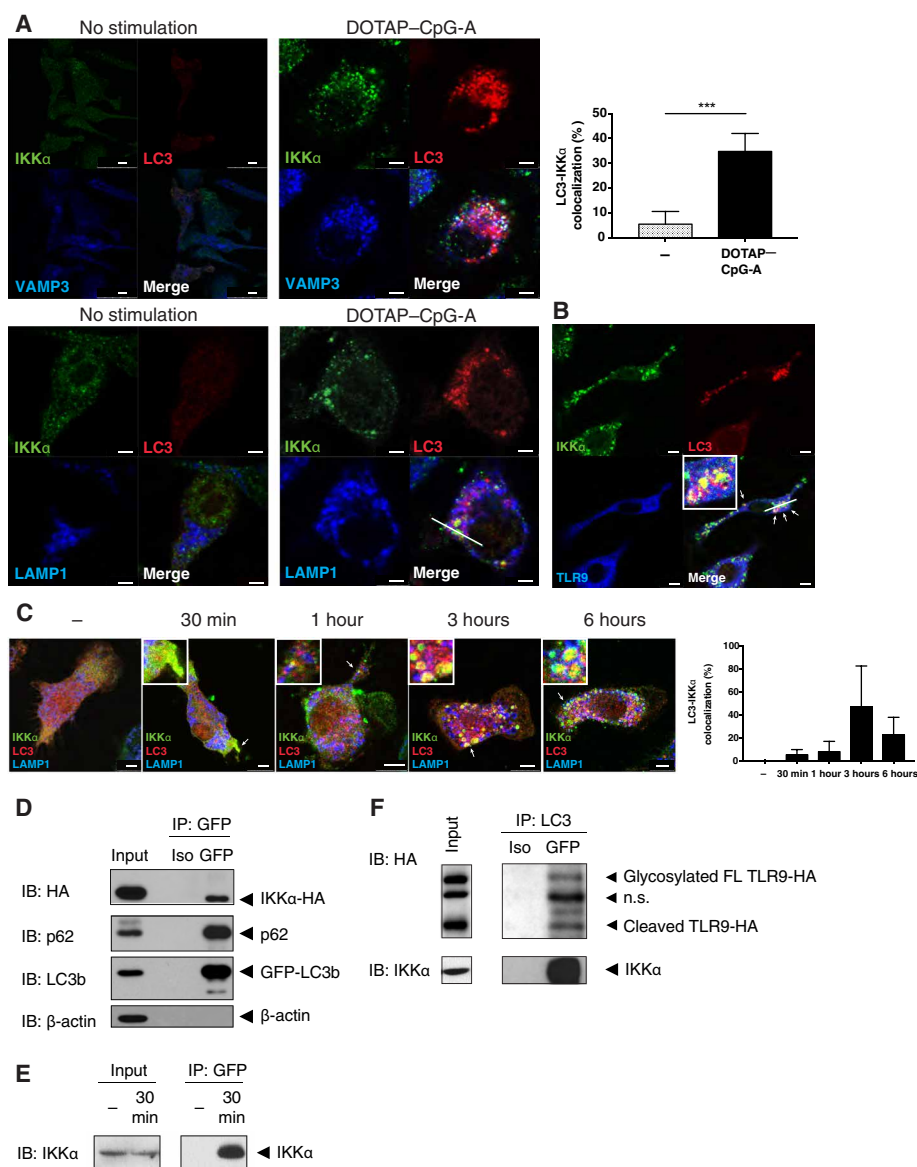


Fig. 1. IKK α binds to LC3 after DOTAP-CpG-A stimulation. (A to C) Confocal microscopy images from *Ikk α* knockout (KO) fetal liver macrophages that were retrovirally co-transduced with wild-type (WT) IKK α -HA, TLR9-GFP, and Cherry-LC3 plasmids and then stimulated with 3 μ M DOTAP-CpG-A. Images are representative of three or more independent experiments. Scale bars, 2.5 μ m. Quantified co-localization data are means \pm SEM. *** P < 0.001 by t test. (D) HEK293T cells were co-transfected with WT IKK α -HA and GFP-LC3, and then cell lysates were immunoprecipitated (IP) for GFP and blotted for the indicated protein. Blots are representative of three or more independent experiments. IB, immunoblot. (E) Day 7 BMMs were stimulated with DOTAP-CpG-A for 30 min, and then the pre-nuclear cell lysates were immunoprecipitated for GFP-LC3 and blotted for the indicated proteins. Blots are representative of three or more independent experiments. (F) RAW264.7 cells stably expressing TLR9-HA and Cherry-LC3 were stimulated with DOTAP-CpG-A for 6 hours, and then cell lysates were immunoprecipitated for LC3 and blotted for the indicated proteins. Blots are representative of three or more independent experiments. n.s., nonspecific.

confirmed in primary bone marrow-derived macrophages (BMMs) and was found within 30 min of stimulation with DOTAP-CpG-A (Fig. 1E). Furthermore, IKK α -LC3 complexes also contained TLR9 (Fig. 1F). These data indicate that IKK α was mobilized upon DOTAP-CpG-A stimulation to bind to LC3 on TLR9-containing endosomes.

Canonical autophagy induction is not sufficient to recruit IKK α to LC3

To determine whether LC3-IKK α interactions occurred as a consequence of the activation of canonical autophagy, we stimulated autophagy in BMMs and examined the colocalization of IKK α and LC3. Autophagy can be induced in various ways, such as serum starvation or treatment with a Tat-Beclin-1 peptide (16). Although confocal analysis showed that autophagy was induced in response to both stimuli, as expected, as measured by the formation of LC3 puncta (Fig. 2, A to D), colocalization between IKK α and LC3 was not observed under these conditions. Similar results were obtained when we analyzed BMMs for IKK α and LC3 colocalization after treatment with DOTAP alone (Fig. 2A). These data indicate that neither ca-

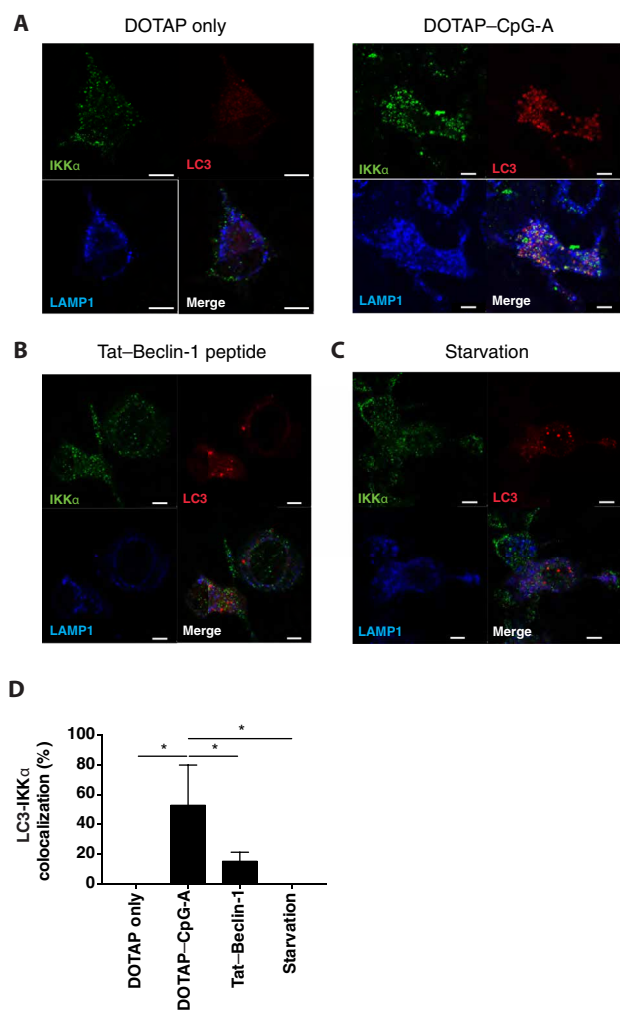


Fig. 2. Autophagy induction is not sufficient to trigger LC3-IKK α interaction. (A to C) WT hematopoietic stem cells (HSCs) were retrovirally transduced with retrovirus expressing Cherry-LC3 and then differentiated into BMMs that were either treated with DOTAP alone (A), 3 μ M DOTAP-CpG-A (A), or 1 μ M Tat-Beclin-1 peptide (B), or starved in serum-deprived media (C) for 6 hours. After treatment, cells were stained for the indicated proteins and analyzed by confocal microscopy. Images are representative of three or more independent experiments. Scale bars, 5 μ m. (D) Percentage of IKK α -LC3 colocalization from confocal analysis BMMs expressing Cherry-LC3 (A to C). Data are means \pm SEM from three or more independent experiments. * P < 0.05, ** P < 0.01 by Student's t test.

nonical autophagy induction alone nor exposure to cationic lipid was sufficient to trigger IKK α -LC3 complex formation. Furthermore, these results suggest that the interaction between IKK α and LC3 on endosomes required CpG stimulation.

IKK α binding to LC3 requires LAP but not canonical autophagy

Because noncanonical autophagy is required for LAP formation in response to TLR9 stimulation (12), we tested the involvement of this pathway in TLR9-mediated IKK α -LC3 colocalization. ATG5 is critical for both canonical autophagy (17) and for LAP formation (18). When fetal liver macrophages from *Atg5*^{-/-} mice were stimulated with DOTAP-CpG-A, no colocalization between IKK α and LC3 was found (Fig. 3A). However, ATG5 did not interact with IKK α or LC3 in immunoprecipitation studies (fig. S2), suggesting that the requirement for ATG5 for IKK α -LC3 complex formation is likely upstream of complex formation. As shown previously (19), *Atg5*-deficient macrophage cells were unable to secrete IFN- α but secreted similar amounts of proinflammatory cytokine IL-12 to WT cells (Fig. 3B). These data suggested that although IRF7-mediated IFN- α production was impaired in the absence of ATG5, proinflammatory cytokine production downstream of NF- κ B activation was not. In contrast to ATG5, activation of the autophagy preinitiation complex that includes FIP200 is required for the induction of autophagy but not LAP formation (12). Note that we found that IKK α still colocalized with LC3 in *Fip200*^{-/-} macrophages stimulated with DOTAP-CpG-A (Fig. 3C). Similarly, when *Fip200*-deficient bone marrow-derived dendritic cells (BMDCs) were stimulated through TLR9 with DOTAP-CpG-A, we found no defect in the production of IFN- α , IFN- β , or IL-12p40 (Fig. 3D). Together, these data indicate that LAP formation, but not canonical autophagy, was required for the interaction between LC3 and IKK α upon TLR9 stimulation and for the signaling required for type I IFN production.

The LC3-IKK α complex recruits TRAF3 and IRF7

To determine whether LC3-IKK α complexes recruited other key molecules involved in type I IFN production, HEK293T cells were co-transfected with plasmids encoding IKK α -HA, GFP-LC3b, and TRAF3. When cell lysates were subjected to immunoprecipitation of IKK α , we found that IKK α associated with TRAF3 and that this association was enhanced in the presence of LC3 (Fig. 4A). We also examined the recruitment to the LC3-IKK α complex of IRF7, the key transcription factor for the expression of type I IFN genes downstream of TLR9. Using a similar approach, we co-transfected HEK293T cells with plasmids encoding IKK α -HA, GFP-LC3b, and IRF7-Flag, as well as plasmids encoding MyD88, TRAF6, and TRAF3, to better mimic TLR9 signaling (6). Cell lysates were subjected to immunoprecipitation of the Flag-tagged proteins, and Western blotting results suggested that the association between IKK α and IRF7 was enhanced in the presence of LC3 (Fig. 4B). Thus, these data suggest that LC3 recruits IKK α and that this LC3-IKK α complex may further assemble a larger LAP-dependent signalosome (LAPosome) involving TRAF3 and IRF7.

Interactions between IKK α and LC3 depend on the LC3-interacting regions in IKK α

The LC3-interacting region (LIR), which is defined by a conserved hydrophobic motif of WXXL, where X is any amino acid, is used by

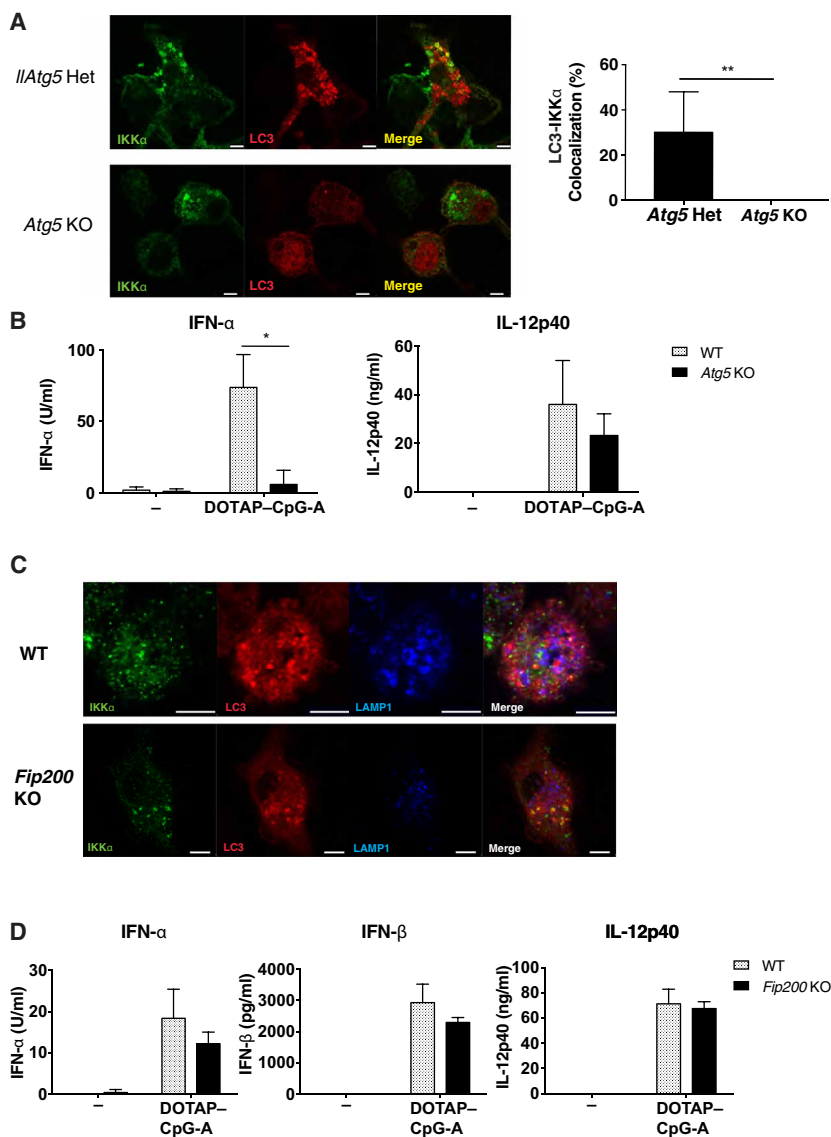


Fig. 3. LC3- $IKK\alpha$ interaction depends on LAP but not canonical autophagy. (A) *Atg5*^{+/−} or *Atg5*^{−/−} fetal liver HSCs were retrovirally transduced with WT $IKK\alpha$ -HA and Cherry-LC3, differentiated into macrophages, and stimulated with 1 μ M DOTAP-CpG-A for 6 hours. After stimulation, cells were stained for the indicated proteins and analyzed by confocal microscopy. Images are representative of three or more independent experiments. Scale bars, 5 μ m. ** $P < 0.01$. het, heterozygous. (B) WT or *Atg5*^{−/−} fetal liver macrophages were stimulated with DOTAP-CpG-A for 24 hours, and secretion of IFN- α and IL-12 was measured by enzyme-linked immunosorbent assay (ELISA). Data are means \pm SEM from three to five independent experiments. * $P < 0.05$ by *t* test. (C) WT and *Fip200*^{f/f} \times *LysM-Cre* HSCs were retrovirally transduced with Cherry-LC3, differentiated to BMDCs, and stimulated with DOTAP-CpG-A for 6 hours. After treatment, cells were stained and analyzed by confocal microscopy. Images are representative of three or more independent experiments. Scale bars, 5 μ m. (D) WT and *Fip200*^{f/f} \times *LysM-Cre* BMDCs were stimulated with 1 μ M DOTAP-CpG-A for 24 hours, and the cytokine accumulation in the supernatants was assayed by ELISA. Data are means \pm SEM from three or more independent experiments.

various proteins to interact with LC3 (20, 21). Sequence analysis of $IKK\alpha$ uncovered three putative LIRs, which we designated as LIR-1 (residues 275 to 278), LIR-2 (residues 651 to 654), and LIR-3 (residues 740 to 743) (fig. S3A). We mutated each LIR sequence by replacing the tryptophan residue in the motif with an alanine residue to examine their role in LC3- $IKK\alpha$ complex formation. After

co-transfection of HEK293T cells with plasmids encoding each individual LIR-3 mutant $IKK\alpha$ -HA and LC3-GFP, Western blotting analysis showed an increase in the abundance of $IKK\alpha$ protein (Fig. 5A). These data suggest that mutation of the LIR-3 sequence of $IKK\alpha$ may have affected protein stability. Nevertheless, $IKK\alpha$ -HA was immunoprecipitated from all cell lysates in equivalent abundance. Western blotting analysis of immunoprecipitates of LC3 showed that mutation of the LIR-2 and LIR-3 sequences but not the LIR-1 sequence abolished the interaction between $IKK\alpha$ and LC3. These data suggested that LIR-2 and LIR-3 might be involved in the binding of $IKK\alpha$ to LC3 (Fig. 5A). We confirmed this observation by confocal microscopy, which showed that WT $IKK\alpha$ and the LIR-1 mutant $IKK\alpha$, but not LIR-2 and LIR-3 mutant $IKK\alpha$, colocalized with LC3 (Fig. 5B). Together, these results suggest that LIRs near the C terminus of $IKK\alpha$ are required for binding to LC3 and recruitment to the LAPosomes after TLR9 stimulation.

Mutations in the LIR motifs do not affect the kinase activity of $IKK\alpha$

The kinase activity of $IKK\alpha$ is required for noncanonical NF- κ B signaling by mediating the ubiquitin-dependent degradation of NF- κ B p100 and its processing into p52 (22). A subset of TNF receptor superfamily members, such as lymphotoxin β receptor (LT β R), B cell-activating factor of the TNF family (BAFF), and CD40, induce noncanonical NF- κ B activation (23). To examine the possibility that LIR mutations interfered with the kinase activity of $IKK\alpha$, we transduced $IKK\alpha$ -deficient mouse embryonic fibroblasts (MEFs) with retroviruses expressing WT $IKK\alpha$ or $IKK\alpha$ -LIR mutants and analyzed p52 nuclear translocation after LT β R stimulation. The mutant $IKK\alpha$ constructs were expressed; however, more $IKK\alpha$ protein accumulated in cells expressing the LIR-3 mutant than in cells expressing WT $IKK\alpha$ (fig. S3). Agonistic antibody cross-linking of LT β R induced p52 nuclear translocation in MEFs expressing either WT or LIR mutant $IKK\alpha$ but not in control $IKK\alpha$ -deficient MEFs (fig. S3B). These data suggest that LIR mutant $IKK\alpha$ constructs retain kinase activity.

Interaction between $IKK\alpha$ and LC3 is required for type I IFN production

Finally, we sought to identify the physiological importance of the interaction between $IKK\alpha$ and LC3. $IKK\alpha$ is specifically required for type I IFN production in response to TLR9 activation because $IKK\alpha$ deficiency in both pDCs and conventional dendritic cells results in impaired IFN production but intact IL-12p40 production (6, 24). Therefore, we reconstituted *Ikka*^{−/−} BMDCs with WT or LIR mutant $IKK\alpha$ to examine whether LIR sequence mutations had any effect on type I IFN production after TLR9 stimulation. Consistent with previous results (6), *Ikka*^{−/−} pDCs showed significantly reduced IFN- α production upon TLR9 stimulation but

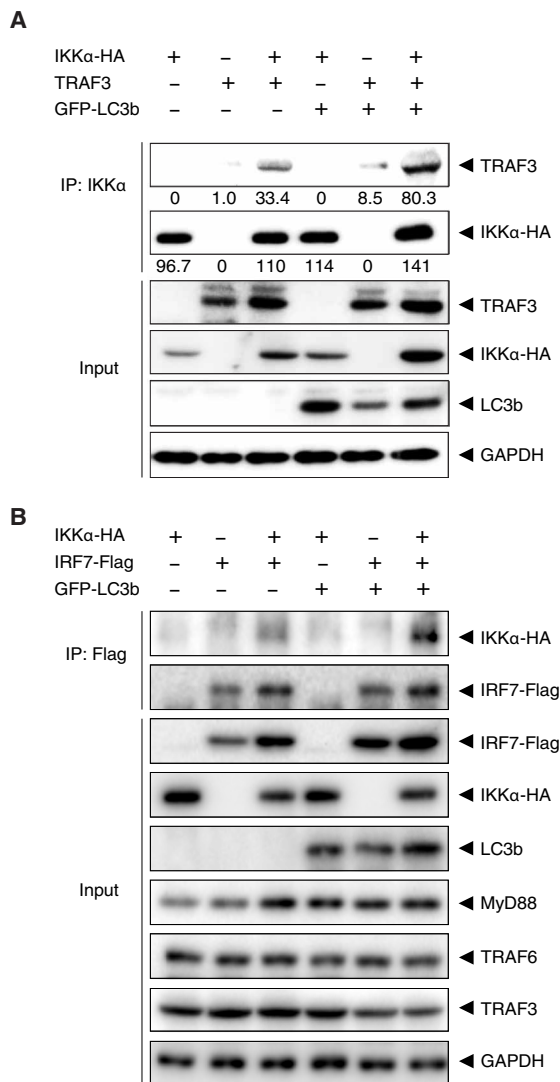


Fig. 4. IKK α -LC3 complex recruits TRAF3 and IRF7. (A) HEK293T cells were transfected with plasmids encoding WT IKK α -HA, GFP-LC3, and TRAF3. Cell lysates were immunoprecipitated for IKK α and blotted for the indicated proteins. Blots are representative of three independent experiments. GAPDH, glyceraldehyde-3-phosphate dehydrogenase. (B) HEK293T cells were transfected with plasmids encoding WT IKK α -HA, GFP-LC3b, MyD88, TRAF3, TRAF6, and IRF7-Flag. Cell lysates were subjected to immunoprecipitated for the Flag-tag and blotted for the indicated proteins. Blots are representative of three independent experiments.

had no defect in IL-12p40 production (fig. S4). Similarly, when BMDCs from *Ikka*^{-/-} fetal liver chimeras were stimulated through TLR9, they produced substantially less IFN- β than did WT cells but secreted normal amounts of IL-12p40 (Fig. 6). Upon reconstitution of *Ikka*^{-/-} BMDCs with WT or mutant IKK α constructs, only those cells expressing WT or LIR-1 mutant IKK α exhibited restored IFN- β production (Fig. 6). Furthermore, in HEK293T cells overexpressing LC3, TRAF3, TRAF6, MyD88, and IRF7, only those expressing either WT IKK α or the LIR-1 mutant IKK α , but not LIR-2 or LIR-3 mutant IKK α , exhibited phosphorylation of IRF7 (fig. S5). Computational modeling using SWISS-MODEL suggests that LIR-2

could be located on the surface of the protein, whereas it appears that LIR-1 may be buried close to the kinase core (fig. S6). Thus, the LIR-2 domain but not the LIR-1 domain of IKK α may be accessible for binding to LC3. Together, our results suggest that IKK α recruitment and interaction with LC3 and TLR9 on LAPosomes are critical for stimulating type I IFN production.

DISCUSSION

Here, we provide evidence for a link between two previously identified pathways required for the TLR9 signaling that stimulates type I IFN production. We showed that shortly after stimulation of macrophage and dendritic cells with CpG-A, IKK α was recruited to the LC3 embedded in TLR9-positive endosomes. This process did not occur after the induction of canonical autophagy. In addition, macrophages that lacked FIP200, a critical component of autophagosome formation, were still capable of exhibiting IKK α recruitment to LC3-positive endosomes and signaling for type I IFN production downstream of TLR9. These data suggest that noncanonical autophagy, namely, LAP, is important for the IKK α recruitment and TLR9 signaling. To gain a better understanding of the IKK α -LC3 interaction, we identified and examined three putative LIRs in IKK α . Alanine replacement of the critical tryptophan motif of the LIR (WXXL) suggested that mutations in the LIR-2 and LIR-3 domains reduced the binding of IKK α to LC3. Furthermore, reconstitution of *Ikka*^{-/-} BMDCs with these mutants failed to restore type I IFN production. In contrast, WT IKK α and the LIR-1 mutant IKK α bound to LC3 and restored the TLR signaling required for type I IFN production in *Ikka*^{-/-} BMDCs. Collectively, our study highlights a link between two known pathways for IFN induction by TLR9, namely, LAP and IKK α .

Although this type of experiment cannot rule out the possibility that a loss of binding may be secondary to protein structural distortion, the IKK α mutants exhibited no defects in the activation of non-canonical NF- κ B signaling (fig. S3B). Specifically, all of our IKK α mutants were capable of p52 activation downstream of LT β R signaling mediated by NF- κ B-inducing kinase (NIK) (fig. S3B). Because NIK is required for the phosphorylation of IKK α (25), these data suggest that none of the LIR mutations that we generated interfered with phosphorylation by NIK. However, we did not directly examine this possibility in our current study.

IKK α is required for canonical autophagy induced by starvation or inflammatory signals. IKK α and IKK β are phosphorylated upon starvation, and cells deficient in these genes have impaired autophagy induction (26, 27). Downstream of TNF receptor signaling, activated IKK α also phosphorylates and stabilizes Atg16L1 and prevents its degradation (28). In our current study, canonical autophagy induced by starvation or Tat-Beclin-1 peptide treatment did not result in IKK α recruitment to the LC3-coated LAPosomes (Fig. 2), suggesting that IKK α phosphorylation induced by these signals is not sufficient for IKK α -LC3 interaction. In addition, IKK α -deficient BMDC reconstituted with LIR-2 and LIR-3 mutants had only slightly fewer LC3 puncta per cell (Fig. 5B). Thus, it is possible that IKK α also participates in LAP formation and that these mutations may have interfered with that process. Nevertheless, there was little association of IKK α LIR mutants with LC3 puncta (Fig. 5C), which suggests that these LIR regions in IKK α control their recruitment to LAPosomes.

On the basis of our current study, we propose a revised model for TLR9 signaling (Fig. 6). After stimulation, ligand-bound TLR9

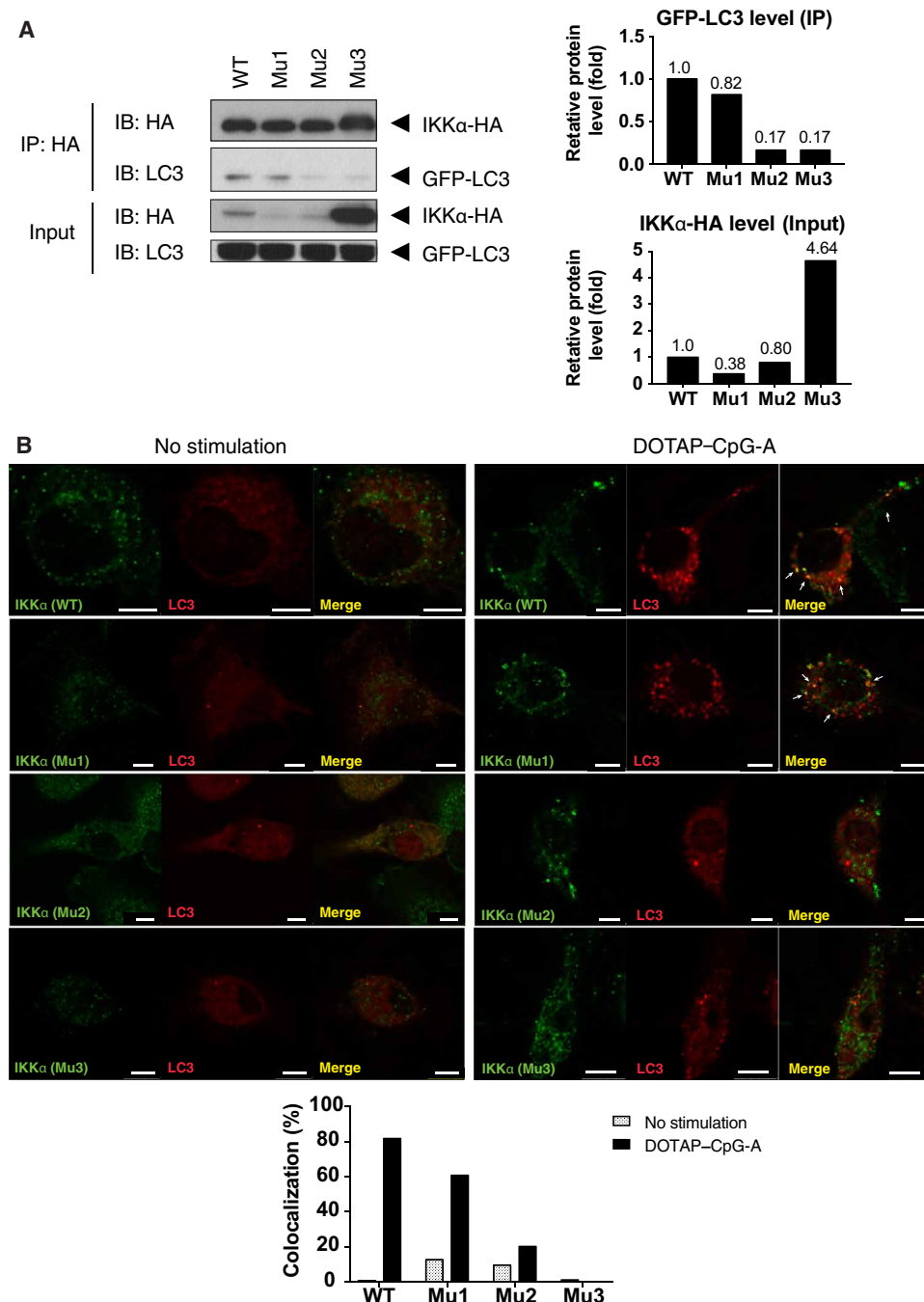


Fig. 5. Interaction between IKK α and LC3 depends on the LC3-interacting region in IKK α . (A) HEK293T cells were transfected with different LIR mutants, immunoprecipitated for LC3, and blotted for the indicated proteins. Blots are representative of three or more independent experiments. Mu1, LIR-1 mutant; Mu2, LIR-2 mutant; Mu3, LIR-3 mutant. (B) IKK α -deficient bone marrow cells were retrovirally reconstituted with IKK α -HA (WT, LIR-1, LIR-2, or LIR-3 mutant) and Cherry-LC3, differentiated into BMDCs, and stimulated with 0.5 μ M DOTAP-CpG-A for 6 hours. After treatment, cells were analyzed by confocal microscopy. Images are representative of three independent experiments. Scale bars, 5 μ m.

dimerizes and recruits the signaling adaptor MyD88 and downstream signaling molecules, including IRAK4 and the E3 ubiquitin ligase TRAF6. Autoubiquitylation of TRAF6 mediates recruitment of the kinase TAK1 and its regulatory proteins TAB2 and TAB3, which

phosphorylates and activates IKK β . This induces the NF- κ B-mediated transcription of proinflammatory genes (NF- κ B endosome; Fig. 7). Activation of TLR9 also stimulates the ATG5-dependent recruitment of LC3 to the endosomal membrane through the LAP process (12). Our data suggest that IKK α is recruited to the LC3 through the LIR-2 and LIR-3 domains. This likely brings IKK α into proximity with IRF7, which it may phosphorylate to activate the transcription of genes encoding type I IFNs.

These data provide a link between IKK α and LAP for type I IFN production. We suggest that the direct recruitment of IKK α is LAP-dependent because the recruitment was lost in *Atg5*^{-/-} BMMs but not in *Fip200*^{-/-} BMMs. Our data suggest that LC3 acts as an anchor for IKK α on TLR9-positive endosomes and is necessary for the production of type I IFN (Fig. 7). Although ATG8 family proteins (including LC3) recruit various cargoes into autophagosomes by interacting with adaptor proteins (21), our work suggests that IKK α may be directly recruited to LAPosomes through its interaction with LC3. In addition, because LC3 decorates the outer membrane of TLR9-containing endosomes, this enables IKK α to be recruited to a signaling platform on the cytosolic face of the membrane. This suggests that LC3 may also mediate the recruitment of other molecules to the outer membrane of autophagosomes or the LAPosome for various cellular processes.

Second, our results contribute to the expanding list of potential roles of LAP in regulating molecular and cellular processes. The central component of LAP is LC3, which interacts with various proteins through its LIRs. The role of LIRs in the selective recruitment of autophagic cargoes has been extensively studied because LIRs provide docking sites for adaptor proteins, such as p62, a ubiquitin-binding protein, neighbor of BRCA1 gene 1, nuclear domain 10 protein 52, Nip-like protein X, and ATG19 (21). A proteomics analysis showed that the ATG8 family, which includes the LC3 and gamma-aminobutyric acid receptor-associated protein (GABARAP) subfamilies, has 67 high-confidence interactions with other cellular proteins (29). In addition, proteomic analysis identified Rubicon and UVRAG-containing class III PI3K complex proteins as being associated with the TLR2-induced LAPosome (14). Our study identifies IKK α as another LC3-binding, LAPosome-associated protein

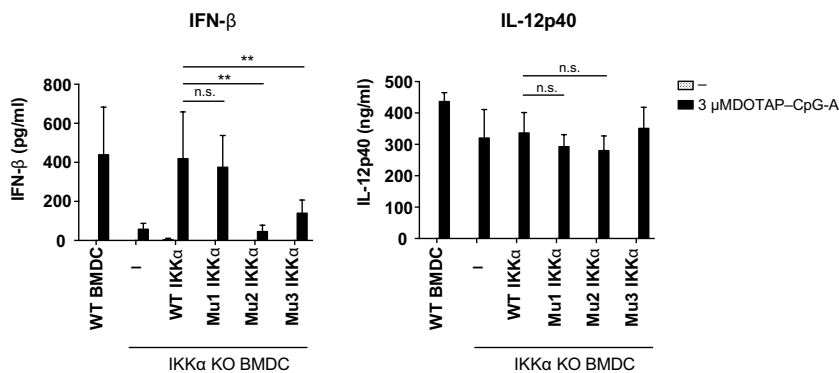


Fig. 6. Interaction between LC3 and IKK α is crucial for type I IFN production in granulocyte-macrophage colony-stimulating factor–differentiated BMDCs. *Ikk α* ^{-/-} bone marrow cells were retrovirally reconstituted with IKK α -HA (WT, LIR-1, LIR-2, or LIR-3 mutant), differentiated into BMDCs, and stimulated with DOTAP–CpG-A. After 24 hours, the supernatants were harvested, and cytokine production was assayed by ELISA. Data are means \pm SEM from three independent experiments. ***P* < 0.01 by *t* test. n.s., not significant.

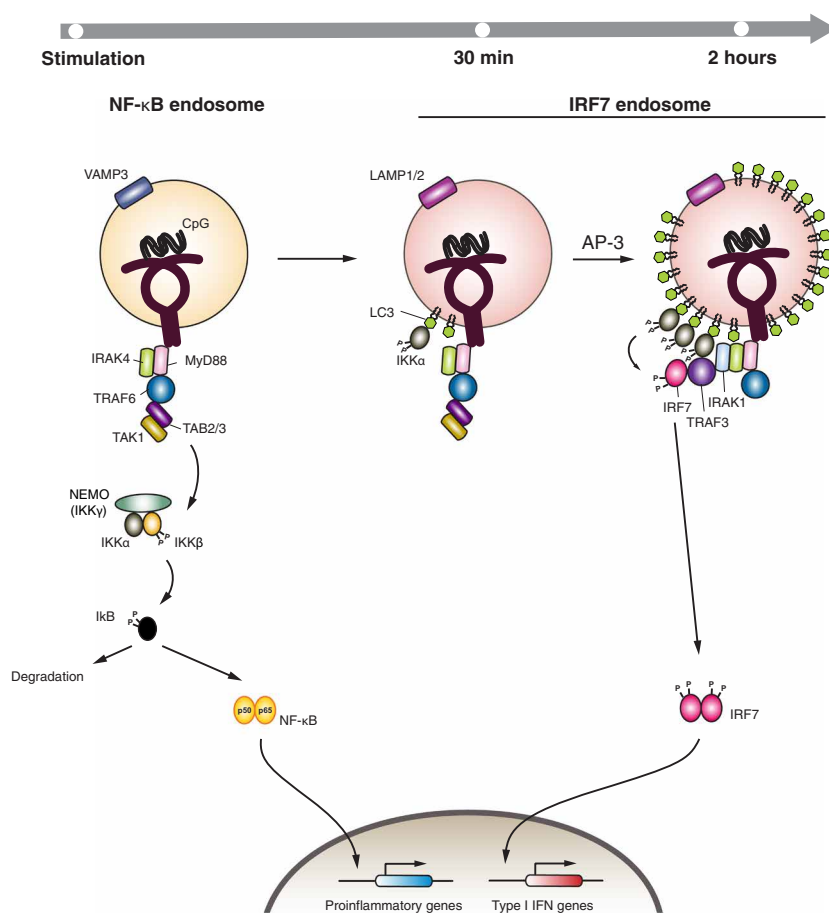


Fig. 7. Schematic of IKK α complex recruitment to LC3-containing LAPosomes necessary for TLR9 signaling and IFN production. After stimulation, ligand-bound TLR9 dimerizes and recruits signaling adaptor MyD88 and downstream signaling molecules including IRAK4 and the E3 ligase TRAF6. Autoubiquitination of TRAF6 mediates recruitment of TAK1 and its regulatory proteins TAB2/3, which phosphorylates and activates IKK β . This induces NF- κ B–mediated transcription of proinflammatory genes (NF- κ B endosome). Activation of TLR9 also triggers ATG5-dependent recruitment of LC3 to the endosomal membrane through the process of LAP. IKK α is recruited to the LC3 through LIR-2 and LIR-3 domains. Adaptor protein 3 (AP-3) is required for the formation of lysosome-related organelle decorated with LAMP2. IKK α recruitment to the membrane-bound LC3 brings this kinase to the proximity of IRF7, where it phosphorylates IRF7 and activates type I IFN gene transcription. I κ B, NF- κ B inhibitor β .

and suggests that the LIR-LC3 interaction may be involved in kinase recruitment to the signaling platform on the endosomal membrane.

Our results suggest that both LIR-2 and LIR-3 may be important for the ability of IKK α to interact with LC3 after TLR9 stimulation. Although the crystal structure of IKK α has not been solved yet, its likely structure can be extrapolated from that of IKK β due to the similarity between the two molecules (52% identical in amino acid sequence) (30). Our prediction of the IKK α structure using SWISS-MODEL suggests that LIR-2 could be located near the surface of the protein, which has higher accessibility for other protein interactions (fig. S6). Unfortunately, SWISS-MODEL could not predict the likely structure of LIR-3 because no corresponding region exists in IKK β . Whereas yeast and other fungi have only one ATG8-encoding gene, humans have multiple ATG8 subfamily genes, including a single GATE-16 gene, two GABARAP genes, and four LC3 genes with splice variants (31). Whereas all ATG8 subfamily proteins are recruited to autophagosomes, it remains unknown if they play redundant roles in mediating TLR9 signaling. It is possible that only a subset of ATG8 proteins form specific interactions with IKK α .

In autoimmune disease, recognition of self-RNA and DNA by a variety of PRRs may trigger excessive amounts of type I IFN to be produced. The activation of TLR7 and TLR9 is linked to the exacerbation of autoimmune diseases such as systemic lupus erythematosus (SLE) (32) and psoriasis (33). Similarly, a universal deficiency in LAP components impaired clearance of dying cells and exacerbated SLE pathogenesis in mice (34). On the basis of our current study, inhibiting LAPosome signaling in pDCs may ameliorate SLE symptoms by reducing the amount of type I IFN produced without affecting phagocytic clearance. Specific inhibition of IKK α -LC3 interaction may provide a selective means to reduce the amount of type I IFN produced by pDCs in patients with SLE or psoriasis, while leaving the LAP-dependent phagosomal degradation intact. Further investigation is still needed to determine the cell type differences in LAP usage as a potential target of intervention in SLE pathogenesis.

MATERIALS AND METHODS**Mice**

Age- and sex-matched C57BL/6 (wild type) mice from the National Cancer Institute were used for isolating bone marrow. CD45.1 congenic C57BL/6 mice were purchased from Charles River for fetal liver chimera generation. All procedures used in this study complied with federal guidelines and the institutional policies of the Yale Animal Care and Use Committee. *Ikkα*^{-/-} mice were a gift from M. Karin at University of California, San Diego. *Atg5*^{+/-} mice (35) were bred in the Yale University Animal Facility.

Fetal liver chimera generation

Ikkα and *Atg5* fetal liver chimeric mice were generated by standard protocols, as described previously (19). Briefly, 2×10^6 to 3×10^6 liver cells from embryonic day 15.5 (E15.5) to E18.5 embryos born to *Ikkα*^{+/-} or *Atg5*^{+/-} parents were screened for the *Ikkα*^{-/-} or *Atg5*^{-/-} genotype and injected into recipient C57BL/6 mice through the tail vein. Before the transfer of liver cells, the recipient mice were lethally irradiated with 4.75 Gy (gray) of irradiation twice, 3 hours apart. This procedure led to complete chimerism of hematopoietic cells by 8 weeks after transplantation.

Cell lines and primary cell culture

RAW267.4 cells were purchased from the American Type Culture Collection and cultured in RPMI supplemented with 10% fetal bovine serum, penicillin/streptomycin, 10 mM Hepes (Invitrogen), and 2-mercaptoethanol (fetal bovine serum). TLR9-HA- and Cherry-LC3b-coexpressing cells were prepared by transduction with MSCV2.2 retrovirus, and the cells were selected with puromycin (2 μg/ml). Bone marrow-derived pDCs (36), BMDCs (37), and BMM (11) were prepared as previously described. HSCs were transduced with retroviruses expressing the gene of interest at day 0 and day 1 by spinoculation at 1500g for 2 hours. After spinoculation, viral medium was removed, and cells were resuspended with fresh medium to be plated for further cell differentiation. Cell culture medium was replaced at day 3 and day 5 throughout the 7-day BMM and BMDC differentiation. Bones from *Fip200*^{fl/fl} × *LysM-Cre* mice were a gift from H. W. Virgin (Washington University at St. Louis). *IKKα*-deficient MEFs were generated from E14.5 fetuses as previously described (38).

Reagents and antibodies

LPS was purchased from List Biological Laboratories, CpG 2216 (CpG-A) was purchased from TriLink BioTechnologies, and both polyinosinic:polycytidylic and R848 were purchased from Invivogen. DOTAP was obtained from Roche. The Tat-Beclin-1 peptide was purchased from Sigma-Aldrich. Mouse anti-HA tag (H3663), mouse anti-Flag-tag M2 (F1804), and mouse anti-β-actin (A1978) antibodies were from Sigma-Aldrich. Mouse anti-glyceraldehyde-3-phosphate dehydrogenase (GTX627408) antibody was from GeneTex. Rabbit anti-LC3b (ab48394), rabbit anti-Myd88 (ab2064), and mouse anti-GFP (ab1218) were from Abcam. Rabbit anti-*IKKα* (2682), rabbit anti-phosphorylated *IKKα/β* (2697), rabbit anti-p100/p52 (4882), rabbit anti-TRAF3 (4729), rabbit anti-TRAF6 (8028), and rabbit anti-Flag-tag horseradish peroxidase (HRP) (2044) antibodies were obtained from Cell Signaling Technology. HRP-conjugated goat antibody against mouse immunoglobulin G (IgG) and HRP-conjugated antibody against rabbit IgG were purchased from Jackson ImmunoResearch and Cell Signaling Technology, respectively. TrueBlot ULTRA HRP anti-mouse (13-8817-80) and TrueBlot HRP anti-rabbit (18-8816-31) antibodies

were obtained from Rockland. For flow cytometry, anti-phosphorylated IRF7 antibody (558630) was from BD Biosciences. Agonistic LTβR antibody (5G11) was purchased from Abcam.

Plasmids

TRAF3-HA was constructed as described previously (11). TLR9-HA and the MyD88 plasmids were a gift from R. Medzhitov. Plasmid encoding GFP-LC3b was a gift from N. Mizushima. Plasmid encoding the mIRF7-Flag was a gift from K. Hoshino and T. Kaisho. The plasmids pcDNA3-*IKKα*-HA (#23296), pMIG-II (#52107), TRAF6 (#21624), and c-Flag pcDNA3 (#20011) were purchased from Addgene. Human IRF7 plasmid (LV798857) was purchased from Applied Biological Materials Inc. The ATG5-Flag was constructed by subcloning the gene from our previously generated ATG5-IRES-EGFP into the c-Flag pcDNA3 vector. Cherry-LC3b and the mutant *IKKα* (mutants 1 to 3, AA mutant) were subcloned by standard polymerase chain reaction protocols. Retrovirus vectors containing TLR9-HA and Cherry-LC3b were generated by subcloning the genes into pMSCV2.2 and pMSCVpuro, respectively. Retrovirus vectors containing mutant *IKKα* were generated by subcloning the genes into the pMIG-II vector.

Immunoprecipitation and Western blotting

Cells were lysed with lysis buffer containing 20 mM tris-HCl (pH 7.5), 1% NP-40, 150 mM NaCl, proteinase inhibitor cocktail (Roche), and phosphatase inhibitor (PhosSTOP tablets, Roche) for 30 min (HEK293T cells) or overnight (RAW264.7 and BMMs). Lysates were harvested and centrifuged at 21,130g for 10 min to obtain a postnuclear supernatant. The supernatant was then subjected to immunoprecipitation assay with antibodies and Dynabeads Protein G (Thermo Fisher Scientific). Beads were then washed with wash buffers, followed by Western blotting analysis, which was performed as previously described (11).

Immunofluorescence microscopy

Transduced RAW264.7 and BMMs were stimulated on day 7 of differentiation with 3 μM DOTAP-CpG-A or 1 μM Tat-Beclin-1 peptide for the times indicated in the figure legends and then stained for immunofluorescence microscopy as previously described (39). The following antibodies were used for immunofluorescence microscopy analysis: mouse anti-*IKKα* (from Santa Cruz Biotechnology); rabbit anti-HA, mouse anti-GFP, and rabbit anti-LAMP1 (from Abcam); rabbit anti-VAMP3 (from Synaptic Systems); rat anti-red fluorescent protein (5F8) (from ChromoTek); Alexa Fluor 488 anti-mouse antibody (Invitrogen); and biotin anti-mouse, streptavidin-A488, streptavidin-Cy5, Cy3 anti-rabbit, Cy3 anti-rat, Cy5 anti-rabbit, and A647 anti-rabbit antibodies (from Jackson ImmunoResearch). After staining, coverslips were mounted on slides with ProLong Gold (Invitrogen) and imaged with a Leica TCS SP8 confocal microscope using a 64× objective lens. A minimum of 15 to 20 cells were examined per slide.

Enzyme-linked immunosorbent assay

The amounts of IFN-α protein present in the cell culture medium and serum were measured by ELISA as previously described (40). The amounts of IFN-β protein present in the cell culture medium and serum were measured with a VeriKine mouse IFN-β ELISA (PBL Assay Science). The amounts of IL-12p40 protein present in the cell culture medium and serum were measured according to the manufacturer's instructions with specific antibodies (BioLegend).

Flow cytometry

HEK293T cells were transfected with plasmids encoding WT or mutant IKK α constructs, human IRF7, Myd88, TRAF3, TRAF6, and GFP-LC3b. The cells were then fixed, permeabilized in 200 μ l of BD Cytotfix/Cytoperm (BD Biosciences) for 20 min at 4°C, and stained with antibodies Alexa Fluor 647-conjugated anti-pIRF7 antibody (pS477/pS479, BD Biosciences) for 30 min in BD Perm/Wash buffer at 4°C. Mean fluorescent intensity of phosphorylated IRF7 in each sample was analyzed by FlowJo (Tree Star).

Statistical analysis

Normally distributed continuous variable comparisons were performed with the Student's *t* test using Prism software (GraphPad). Data are presented as means \pm SEM. *P* < 0.05 was considered statistically significant.

SUPPLEMENTARY MATERIALS

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 Fig. S1. LC3 binds to TLR9 and IKK α after DOTAP-CpG-A stimulation.
 Fig. S2. ATG5 does not interact with LC3 or IKK α .
 Fig. S3. LIR mutants of IKK α maintain intact kinase activity.
 Fig. S4. Fms-related tyrosine kinase 3 ligand-derived IKK α -deficient pDCs show impaired IFN- α production but intact IL-12p40 after CpG-A stimulation.
 Fig. S5. WT and LIR-1 but not LIR-2 or LIR-3 mutant IKK α induces IRF7 phosphorylation.
 Fig. S6. LIR prediction based on IKK β structure by SWISS-MODEL.

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The interaction between IKK α and LC3 promotes type I interferon production through the TLR9-containing LAPosome

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LC3 recruits IKK α after TLR9 activation

Toll-like receptor 9 (TLR9) is a pattern recognition receptor involved in innate sensing of extracellular DNA within the endosomes of phagocytes. Hayashi *et al.* found that IKK α kinase associated with the autophagy protein LC3 deposited on endosomes after TLR9 stimulation of macrophages. Formation of this LC3-IKK α signaling platform through noncanonical autophagy also recruited other effectors involved in type I interferon signaling. Mutagenesis of LC3-interacting regions in IKK α suggested that these effects were mediated by a direct interaction of the kinase with LC3. Together, these data identify IKK α as a critical component of LC3-associated phagocytosis-mediated endosome-based signaling platforms necessary for stimulating type I interferon production after TLR9 engagement.

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