Proinflammatory NF-κB activation requires the IκB (inhibitor of NF-κB) kinase (IKK) complex that contains two catalytic subunits named IKKα and IKKβ and a regulatory subunit named NF-κB essential modulator (NEMO). NEMO and IKKβ are essential for tumor necrosis factor (TNF)-induced NF-κB activation, and we recently demonstrated that NEMO and IKKα are sufficient for interleukin (IL)-1-induced signaling. IKKα and IKKβ both contain a functional NEMO-binding domain (NBD); however, the role of NEMO association with each kinase in NF-κB signaling and IKK complex formation remains unclear. To address this question, we stably reconstituted IKKα−/− and IKKβ−/− murine embryonic fibroblasts (MEFs) with wild-type (WT) or NBD-deficient (ΔNBD) versions of IKKα and IKKβ, respectively. TNF-induced classical NF-κB activation in IKKβ−/− MEFs was rescued by IKKβWT but not IKKβΔNBD, whereas neither IKKαWT nor IKKαΔNBD affected IL-1-induced NF-κB signaling. As previously described, classical NF-κB transcriptional activity was absent in IKKα−/− cells. Reconstitution with either IKKαWT or IKKαΔNBD rescued both IL-1 and TNF-induced transcription, demonstrating that NEMO association is not required for IKKα-dependent regulation of NF-κB-dependent transcription. Stably expressed IKKαWT or IKKβWT associated with endogenous IKKs and NEMO in IKKα−/− or IKKβ−/− MEFs, respectively, resulting in formation of the heterotrimetric IKKα-IKKβ-NEMO complex. In contrast, although the IKKαΔNBD and IKKβΔNBD mutants associated with endogenous IKKs containing an NBD, these dimeric endogenous IKK-IKKβ-NEMO complexes did not associate with NEMO. These findings therefore demonstrate that formation of the heterotrimetric IKKα-IKKβ-NEMO holocomplex absolutely requires two intact NEMO-binding domains.

NF-κB² describes a family of transcription factors that regulate the inducible expression of many genes essential for innate and adaptive immunity, inflammation, and cell survival. A wide range of stimuli activates NF-κB, including ligation of innate immune receptors (e.g., TLRs), antigen receptor engagement (B cell receptor and T cell receptor), and proinflammatory cytokines (e.g., IL-1 and TNF) (1). NF-κB activation by these stimuli is normally rapid and transient; however, constitutive NF-κB activity occurs in some chronic inflammatory diseases, solid tumors, leukemias, and lymphomas (1, 2). Understanding the molecular mechanisms that regulate NF-κB activity will therefore reveal novel targets for blocking pathophysiological NF-κB signaling (3–5).

The five NF-κB family members are NF-κB1/p105 and NF-κB2/p100 that are processed to generate p50 and p52, respectively, p65 (RelA), c-Rel, and RelB (1). These proteins homo- or heterodimerize to form transcriptionally active (e.g., p50-p65) or repressive (e.g., p50-p50) dimers that are retained in the cytosol of resting cells by members of the inhibitory family of IκB proteins. The IκBs include IκBα, IκBβ, IκBε, and the C termini of p105 and p100. The prototypic NF-κB-IκB complex expressed in most cell types is a heterodimer of p50 and p65 associated with IκBα (1). Following cell stimulation, the IκBs are rapidly phosphorylated, ubiquitinated, and then degraded by the 26 S proteasome. Free NF-κB dimers then migrate to the nucleus, where they bind target gene promoters and regulate transcription (1).

The IκB proteins are phosphorylated by the high molecular weight heterotrimeric IκB (inhibitor of NF-κB) kinase (IKK) complex (1, 6, 7). The IKK complex contains two kinases named IKKα (IKK1) and IKKβ (IKK2) and a noncatalytic subunit named NEMO (NF-κB essential modulator) or IKKγ (1, 6, 7). NEMO is critical for proinflammatory IKK activation (8–12), and we previously identified a domain within the C termini of both IKKα and IKKβ that facilitates their association with NEMO (13, 14). A cell-permeable peptide spanning this NEMO-binding domain (NBD) disrupts the IKK complex and blocks proinflammatory NF-κB activation, confirming the crucial role of NEMO association for IKK complex activation (13).

Despite their significant structural similarities, genetic analyses of IKKα and IKKβ revealed distinct roles for the kinases during NF-κB activation (1, 6, 7, 15). In this regard, TNF-induced IκBα degradation is dependent upon IKKβ and also requires NEMO (8, 16, 17). This mechanism is termed the classical NF-κB pathway and is defined as NEMO- and IKKβ-dependent IκB phosphorylation and degradation releasing canonical NF-κB complexes typified by the ubiquitous p50-p65 heterodimer. IKKα plays a separate role in mediating NIK (NF-κB-inducing kinase)-dependent processing of NF-κB2/p100 to generate p52 (18–22). This mechanism is activated in the absence of both IKKβ and NEMO and is named the noncanoni-

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classical NF-κB pathway (15). Ligation of only a subset of receptors, including the lymphotxin-β receptor, CD40, and BAFF-R, activates the IKKa-dependent noncanonical pathway, and the resulting p52, together with RelB, regulates a panel of chemokine and cytokine genes required for lymphoid organogenesis and B-cell maturation (18–23). IKKa has also been shown to play several roles in regulating the transcriptional activity of classical NF-κB that are separate from its upstream signaling function as an IκB kinase (24–29).

Since TNF-induced IκBα degradation and classical NF-κB nuclear translocation occurs in the absence of IKKa, a model of classical pathway activation has emerged in which IKKa is redundant (15). It remains, however, that IKKa associates via its NBD with NEMO (14), and we previously questioned whether this association plays a functional role in classical NF-κB signaling (30). Surprisingly, we found that although TNF-induced IκBα degradation requires NEMO and IKkB, IL-1-induced classical pathway activation is intact in cells lacking IKkB. Furthermore, IL-1-induced NF-κB activation in IKkB-deficient cells was blocked by the NBD peptide, demonstrating that a complex of only IKKa and NEMO is sufficient for IL-1- but not TNF-induced classical NF-κB activation (30). Intriguingly, Lam et al. (31) recently demonstrated that IKKa plays a crucial compensatory role in regulating constitutive classical NF-κB pathway activation in a subset of diffuse large B-cell lymphoma cells in which IKkB has been pharmacologically inhibited. These findings therefore identify differences in the absolute requirements for the separate IKK subunits activated in a NEMO-dependent manner by distinct stimuli and suggest that targeting only IKkB may not effectively block dysregulated classical NF-κB activation. Consequently, determining the role of the interaction of each IKK subunit with NEMO will provide novel insight into the mechanisms that regulate NEMO-dependent classical NF-κB activation.

To address this question, we determined the effects of individually deleting the NBD in IKKa and IKkB on classical NF-κB signaling and IKK complex formation. We stably reconstituted IKKaΔNBD and IKkBΔNBD murine embryonic fibroblasts (MEFs) with wild-type (WT) or NBD-deficient (ΔNBD) versions of IKKa and IKkB, respectively. Reconstitution of IKKaΔNBD/IKkBWT with IKKBWT but not IKKBΔNBD rescued TNF-induced classical NF-κB activation, confirming the requirement for the NEMO-IKKβ association for TNF signaling. IL-1 signaling was intact in IKKaΔNBD cells, and this was not affected by expression of either IKkBWT or IKKBΔNBD. Classical NF-κB transcriptional activity was absent in IKKaΔNBD cells, and reconstitution with either IKKaWT or IKKAΔNBD rescued both IL-1- and TNF-induced transcription. This therefore demonstrates that association with NEMO is not necessary for IKKα-dependent regulation of NF-κB-dependent transcriptional activity. Immunoprecipitation analysis and size exclusion chromatography revealed that stably expressed IKKAWT or IKKAΔNBD associated with endogenous IKKα and NEMO in IKKaΔNBD or IKKBΔNBD MEFs, respectively, and this association resulted in the formation of the heterotrimeric IKKα-IKKβ-NEMO complex. In contrast, despite the ability of the IKKαΔNBD and IKKβΔNBD mutants to associate with endogenous IKKαs containing an NBD, these mutants formed dimeric endogenous IKKβΔNBD complexes that did not associate with NEMO. These findings therefore demonstrate that formation of the heterotrimeric IKKα-IKKβ-NEMO holocomplex absolutely requires the presence of two intact NEMO-binding domains.

**EXPERIMENTAL PROCEDURES**

**Reagents and Cell Culture**—Recombinant human IL-1α was obtained from Peprotech (Rocky Hill, NJ). Recombinant human TNF and recombinant mouse LTαβ2 were from R&D Systems (Minneapolis, MN). Polyclonal rabbit anti-IKKα (sc-7218), rabbit anti-NEMO (sc-8330), goat anti-NEMO (sc-8256), rabbit anti-p100/p52 (sc-298), and rabbit anti-IKKβ (sc-8014) antisera were from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA). Monoclonal anti-α-tubulin (T5168) was from Sigma. Anti-phospho-p65 (Ser536) (catalog number 2694), anti-phospho-IKKαβ (Ser176/180) (catalog number 3033), anti-phospho-IKKα/β (Ser176/180) (catalog number 2694), and anti-histone 3 (catalog number 9715) were from Cell Signaling Technologies (Beverly, MA). Normal rabbit IgG (sc-2027), normal goat IgG (sc-2028), and donkey anti-mouse IgG (sc-2518), used as nonspecific antibodies in immunoprecipitations, were from Santa Cruz Biotechnology. Immobilized Protein A/G beads were from Pierce, and Protein G-Sepharose beads were from Amersham Biosciences. Horseradish peroxidase-conjugated secondary antibodies against either rabbit or mouse IgG, AffiniPure goat anti-mouse IgG light chain-specific, and IgG fraction monoclonal mouse anti-rabbit IgG light chain-specific secondary antibodies were from Jackson ImmunoResearch Laboratories (West Grove, PA).

WT, IKKaΔNBD, and IKkBΔNBD MEFs were generously provided by Dr. Inder Verma (Salk Institute for Biological Studies, La Jolla, CA). Plat-E cells were kindly provided by Dr. Tadaichi Kitamura (Institute of Medical Science, University of Tokyo, Japan), and Phoenix cells were provided by Dr. Garry Nolan (Stanford University, Stanford, CA). All cells were maintained in Dulbecco’s modified Eagle’s medium (Invitrogen) supplemented with 10% fetal calf serum, 2 mM l-glutamine, penicillin (50 units/ml), and streptomycin (50 μg/ml). For all experiments, unless otherwise indicated, cells were cultured in either 6-well tissue culture trays or 100-mm dishes and were stimulated with IL-1α (10 ng/ml) or TNF (10 ng/ml) when they reached 80% confluence.

**Generation of Stable Cell Lines**—All cloning procedures were performed by PCR using cloned Pfu DNA polymerase (Stratagene, La Jolla, CA). Complementary DNA encoding full-length IKKβ or IKKB1–733 (IKKBΔNBD) were subcloned into the HindIII and NotI sites of the LZRS-pBMN-lacZ retroviral vector (kindly provided by Dr. Garry Nolan). Resulting LZRS-IKKβWT and LZRS-IKKβΔNBD were transiently transfected using Fugene6 into Phoenix cells and selected for gene expression 24 h after transfection using puromycin (1 μg/ml). Puromycin-resistant cells were used to derive conditioned medium to provide a retroviral stock for MEF transduction. For cell transduction, IKKBΔNBD MEFs were washed and incubated for 8 h with retrovirus-conditioned medium containing Polybrene (8 μg/ml; Sigma). After incubation, retrovirus was removed and replaced with normal growth medium. The transduction process was repeated a further three times until cells became positive for IKKB when visualized by immunoblotting.
Role of the NBD in IKKα and IKKβ

To generate stably transduced IKKα<sup>WT</sup> and IKKα<sup>NBD</sup> MEFs, full-length IKKα or IKKα<sup>NBD</sup> cDNA was cloned into the EcoRI and XhoI restriction sites of retroviral GFP-MIGR1 vector (kindly provided by Dr. Warren Pear, University of Pennsylvania, Philadelphia, PA). The resulting MIG-IKKα<sup>WT</sup> and MIG-IKKα<sup>NBD</sup> were transiently transfected using Fugene6 into Plat-E cells to produce ectropic virus that was derived from conditioned medium containing Polybrene (8 µg/ml). For cell transduction, IKKα<sup>−/−</sup> MEFs were washed and incubated for 8 h with retrovirus-conditioned medium. After incubation, retrovirus was removed and replaced with normal growth medium. The transduction process was repeated a further three times until cells became positive for IKKα, as assessed by FACS analysis using Cell Quest software (FACSort; BD Biosciences).

For cell sorting, transduced cells were trypsinized and washed in FACS buffer (sterile phosphate-buffered saline, 0.5 mM EDTA, and 0.5% bovine serum albumin). Evaluation of green fluorescent protein was performed on a three-laser (argon (488 nm), krypton (407 nm), and dye laser (tuned to 600 nm)), 10-parameter FACSVantage<sup>TM</sup> obtained from BD Biosciences. Compensation and data analyses were performed using Flowjo software (Tree Star, Ashland, OR).

**Immunoblotting and Immunoprecipitation**— Cells were washed once with phosphate-buffered saline and then incubated for 10 min at 4 °C in 100 µl of TNT lysis buffer (50 mM Tris-Cl, pH 7.5, 150 mM NaCl, and 1% Triton X-100) and a complete miniprotease inhibitor mixture (Roche Applied Science). Samples were then scraped and harvested into 1.5-ml microcentrifuge tubes, vortexed for 30 s, and then centrifuged (425 × g for 10 min). Protein levels in supernatants were determined using a Coomassie protein assay kit (Bio-Rad), and 20 µg of protein from each sample was separated by SDS-PAGE (10%) followed by immunoblotting, as described above. The beads were washed three times with lysis buffer, and then samples were analyzed by SDS-PAGE (10%) followed by immunoblotting, as described above.

**Immune Complex Kinase Assay**—For immune complex kinase assays, MEF lysates were prepared, and immunoprecipitations were performed as described above. The resulting immunoprecipitates were washed extensively in TNT and then kinase buffer (20 mM HEPES, pH 7.5, 20 mM MgCl₂, 1 mM EDTA, 2 mM NaF, 2 mM β-glycerophosphate, 1 mM dithiothreitol, 10 mM ATP). Precipitates were then incubated for 15 min at 30 °C in 20 µl of kinase buffer containing glutathione-S-trans- ferase-IκBo (amino acids 1–54; a generous gift from Dr. Serge Fuchs, University of Pennsylvania) and 10 µCi of [γ-<sup>32</sup>P]ATP (Amersham Biosciences). The substrate was then precipitated using glutathione-agarose (Amersham Biosciences) and washed extensively with TNT. Beads were then suspended in 20 µl of sample buffer, and samples were separated by SDS-PAGE (10%). The resulting gel was stained with Coomassie Blue solution (0.05% Brilliant Blue G250, 40% methanol, 10% acetic acid) and then destained (40% methanol, 10% acetic acid) and dried. Kinase activity was determined by autoradiography.

**Generation of Nuclear Lysates for Immunoblotting**—MEFs were stimulated with TNFα (10 ng/ml) for the indicated times and then scraped into phosphate-buffered saline at 4 °C and pelleted (425 × g, 10 min). Pellets were resuspended and swollen for 10 min on ice in 100 µl of Buffer A (10 mM HEPES, pH 7.9, 10 mM KCl, 0.1 mM EDTA, 2 mM NaF, 2 mM β-glycerophosphate, and complete miniprotease inhibitors) plus 0.1% Nonidet P-40 and centrifuged (3800 × g) for 1 min. Supernatants (cytoplasmic fraction) were centrifuged at 20,000 × g for 1 h at 4 °C, and the resulting supernatants were snap frozen and retained for analysis. Pelleted nuclei were washed four times with 100 µl of Buffer A plus 0.1% Nonidet P-40 buffer before being lysed in 30 µl of 1% TNT, 1% SDS buffer plus complete mini protease inhibitors with a 26-gauge (half-inch) needle. Nuclear lysates were then centrifuged for 20 min (20,000 × g) at room temperature and then either used immediately or snap frozen and stored at −80 °C. Lysates were immunoblotted as described above.

**Transfections and Luciferase Reporter Assays**—WT MEFs and stable cell lines grown in 12-well plates (2.5 × 10⁵/well) were transiently transfected using Fugene6 (Roche Applied Science) following the manufacturer's protocol. Cells were transfected with a total of 0.11 µg of DNA/well, consisting of the NF-κB-dependent firefly reporter construct pBIIx-luciferase (0.2 µg/well) and a Renilla luciferase reporter (0.02 µg/ml). Cells were stimulated with TNFα or IL-1α for 5 h and then lysed in passive lysis buffer (Promega, San Luis Obispo, CA) 24–36 h after transfection. Samples were assayed using a Luminoscan 96-well automated luminometer (Thermo Labsystems, Franklin, MA), and firefly/Renilla luciferase ratios were calculated using Ascent software (Thermo LabSystems).

**Electrophoretic Mobility Shift Assays (EMSAs)**—MEFs were stimulated with TNFα (10 ng/ml) or IL-1α (10 ng/ml) for the indicated times and then scraped into phosphate-buffered saline at 4 °C and pelleted (425 × g, 10 min). Pellets were resuspended and swollen for 30 min on ice in 100 µl of Buffer A (10 mM HEPES, pH 7.9, 10 mM KCl, 0.1 mM EDTA, 2 mM NaF, 2 mM β-glycerophosphate, and complete mini protease inhibitors), incubated a further 5 min on ice in 0.1% Nonidet P-40, and then vortexed and centrifuged (3800 × g) for 1 min. Supernatants (cytoplasmic fraction) were centrifuged at 20,000 × g for 1 h at 4 °C, and the resulting supernatants were snap frozen and retained for analysis. Pelleted nuclei were washed once with 100 µl of Buffer A buffer before being vortexed in 30 µl of NaCl buffer (20 mM HEPES, pH 7.9, 0.4 mM NaCl, pH 8, 1 mM EDTA, pH 8, 2 mM NaF, 2 mM β-glycerophosphate, and complete mini protease inhibitors) for 1 h at 4 °C. Nuclear lysates were then centrifuged for 20 min (20,000 × g) at room temperature and then either used immediately or snap frozen and stored at −80 °C.

Single-stranded complementary oligonucleotides encompassing a consensus NF-κB site (upper strand, 5’-AGTT-
GAGGGGACTTCCCCAGGC-3’) or the Oct-1 probe (Santa Cruz Biotechnology) were annealed and then labeled with [γ-32P]ATP using T4 PNK (New England Biolabs, Beverly, MA). Labeled probe was purified using mini-Quick Spin columns (Roche Applied Science) according to the manufacturer’s instructions. For EMSA, 2–5 μg of nuclear extracts supplemented with 1 μg of poly(dI-dC) (Roche Applied Science) was incubated with an equal volume of 2× binding buffer (40 mM Tris-Cl, pH 7.9, 100 mM NaCl, 10 mM MgCl2, 2 mM EDTA, 20% glycerol, 0.2% Nonidet P-40, 2 mM dithiothreitol, 100 μg/ml bovine serum albumin) on ice for 10 min. After incubation, 1 μl of labeled probe was added, and then samples were incubated at room temperature for 20 min. Resulting DNA-NF-κB complexes were separated on 5% polyacrylamide nondenaturing gels by electrophoresis, and then gels were dried and visualized by autoradiography.

**RESULTS**

The IKKβ NBD Is Required for TNF- but Not IL-1-induced NF-κB Activation—To determine the effects of selectively disrupting the IKKβ-NEMO interaction on classical NF-κB signaling and IKK complex formation, we generated retroviral constructs encoding both WT IKKβ and a truncation mutant (residues 1–733) lacking the NBD (IKKβΔNBD) (Fig. 1A). We stably transduced IKKβ-deficient MEFs with these constructs and confirmed by immunoblotting that the cells were reconstituted with IKKβWT and IKKβΔNBD (Fig. 1B). FACS analysis verified that over 95% of the cells were stably transduced with each construct (data not shown). The levels of IKKβWT in multiple cell lines that we generated were consistently higher than endogenous IKKβ in WT MEFs. However, the levels of reconstituted IKKβΔNBD were comparable with endogenous IKKβ levels in WT cells (Fig. 1B, compare lanes 1 and 4). To examine the effects of reconstituted IKKβWT and IKKβΔNBD on classical NF-κB activation, we incubated WT, IKKβ−/−, IKKβWT, and IKKβΔNBD MEFs for a range of times with TNF or IL-1 and then immunoblotted the resulting lysates using anti-ικBα. Following cytokine treatment, ικBα degradation and resynthesis was intact in WT MEFs, and consistent with our earlier findings (30), IL-1 but not TNF induced robust ικBα degradation in IKKβ−/− cells (Fig. 1C). As expected, TNF-stimulated ικBα degradation was restored in IKKβWT-reconstituted MEFs (Fig. 1C). In contrast, TNF did not induce ικBα degradation in IKKβΔNBD-reconstituted cells (Fig. 1C), confirming that the IKKβ NBD is required for TNF-induced classical NF-κB pathway activation.

We previously showed that IKK complexes containing NEMO and either IKKα or IKKβ alone could facilitate IL-1-induced ικBα phosphorylation and classical NF-κB activation (30). Since IKKβ has been shown to be a more potent kinase for ικBα than IKKα (32, 33), we questioned whether reconstituting IKKβ−/− MEFs with IKKβΔNBD inhibited or disrupted NEMO-IKKα-dependent IL-1 signaling. As shown in Fig. 1C (right panels), IL-1-induced ικBα degradation was intact in IKKβ−/− MEFs reconstituted with either IKKβWT or IKKβΔNBD. Furthermore, EMSA analysis demonstrated that the kinetics of IL-1-induced NF-κB nuclear translocation and DNA binding in IKKβ−/−, IKKβWT, and IKKβΔNBD cells were similar to WT MEFs (Fig. 1D). Hence, IKKβΔNBD does not block IL-1-induced classical NF-κB activation.

We next examined basal and IL-1-induced NF-κB transcriptional activity in the panel of MEFs using an NF-κB-specific luciferase reporter assay. Consistent with our previous results (30), IL-1 increased NF-κB activity in IKKβ−/− MEFs to a similar level as that observed in WT cells (Fig. 1E). Both basal and IL-1-stimulated levels of NF-κB activity were slightly enhanced in IKKβWT-reconstituted MEFs, possibly due to the higher levels of IKKβ expression in these cells compared with WT MEFs (Fig. 1B). However, basal and IL-1-induced NF-κB-dependent transcriptional activity was enhanced in IKKβΔNBD MEFs (Fig. 1E) that contained levels of IKKβΔNBD comparable with WT MEFs (Fig. 1B). This finding is consistent with our previous observation of enhanced NF-κB activation in HeLa cells transiently overexpressing NEMO-binding mutants of IKKβ (13, 14).
Role of the NBD in IKKα and IKKβ

FIGURE 1. The IKKβ NBD is required for TNF- but not IL-1-induced classical NF-κB activation. A, the structural domains of wild-type IKKβ (IKKβWT) and IKKβΔNBD are shown. KD, kinase domain; LZ, leucine zipper; ULD, ubiquitin-like domain; HLH, helix-loop-helix. IKKβΔNBD is a truncation mutant encompassing residues 1–733 that lacks the C-terminal 23 amino acids containing the NBD (13, 14). B, lysates from WT, IKKβ−/− (β−/−), IKKβWT (βWT), and IKKβΔNBD (βΔNBD) MEFs were immunoblotted using the antibodies indicated (right). C, WT, IKKβ−/−, IKKβWT, and IKKβΔNBD MEFs were incubated with either TNF (10 ng/ml) (left) or IL-1α (10 ng/ml) (right) for the indicated times, and then lysates were immunoblotted using anti-IκBα or anti-tubulin (Tub.) as a loading control. D, the same panel of MEFs was treated with IL-1α for the indicated times, and then nuclear extracts were prepared for EMSA. Assays were performed using either a consensus NF-κB binding site probe (top) or an Oct-1 probe as a loading control (bottom). E, MEFs were transiently transfected with the NF-κB-dependent reporter pBlix-firefly luciferase together with β-actin Renilla luciferase. Twenty-four hours later, cells were either left untreated or treated for a further 5 h with IL-1α, and then NF-κB activity was determined by a dual luciferase assay.

FIGURE 2. IKKβΔNBD associates with endogenous IKKα but not with endogenous NEMO. IKK complexes in whole cell lysates of WT, IKKβ−/− (β−/−), IKKβWT (βWT), and IKKβΔNBD (βΔNBD) MEFs were immunoprecipitated (IP) using anti-NEMO (A), anti-IKKα (B), or anti-IKKβ (C). Immunoprecipitated material was immunoblotted using anti-IKKα, anti-IKKβ, and anti-NEMO as indicated (right). D, samples of lysates saved prior to immunoprecipitation (Pre-IP) were immunoblotted using anti-IKKα, anti-IKKβ, and anti-NEMO as shown.

14) and supports our earlier conclusion that the IKKβ NBD plays a role in maintaining the basal activity of the IKK complex. IKKβΔNBD is not incorporated into the IKK Holocomplex—Within the tripartite IKK holocomplex, IKKα and IKKβ interact via their respective leucine zipper domains (34). Each IKK subunit in turn associates via its C-terminal NBD with NEMO (13, 14, 35). We therefore questioned whether IKKβΔNBD was incorporated into a trimeric IKK complex in IKKβ−/− MEFs via interaction with endogenous IKKα associated with NEMO. To determine the effects of deleting the IKKβ NBD on complex formation, we immunoprecipitated IKK complexes from WT, IKKβ−/−, IKKβWT, and IKKβΔNBD MEFs using anti-NEMO, anti-IKKα, and anti-IKKβ. As shown in Fig. 2A, IKKα and IKKβ both co-immunoprecipitated with NEMO from WT and IKKβWT-reconstituted MEFs. In contrast, only IKKα associated with NEMO in IKKβ−/− and IKKβΔNBD-reconstituted cells. When immunoprecipitations were performed using anti-IKKα, both NEMO and IKKβ co-precipitated with IKKα from WT, IKKβWT, and IKKβΔNBD MEFs, whereas only NEMO associated with IKKα in IKKβ−/− cells (Fig. 2B). Immunoprecipitation using anti-IKKβ pulled down all three IKK complex sub-
units from WT and IKKβ<sup>WT</sup>-reconstituted MEFs, but as expected, anti-IKKβ did not precipitate any of these proteins from IKKβ<sup>−/−</sup> cells (Fig. 2C, lane 2). Intriguingly, anti-IKKβ pulled down complexes consisting of IKKβ<sup>ΔNBD</sup> and IKKα but not NEMO from IKKβ<sup>ΔNBD</sup>-reconstituted MEFs (Fig. 2C, lane 4). These findings therefore suggest that reconstituted IKKβ<sup>WT</sup> is incorporated into the heterotrimeric and IKK complex, whereas IKKβ<sup>ΔNBD</sup> associates only with endogenous IKKα in a separate complex that does not contain NEMO.

To further explore the nature of the IKK complexes in the reconstituted cell lines, we performed size exclusion chromatography of S100 lysates. As shown in Fig. 3A (top), and consistent with previous reports (10, 34, 36, 37), IKKα, IKKβ, and NEMO in WT MEFs co-eluted in fractions corresponding to a predicted molecular mass of 600–900 kDa. Immunoprecipitation of complexes from these fractions using anti-NEMO or anti-IKKα confirmed that all three subunits co-precipitated, with the majority of the IKK holocomplex appearing in fractions 18 and 19 (Fig. 3A, bottom). Fractionation of lysates from IKKβ<sup>−/−</sup> cells showed that IKKα and NEMO elute together in fractions 18–21, suggesting a complex of ~500–900 kDa (Fig. 3B, top). In addition, IKKα eluted in fractions 22–24, which did not contain any NEMO. Immunoprecipitation of complexes from fractions 17–23 using anti-NEMO demonstrated association of IKKα with NEMO in fractions 18–21 (Fig. 3B, bottom). These data therefore demonstrate that in IKKβ<sup>−/−</sup> MEFs, an IKK complex consisting of only NEMO and IKKα exists that is similar in size to the tripartite complex in WT cells. In addition, a portion of IKKα does not associate with NEMO in the absence of IKKβ but instead elutes in fractions corresponding to a lower molecular weight NEMO-independent complex.

All three IKK complex subunits eluted in high molecular weight fractions (fractions 17–20) in IKKβ<sup>WT</sup>-reconstituted IKKβ<sup>−/−</sup> MEFs (Fig. 3C, top), and immunoprecipitation demonstrated that IKKα, IKKβ, and NEMO associate in these fractions (Fig. 3C, bottom). Similar to IKKβ<sup>−/−</sup> cells, IKKα also appeared in fractions 21–24, which did not contain NEMO, and NEMO-IKKα complexes could not be immunoprecipitated in these fractions. Hence, reconstituted IKKβ<sup>WT</sup> incorporates into the high molecular weight tripartite IKK holocomplex in IKKβ<sup>−/−</sup> MEFs, and lower molecular weight endogenous IKKα-containing complexes are also detected in these cells. We also consistently observed IKKβ in later eluting fractions up to sample number 27. IKKβ in these fractions was not associated with either IKKα or NEMO (Fig. 3C, bottom) (data not shown), suggesting that it is an IKKβ homodimeric complex possibly resulting from the high level of overexpression in these cells (Fig. 1B).
Role of the NBD in IKKα and IKKB

Similar to IKKB\(^{-/-}\) MEFs, IKKα and NEMO eluted together in fractions 17–21 in IKK\(^{\text{BND}}\) cell lysates, suggesting that they exist as a complex comparable in size with native IKK in WT MEFs (Fig. 3D, top). IKK\(^{\text{BND}}\) did not co-elute with the majority of IKKα and NEMO but instead eluted in two pools with peaks in fractions 20–22 and 26–27, respectively. The first pool of IKK\(^{\text{BND}}\) co-eluted with IKKα in fractions 19–23, whereas the second pool did not co-elute with either IKKα or NEMO. When immunoprecipitations were performed using anti-NEMO (Fig. 3D, bottom), IKKα but not IKK\(^{\text{BND}}\) associated with NEMO in the high molecular weight fractions. Similarly, NEMO co-precipitated with anti-IKKα in fractions 17–21, but IKK\(^{\text{BND}}\) only associated with IKKα in fractions 20–23.

Collectively, these data support our findings in Fig. 2 and demonstrate that reconstituted IKK\(^{\text{BND}}\) does not incorporate into the tripartite IKK complex in IKK\(^{-/-}\) MEFs. In these cells, IKKα and NEMO form the high molecular weight IKK complex, whereas IKK\(^{\text{BND}}\) associates with IKKα in a separate NEMO-independent complex. In addition, IKK\(^{\text{BND}}\) appears in a third, lower molecular weight complex that does not contain either IKKα or NEMO.

IKK\(^{\text{BND}}\) Restores Noncanonical NF-κB Signaling in IKK\(^{-/-}\) MEFs—To establish the effects of selectively disrupting the IKKα-NEMO interaction on NF-κB signaling and IKK complex formation, we generated retroviral constructs encoding both WT IKKα and a truncation mutant (residues 1–745) lacking the NBD (IKKα\(^{\text{BND}}\)) (Fig. 4A) and used these to stably transduce IKKα-deficient MEFs. Immunoblotting demonstrated that the cells were reconstituted with IKKα\(^{\text{WT}}\) and IKKα\(^{\text{BND}}\) (Fig. 4B), and FACS analysis confirmed that over 95% of the cells were stably transduced with each construct (not shown). The levels of IKKα\(^{\text{WT}}\) and IKKα\(^{\text{BND}}\) in multiple stable cells lines generated were consistently similar to those of endogenous IKKα in WT MEFs.

Noncanonical NF-κB signaling requires IKKα but is intact in NEMO-deficient cells (18–20, 22). However, it remains unclear whether disrupting the IKKα NBD in cells containing NEMO affects noncanonical signaling. To address this question, we incubated our panel of MEFs with heterotrimeric lymphotoxin (LT\(\alpha\)β2) and then immunoblotted the resulting lysates using anti-p100/p52. As shown in Fig. 4C (lanes 1–3), LT\(\alpha\)β2 induced the appearance of p52 in WT MEFs, indicating activation of the noncanonical pathway. As expected, p52 was absent in LT\(\alpha\)β2-stimulated IKKα\(^{-/-}\) cells (lanes 4–6) but was present in IKKα\(^{\text{WT}}\)-reconstituted MEFs (lanes 7–9). Similarly, IKKα\(^{\text{BND}}\) reconstitution of IKKα\(^{-/-}\) MEFs restored LT\(\alpha\)β2-induced p100 processing to p52, demonstrating that the IKKα NBD is not required for noncanonical NF-κB activation.

IKKα\(^{\text{BND}}\) Rescues Classical NF-κB Activity in IKKα\(^{-/-}\) MEFs—

We next questioned whether loss of the IKKα NBD affects proinflammatory cytokine-induced classical NF-κB activation. Since our earlier study demonstrated that a NEMO-IKKα complex facilitates IL-1-induced classical NF-κB activation (30), we asked whether IKKα\(^{\text{BND}}\) might function as a dominant negative inhibitor of IL-1 signaling. However, consistent with our findings with IKKα\(^{\text{BND}}\) (Fig. 1C), both IL-1 and TNF induced IkBα degradation in IKKα\(^{\text{WT}}\) and IKKα\(^{\text{BND}}\)-reconstituted MEFs (Fig. 5A). Hence, IKKα\(^{\text{BND}}\) does not inhibit the ability of IKKβ to phosphorylate IkBα, leading to its degradation.

Although TNF- and IL-1-induced IkBα degradation is intact in IKKα\(^{-/-}\) cells, we found previously that DNA binding of nuclear NF-κB in response to IL-1 is diminished in the absence of IKKα compared with either wild-type or IKKβ\(^{-/-}\) MEFs (30). To determine whether IKKα\(^{\text{WT}}\) or IKKα\(^{\text{BND}}\) could rescue defective NF-κB activation in IKKα-deficient MEFs, we performed EMSAs using our panel of cells. As shown in Fig. 5B, IL-1-induced NF-κB activation was maximal in WT MEFs after 15 min and returned to basal levels by 60 min (lanes 1–4). Consistent with our previous data (30), DNA binding in IKKα\(^{-/-}\) MEFs was less robust than that observed in WT MEFs (compare lanes 1–4 and 5–8). However, reintroduction of either IKKα\(^{\text{WT}}\) or IKKα\(^{\text{BND}}\) into IKKα\(^{-/-}\) MEFs restored IL-1-induced NF-κB DNA binding to WT levels (Fig. 5B; lanes 9–16). Surprisingly, we also found that TNF-induced NF-κB activation was diminished in IKKα\(^{-/-}\) MEFs (Fig. 5C, lanes 1–8), and similar to IL-1 signaling, this defect was rescued by reintroduction of either IKKα\(^{\text{WT}}\) or IKKα\(^{\text{BND}}\) (lanes 9–16).

Previous studies have demonstrated that IKKα enters the nucleus and regulates NF-κB transcriptional activity via mechanisms including phosphorylation of p65 and histone H3 (24–29, 38). We therefore performed NF-κB-dependent luciferase reporter assays to determine the effects of deleting the IKKα NBD on transcriptional activation of NF-κB. Confirming earlier reports (24, 29, 38), we found that neither IL-1 nor TNF could activate NF-κB-dependent transcription in IKKα\(^{-/-}\) MEFs (Fig. 5D). Stable re-expression of IKKα\(^{\text{WT}}\) led to an increase in basal transcriptional activity compared with WT MEFs but also rescued the ability of IL-1 and TNF to up-regulate transcription in these cells (Fig. 5D). Furthermore, IKKα\(^{\text{BND}}\) also restored basal and cytokine-induced NF-κB

FIGURE 4. The IKKα NBD is not required for noncanonical NF-κB activation. A, the structural domains of wild-type IKKα (IKKα\(^{\text{WT}}\)) and IKKα\(^{\text{BND}}\) are shown. KD, kinase domain; LZ, leucine zipper; HLH, helix-loop-helix. IKKα\(^{\text{BND}}\) is a truncation mutant encompassing residues 1–737 that lacks the C-terminal 8 amino acids containing the NBD (13, 14). B, lysates from WT, IKKα\(^{-/-}\) (α\(^{-/-}\)), IKKα\(^{\text{WT}}\) (α\(^{\text{WT}}\)), and IKKα\(^{\text{BND}}\) (α\(^{\text{BND}}\)) MEFs were immunoblotted using the antibodies indicated (right). C, WT, IKKα\(^{-/-}\), IKKα\(^{\text{WT}}\), and IKKα\(^{\text{BND}}\) MEFs were either untreated or incubated with LT\(\alpha\)β2 (L7) for the times indicated, and then lysates were immunoblotted using either anti-p100/p52 (top), anti-IKKα, or anti-tubulin (Tub.) as indicated (right).
The ability of IKKα to regulate NF-κB-dependent transcription has been shown to involve phosphorylation of p65 at Ser536 (28, 38, 39), we immunoblotted cell lysates using an antibody that recognizes phosphorylated Ser536 on p65. As shown in Fig. 5E, Ser536 phosphorylation in response to TNF was severely diminished in IKKα−/− cells, whereas re-expression of either IKKαWT or IKKαNBD restored TNF-induced p65 phosphorylation to levels similar to that in WT MEFs (Fig. 5E).

In addition to IKKα, NEMO has also been shown to shuttle between the cytoplasm and nucleus (40–43). We therefore questioned whether the ability of IKKα to enter the nucleus, where it regulates NF-κB transcriptional activity, requires its association with NEMO. As shown in Fig. 5F (top), IKKα was present in the nucleus of untreated WT, IKKαWT, and IKKαNBD MEFs, and neither TNF nor IL-1 (not shown) stimulation further enhanced its nuclear localization. Furthermore, NEMO was present in the nucleus in WT, IKKα−/−, and IKKαWT MEFS but was severely depleted in the nucleus of IKKαNBD cells. As previously shown (24, 29), IKKβ was absent in the nucleus of WT cells. Surprisingly, however, we detected elevated nuclear IKKβ in IKKα−/− MEFS that was significantly diminished following reconstitution with either IKKαWT or IKKαNBD. Notably, NEMO was also present in the nucleus of IKKα−/− MEFS.

To determine whether nuclear IKKα in the reconstituted cells is activated in response to proinflammatory stimuli, we immunoblotted nuclear lysates using a phospho-specific antibody that recognizes both IKKα and IKKβ only when they are phosphorylated on crucial serines within their catalytic activation loops. As shown in Fig. 5G, this antibody detected phosphorylated IKKα in WT, IKKαWT, and IKKαNBD cells, demonstrating that IKKα in the nuclei of these cells is activated in response to TNF. However, although IKKβ was present in the nucleus of IKKα−/− cells (Fig. 5G, lanes 3 and 4), we did not detect its phosphorylation, suggesting that nuclear IKKβ is not activated in response to TNF.

Taken together, the findings in Fig. 5 strongly support a model in which the transcriptional regulatory activity of IKKα is independent of its function as a NEMO-associated IκB kinase. Our data also suggest that the ability of IKKα to enter the nucleus and become activated in response to proinflammatory cytokines is NEMO-independent, whereas normal nuclear localization of NEMO requires an intact IKKα NBD. Finally, these results indicate that IKKβ can enter the nucleus but only in the absence of IKKα.

**IKKαNBD Is Not Incorporated into the IKK Holocomplex**

To determine whether IKKαNBD was incorporated into the IKK holocomplex we immunoprecipitated IKK complexes from WT, IKKα−/−, IKKαWT, and IKKαNBD MEFS using anti-NEMO and anti-IKKα. As shown in Fig. 6A, IKKα and IKKβ both co-immunoprecipitated with anti-NEMO from WT and IKKαWT-reconstituted MEFS, whereas only IKKβ associated with NEMO in IKKα−/− and IKKαNBD cells. When anti-IKKα was used for immunoprecipitations, both NEMO and IKKβ co-precipitated with IKKα from WT and IKKαWT MEFS (Fig. 6C).
Role of the NBD in IKKα and IKKβ

(A) WT α<sup>+</sup> α<sup>WT</sup> α<sup>αNBD</sup>
IP: NEMO
IKKα
IKKβ
NEMO
Pre-IP
1 2 3 4

(B) WT α<sup>+</sup> α<sup>WT</sup> α<sup>αNBD</sup>
IP: IKKα
IKKα
IKKβ
NEMO
Pre-IP
1 2 3 4

(C) IKKα<sup>+</sup>
IP: NEMO
IKKα
IKKβ
NEMO
17 18 19 20 21

(D) IKKα<sup>WT</sup>
IP: NEMO
IKKα
IKKβ
NEMO
17 18 19 20 21

(E) IKKα<sup>αNBD</sup>
IP: NEMO
IKKα
IKKβ
NEMO
17 18 19 20 21 22 23 24 25

FIGURE 6. IKKα<sup>αNBD</sup> is not incorporated into the tripartite IKK complex. A and B, IKK complexes in whole cell lysates of wild-type (WT), IKKα<sup>−/−</sup> (α<sup>−/−</sup>), IKKα<sup>WT</sup> (α<sup>WT</sup>), and IKKα<sup>αNBD</sup> (α<sup>αNBD</sup>) MEFs were immunoprecipitated (IP) using anti-NEMO (A) or anti-IKKα (B). Immunoprecipitated material was immunoblotted using anti-IKKα, anti-IKKβ, and anti-NEMO as indicated (right). Samples of lysates saved prior to immunoprecipitation (Pre-IP) were immunoblotted using anti-IKKα, anti-IKKβ, and anti-NEMO as shown. C–E, S100 extracts from IKKα<sup>−/−</sup> (C), IKKα<sup>WT</sup> (D), and IKKα<sup>αNBD</sup> (E) MEFs were fractionated by size exclusion chromatography. Fractions were immunoblotted using the antibodies indicated (left). The column was precalibrated, and the molecular weights of standard proteins are indicated above the appropriate fractions in C. Fractions containing the high molecular weight IKK complex were immunoprecipitated using either anti-NEMO or anti-IKKα. The resulting immunoblots from these immunoprecipitations are displayed below the fractionation profile for each cell type in C–E.

In contrast, only IKKβ associated with IKKα<sup>αNBD</sup>, suggesting that IKKα<sup>αNBD</sup> forms a NEMO-independent complex with endogenous IKKβ (Fig. 6B).

To further investigate the complexes formed in the reconstituted cell lines, we performed size exclusion chromatography. Following separation of lysates from IKKα<sup>−/−</sup> cells, IKKβ and NEMO eluted together in fractions 18–21 (Fig. 6C, top), and immunoprecipitations using anti-NEMO demonstrated that IKKβ associated with NEMO in these fractions (Fig. 6C, bottom). This profile suggests a NEMO-IKKβ complex of similar size (500–900 kDa) to the tripartite IKK complex in wild-type MEFs that predominantly eluted in fractions 18 and 19 (Fig. 3A). When we reconstituted IKKα<sup>−/−</sup> MEFs with IKKα<sup>WT</sup>, the bulk of IKK complexes eluted in fractions 17–20 (Fig. 6D, top), and immunoprecipitation using either anti-NEMO or anti-IKKα confirmed that these complexes contain IKKα, IKKβ, and NEMO (Fig. 6D, bottom). Thus, IKKα<sup>WT</sup> is incorporated into the high molecular weight tripartite IKK complex in IKKα<sup>−/−</sup> cells.

Fractionation of lysates from IKKα<sup>αNBD</sup>-reconstituted MEFs showed that IKKα<sup>αNBD</sup>, IKKβ, and NEMO elute in a broad range of fractions spanning predicted molecular masses from ~250 to 900 kDa (Fig. 6E, top). When immunoprecipitations were performed using anti-NEMO, IKKβ but not IKKα<sup>αNBD</sup> associated with NEMO in fractions 19–24 (Fig. 6E, bottom). In contrast, when anti-IKKα was used to immunoprecipitate complexes, only IKKβ associated with IKKα<sup>αNBD</sup> in fractions 18–25, and NEMO was not co-precipitated in these samples. Hence, these data support our findings in Fig. 6, A and B, and demonstrate that reconstituted IKKα<sup>αNBD</sup> is not incorporated into the tripartite complex. Instead, similar to IKKβ<sup>αNBD</sup> in IKKβ<sup>−/−</sup>-MEFs (Figs. 2 and 3), endogenous IKKβ and NEMO form a high molecular weight IKK complex, whereas IKKα<sup>αNBD</sup> associates with IKKβ in a distinct NEMO-independent complex.

In addition to these complexes, our analysis of nuclear extracts demonstrated that IKKα<sup>αNBD</sup> exists alone in the nucleus separate from either IKKβ or NEMO (Fig. 5F). Furthermore, nuclear IKKα<sup>αNBD</sup> was active as measured using an anti-phospho-IKKα/β antibody (Fig. 5G). Since NEMO is absolutely required for activation of the cytoplasmic IKK complex leading to IkBα phosphorylation (8–10, 12), we questioned whether the NEMO-independent IKKβ-IKKα<sup>αNBD</sup> complex could be activated in response to IL-1α or TNF. To test this, we immunoprecipitated IKK complexes from cytoplasmic extracts of IKKα<sup>αNBD</sup> and IKKα<sup>WT</sup> MEFs using either anti-IKKα or anti-
Role of the NBD in IKKα and IKKβ

Contrast, TNF signaling absolutely requires IKKβ, and the classical NF-κB pathway cannot be activated by TNF in IKKβ-deficient cells (16, 17). Since both IKKα and IKKβ associate with NEMO via their C-terminal NBDs, these findings suggest that the ability of both kinases to bind NEMO regulates their function within the classical signaling pathway. We therefore undertook this study to determine the effects of selectively deleting the NBD in IKKα and IKKβ on TNF- and IL-1-induced NF-κB activation and IKK complex formation. Our findings are summarized in the model depicted in Fig. 8.

Consistent with the stringent requirement for NEMO and IKKβ in TNF signaling (8–12), IKKβ^ΔNBD did not rescue defective TNF-induced IkBa degradation in IKKβ^−/− cells. This conclusively demonstrates that direct association of NEMO with IKKβ is absolutely necessary for TNF-induced classical NF-κB activation. Confirming our previous study (30), IL-1-induced IkBa degradation and NF-κB activation was intact in IKKβ^−/− MEFs. In light of the failure of IKKβ^ΔNBD to rescue TNF signaling and previous studies demonstrating that IKKβ is a more efficient kinase for IkBa than IKKα (32, 33), we speculated that IKKβ^ΔNBD might function as a dominant negative and block IL-1 signaling. However, reconstitution of IKKβ^ΔNBD with IKKβ^ΔNBD did not affect IL-1-induced IkBa degradation or NF-κB activation. This is in line with our earlier finding that a catalytically inactive IKKβ mutant that blocks TNF signaling did not affect IL-1-induced classical NF-κB activation (30). We therefore conclude that IL-1-induced NEMO-dependent IKKα activation overcomes the potential dominant negative effects of catalytically inactive or NBD-deficient IKKβ.

These data underscore the ability of IKKα to either compensate in the absence of IKKβ or function in the presence of defective IKKβ in IL-1- but not TNF-induced classical pathway activation.

Although IKKβ^ΔNBD did not affect IL-1-induced IkBa degradation and NF-κB activation in IKKβ^−/− MEFs, NF-κB transcriptional activity was markedly enhanced in IKKβ^ΔNBD cells. IKKβ NEMO binding mutants were shown previously to be constitutively active when overexpressed in HeLa cells (13, 14), and certain deubiquitinases and protein phosphatases that inhibit IKK activity have been reported to associate with NEMO following IKK activation (44–49). Hence, the inability of IKKβ^ΔNBD to recruit these enzymes via NEMO might lead to its increased basal activity. We previously proposed that phosphorylation within the IKKβ NBD is a negative regulatory signal that affects the function of NEMO (13, 14). Several lines of biochemical evidence support this hypothesis (13, 14, 50–53), and Higashimoto et al. (51) demonstrated recently that phosphorylation within the NBD by Polo-like kinase down-regulates IKK activity. Notably, however, basal NF-κB activation measured by EMSA was not enhanced in IKKβ^ΔNBD cells, suggesting that additional regulatory signals control IKK and NF-κB transcriptional activity. In this regard, CUEDC2 was shown recently to directly associate with IKKα and IKKβ to inhibit their activity, and this probably occurs in the absence of NEMO (54).

Clearly, further work is required to understand fully the mechanisms that deactivate the IKK complex; however, these accumulated findings suggest that both NEMO-dependent and -independent mechanisms exist to accomplish this.
Role of the NBD in IKKα and IKKB

The noncanonical NF-κB pathway requires IKKα and can be activated in the absence of either IKKB or NEMO (18–20, 22). We confirmed this by rescuing defective lymphotixin-β receptor-induced p100 processing in IKKα−/− MEFs with IKKα. Moreover, IKKαNBD also rescued p100 processing, providing further evidence that the association of IKKα with NEMO plays no role in noncanonical pathway activation. We therefore conclude that disrupting the interaction of NEMO with the IKKs selectively blocks the classical pathway and does not affect noncanonical NF-κB signaling. This strongly supports the potential use of the NBD peptide or other pharmacological strategies targeting this interaction as highly specific inhibitors of the classical NF-κB pathway (5, 13, 14, 35).

Consistent with our earlier findings (30) and those of others (17, 55), IL-1 and TNF-induced IκBa degradation was intact in IKKα-deficient cells. Furthermore, IKKαNBD did not block either TNF- or IL-1-induced IκBa degradation, demonstrating that IKKαNBD does not function as a dominant negative inhibitor of IKKα. Together with the effects of re-expressing IKKBNBD in IKKβ−/− MEFs described above, this provides further support for our previously proposed model of classical pathway activation (30). In this model, TNF signaling absolutely requires IKKβ associated with NEMO, whereas IL-1 is less stringent and utilizes NEMO with either IKKα or IKKβ to phosphorylate IκBa (30).

Although IL-1 and TNF both induced IκBa degradation in IKKα−/− cells, neither cytokine activated NF-κB transcriptional activity in the absence of IKKα. In addition, DNA binding of nuclear NF-κB was significantly reduced in IKKα-deficient cells. These observations confirm previous reports describing critical roles for IKKα in directly phosphorylating and regulating the transcriptional activity of NF-κB proteins as well as transcriptional co-activators, co-repressors, and histones (24–28, 56). Consistent with these functions, IKKαWT rescued full DNA binding and transcriptional activity in IKKα−/− cells, and intriguingly, activity was also rescued by IKKαNBD. Furthermore, both IKKβWT and IKKαNBD rescued TNF- and IL-1-induced phosphorylation of p65, which is severely diminished in IKKα−/− cells. This therefore demonstrates that association with NEMO is not required for IKKα to regulate the transcriptional activity of NF-κB that occurs, at least in part, via phosphorylation of p65.

The ability of IKKα to regulate NF-κB transcriptional activity depends on its capacity to enter the nucleus (24–28, 56). It has also been shown that NEMO shuttles in and out of the nucleus and plays a key role in DNA damage-induced NF-κB activation (40–42, 57). In addition, NEMO regulates transcription by binding to CBP at NF-κB-dependent promoters (43). We therefore questioned whether nuclear translocation of NEMO requires its association with IKKα and whether nuclear IKKα activity requires NEMO. As previously described, IKKα and NEMO, but not IKKβ, were present in the nuclei of WT MEFs (24, 29); however, cytokine-stimulation did not affect the nuclear levels of either of these proteins. TNF stimulation activated nuclear IKKα in WT MEFs and also activated IKKβ that was present in the nuclei of both IKKαWT and IKKαNBD cells. Since our immune complex kinase analysis demonstrated that cytoplasmic IKKαNBD complexes cannot be activated, these findings strongly suggest that IKKα activation occurs in the nucleus. Further work is clearly required to determine the precise mechanisms of activation of nuclear IKKα; however, our data demonstrate that this occurs independently of its association with NEMO.

Surprisingly, we found IKKβ and NEMO in the nucleus of IKKα−/− cells. Unlike IKKα, IKKβ does not contain a nuclear localization sequence, suggesting that in the absence of IKKα, NEMO chaperones IKKβ into the nucleus (28). Notably, and in contrast to nuclear IKKα, we did not detect any TNF-induced activation of nuclear IKKβ in IKKα−/− cells. When we re-expressed IKKαWT in IKKα−/− cells, the cytoplasmic and nuclear distribution of IKKα, IKKβ, and NEMO was identical to that in

**FIGURE 8. Classical NF-κB signaling and IKK complex formation in IKK-reconstituted cells.** Endogenous proteins are shown as open ovals (N, NEMO; α, IKKα; β, IKKβ), and reconstituted IKKα and IKKβ are depicted in black. IKKαNBD and IKKBNBD are flattened to indicate deletion of the NBD. Activated NF-κB is shown as gray ovals, and the gray arrows indicate transcriptional activity. A, IKKβ−/− MEFs (left) contain NEMO-IKKα complexes that activate classical NF-κB in response to IL-1 (arrow) but not TNF (blunt-ended line). Reconstituted IKKBWT forms a heterotrimeric IKK complex with endogenous NEMO and IKKα and rescues NFκB signaling (middle). In contrast, IKKαNBD neither affects IL-1 signaling nor rescues NFκB signaling (right). IKKβNBD is not incorporated into the heterotrimeric IKK complex but forms complexes with endogenous IKKα that do not contain NEMO. B, IKKα−/− MEFs contain NEMO-IKKκ complexes that activate classical signaling (i.e. IκBa degradation and NFκκ nuclear localization) in response to TNF and IL-1; however, NFκκ transcriptional activity is defective in these cells (left). IKKαWT (middle) and IKKαNBD (right) both rescue transcriptional activity, demonstrating that association with NEMO is not required for the ability of IKKα to regulate transcription. IKKαWT forms a heterotrimeric IKK complex with endogenous NEMO and IKKβ, and a portion of IKKαWT enters the nucleus (middle). IKKαNBD is not incorporated into the heterotrimeric IKK complex but instead forms NEMO-independent complexes with endogenous IKKβ (right). Nuclear localization of IKKα does not require NEMO association as IKKαNBD enters the nucleus.
WT MEFs, demonstrating that IKKα maintains these homeostatic levels. Furthermore, in IKKα\(^{\text{ΔNBD}}\)-reconstituted MEFs we found IKKα in the nucleus in the absence of significant levels of NEMO. Taken together, these findings demonstrate that although NEMO binding is not required for IKKα to enter the nucleus and regulate NF-κB transcription, the IKKα NBD is critical for NEMO nuclear localization in cells containing IKKα. Future efforts to fully dissect the mechanisms regulating nuclear localization of the separate IKK complex subunits are necessary, but our findings suggest that these mechanisms are not mutually exclusive.

IKKα and IKKβ associate via interactions between their respective leucine zipper domains (34), and the most abundant form of the IKK complex is a heterodimer of IKKα and IKKβ associated with NEMO (1, 6, 7). This suggests that IKK heterodimerization is the preferential conformation for the IKKs, although these heterodimers did not bind to NEMO despite the presence of an intact NBD on the endogenous kinase. Furthermore, our immune complex kinase assay and NBD peptide gel filtration analysis confirmed that IKK\(^{\text{WT}}\) re-expressed IKK\(^{\text{ΔNBD}}\) and IKKβ\(^{-/-}\) MEFs, respectively, formed heterotrimeric complexes with endogenous IKKs and NEMO (see model in Fig. 8). In contrast, although IKKα\(^{\text{ΔNBD}}\) and IKKβ\(^{\text{ΔNBD}}\) associated with endogenous IKKβ and IKKα, respectively, these heterodimers did not bind to NEMO despite the absence of an intact NBD on the endogenous kinase. Furthermore, our immune complex kinase assay and NBD peptide experiments clearly demonstrate that in IKKα\(^{\text{ΔNBD}}\) cells, the IKKβ-IKKα\(^{\text{ΔNBD}}\) complex is not activated by IL-1α or TNF, and it is the NEMO-IKKβ complex that responds to stimulation. Consequently, these findings demonstrate that in the absence of one NBD, the heterotrimeric IKK holocomplex cannot assemble, and the resulting NEMO-independent complexes do not respond to proinflammatory cytokine signaling.

Recent reports have revealed that NEMO forms dimers through interactions between specific domains within the N-terminal region of the protein (35, 60). Furthermore, this portion of NEMO is necessary and sufficient for association with the NBDs of IKKα and IKKβ (13, 35, 52, 60). Recent elegant crystallographic studies revealed that this dimeric conformation of the NEMO N terminus forms two parallel 800-Å α-helical IKK-binding pockets into which the NBDs insert (35). Since the heterodimeric complexes formed between endogenous IKK and re-expressed IKK\(^{\text{ΔNBD}}\) did not associate with NEMO, we conclude that the presence of two NBDs is essential for stable formation of this interaction with NEMO. It is intriguing to speculate that the single NBD IKK heterodimer cannot “clamp” into the IKK binding pocket formed by the NEMO dimers; however, extensive structural analysis of these ΔNBD complexes is required to draw any definitive conclusions. Nevertheless, our data suggest that separately targeting the interaction of NEMO with either IKK will disrupt the entire IKK complex.

Role of the NBD in IKKα and IKKβ

In conclusion, we have demonstrated that the interaction of NEMO with both IKKs is necessary for classical NF-κB pathway activation and IKK complex assembly. We have further established that IL-1-induced classical NF-κB activation remains intact in cells lacking functional IKKβ. Tremendous effort has been directed toward developing specific inhibitors of IKKβ (3, 4); however, our findings suggest that this strategy may not effectively block all classical NF-κB activation. Importantly, Lam et al. (31) demonstrated recently that selective inhibition of IKKβ blocks classical pathway-dependent cell survival in a subtype of diffuse large B-cell lymphoma cells only when IKKα is concomitantly ablated. These accumulated findings therefore support efforts to develop NBD-targeting small molecule inhibitors. Such drugs are predicted to inhibit all classical NF-κB activation while retaining both the noncanonical pathway and the NEMO-independent transcriptional regulatory roles of IKKα intact.

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