A Structural Guide to Proteins of the NF- κ B Signaling Module

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The prosurvival transcription factor NF- κ B specifically binds promoter DNA to activate target gene expression. NF- κ B is regulated through interactions with I κ B inhibitor proteins. Active proteolysis of these I κ B proteins is, in turn, under the control of the I κ B kinase complex (IKK). Together, these three molecules form the NF- κ B signaling module. Studies aimed at characterizing the molecular mechanisms of NF- κ B, I κ B, and IKK in terms of their three-dimensional structures have led to a greater understanding of this vital transcription factor system.

NF- κ B is a master transcription factor that responds to diverse cell stimuli by activating the expression of stress response genes. Multiple signals, including cytokines, growth factors, engagement of the T-cell receptor, and bacterial and viral products, induce NF- κ B transcriptional activity (Hayden and Ghosh 2008). A point of convergence for the myriad of NF- κ B inducing signals is the I κ B kinase complex (IKK). Active IKK in turn controls transcription factor NF- κ B by regulating proteolysis of the I κ B inhibitor protein (Fig. 1). This nexus of three factors, IKK, I κ B, and NF- κ B, forms the NF- κ B signaling module—a molecular relay switch mechanism that is conserved across diverse species (Ghosh et al. 1998; Hoffmann et al. 2006). In this article, we introduce the human NF- κ B, I κ B, and IKK proteins, and discuss how they function

from the perspective of their three-dimensional structures.

NF- κ B

Introduction to NF- κ B

NF- κ B was discovered in the laboratory of David Baltimore as a nuclear activity with binding specificity toward a ten-base-pair DNA sequence 5'-GGGACTTCC-3' present within the enhancer of the immunoglobulin κ light chain gene in mature antibody-producing B cells (Sen and Baltimore 1986). The biochemically purified activity was found to be composed of 50 and 65 kilodalton (kDa) subunits. Cloning of the p50 subunit revealed significant amino-acid sequence homology between its amino-terminal 300 amino acids and the oncogene

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T. Huxford and G. Ghosh

Figure 1. The NF-κB signaling module. NF-κB exists in the cytoplasm of resting cells by virtue of its noncovalent association with an IκB inhibitor protein. The IκB kinase (IKK) responds to diverse stimuli by catalyzing the phosphorylation-dependent 26 S proteasome-mediated degradation of complex-associated IκB. Active NF-κB accumulates in the nucleus where it binds with DNA sequence specificity in the promoter regions of target genes and activates their transcription.

from the reticuloendotheliosis virus of turkeys and shared by v-Rel and its cellular proto-oncogene c-Rel. This portion of conserved amino acid sequence was termed the Rel homology region (RHR). As is discussed later, the mRNA responsible for producing the p50 subunit was found to encode a longer precursor protein of 105 kDa in size that possesses the entire p50 amino-acid sequence at its amino-terminal end and its own inhibitor within its carboxy-terminal region. Once the cDNA encoding the p65 subunit (also known as RelA) was sequenced, it was also found to contain an amino-terminal RHR. Two additional NF-κB family subunits, RelB and p52 (the processed product of a longer 100-kDa precursor), were also discovered to harbor the conserved RHR within their amino-terminal regions. These five polypeptides, p50, p65/RelA, c-Rel, p52, and RelB, constitute the entire family of NF-κB subunits encoded by the human genome (Fig. 2A).

Rel Homology Region Structure

The first glimpse at the structure of the RHR was afforded by the successful determination of two x-ray crystal structures of the NF-κB p50:p50 homodimer in complex with related κB DNA (Ghosh et al. 1995; Müller et al. 1995). The structures uncovered a symmetrical protein: DNA complex structure reminiscent of a butterfly with double-stranded DNA comprising the “body” and two-protein subunit “wings” (Fig. 2B,C). These structures revealed a completely novel DNA binding motif in which one entire 300 amino acid RHR from each p50 subunit in the dimer are involved in contacting one whole turn along the major groove of double-stranded DNA (Baltimore and Beg 1995; Müller et al. 1996).

As revealed by the NF-κB p50:DNA complex structures, the RHR consists of two folded domains. The amino-terminal domain (or Rel-N) is approximately 160–210 amino acids in length and exhibits a variant of the immunoglobulin fold. The carboxy-terminal dimerization domain (referred to as Rel-C) spans roughly 100 amino acids and also adopts an immunoglobulin-like fold. The five RHR-containing NF-κB subunits assemble to form various homo- and heterodimer combinations to form active NF-κB transcription factors. The two domains are joined by a short flexible linker approximately 10 amino acids in length. A carboxy-terminal flexible region in which is embedded the nuclear localization signal terminates the conserved RHR portion of the NF-κB subunits. Besides dimerization, this RHR is responsible for sequence-specific DNA binding, nuclear localization, and interaction with IκB proteins.

Outside of the conserved RHR, three NF-κB subunits, p65/RelA, c-Rel, and RelB, contain a transcription activation domain (TAD) at their extreme carboxy-terminal ends. This region, which is poorly understood in protein structural terms, is responsible for the increase in target gene expression that results from induction of NF-κB and, consequently, NF-κB dimers that possess at least one of these subunits function as activators of transcription.

Figure 2. The NF- κ B family. (A) The human genome encodes five polypeptides that assemble in various dimer combinations to form active NF- κ B transcription factors. Each of the subunits contains the Rel homology region (RHR) near its amino terminus. The RHR consists of two folded domains, the amino-terminal domain (NTD) and the dimerization domain (DimD), that are joined by a short flexible linker and a carboxy-terminal flexible region that contains the nuclear localization signal (L). Three of the subunits, p65, c-Rel, and RelB, also contain a transcription activation domain (TAD) at their carboxy-terminal ends. RelB contains a predicted leucine zipper motif (LZ) amino-terminal to its RHR. The NF- κ B subunits p50 and p52 lack transactivation domains and have glycine-rich regions (G). (B) A ribbon diagram representation of the RHR from p50 in its DNA-bound conformation. (C) The NF- κ B p50:p65/RelA heterodimer bound to κ B DNA. (D) Another view of the complex. (E) The NF- κ B p50:p65/RelA heterodimer dimerization domains with key amino acid side chains labeled. (F) κ B DNA from the NF- κ B:DNA complex with key base-contacting amino acid residues labeled.

RelB also contains a predicted leucine zipper motif amino-terminal to its RHR. The NF- κ B subunits p50 and p52 lack transactivation domains. Rather, their extreme carboxy-terminal ends are rich in glycine. Consequently, NF- κ B dimers consisting exclusively of p50 and p52 subunits are capable of nuclear localization and DNA binding but fail to activate target gene expression and, in fact, function both in vitro and in vivo as repressors of transcription (Franzoso et al. 1992).

NF- κ B Dimerization

Assembly of individual NF- κ B subunits into dimers capable of sequence-specific DNA

binding and activating target gene expression is mediated entirely by the dimerization domain. In theory, a total of 15 unique homo- and heterodimers are possible from combinatorial dimerization of the five NF- κ B subunits (Hoffmann et al. 2006). Of these, 12 have been identified in vivo. Three that are not known to exist are the RelB:RelB, RelB:c-Rel, and p52:c-Rel. However, of these three, only RelB homodimer fails to exist in vivo, and the status of the other two dimers is uncertain; they could exist under specialized conditions such as the p65/RelA:RelB heterodimer (Marienfeld et al. 2003).

The homo- and heterotropic interactions between NF- κ B family subunits follow the

T. Huxford and G. Ghosh

central dogma of all protein–protein complexes: A set of amino acid residues from each polypeptide participate in direct contacts with one another, forming the interface, whereas a different set of residues present outside of the interface indirectly affects the stability of the interface by modulating the local environment. One of the most stable NF- κ B dimer interfaces is created by the p50:p65/RelA heterodimer (Fig. 2D). The p50:p65/RelA heterodimer assembles with a significantly greater stability than either of the respective p50:p50 or p65/RelA:p65/RelA homodimers (Huang et al. 1997). Differences in the amino acid sequences of p50 and p65/RelA at two positions partially explain the variation in dimerization stability observed in the three dimers. The positions of Tyr-269 and Asp-256 in p50 are occupied by Phe-213 and Asn-200, respectively, in p65. Within the heterodimer, the Asp and Asn approach one another symmetrically at the interface and stabilize the heterodimer through formation of a highly stable hydrogen bond. In contrast, in the homodimers, the juxtaposition of Asp-Asp and Asn-Asn at the interface are detrimental to dimer stability. Similarly, the hydroxyl group on Tyr-269 of p50 hydrogen bonds at the dimer interface, contributing to stabilization of both the p50:p50 homodimer as well as the p50:p65/RelA heterodimer. The substitution of Phe at this position serves to weaken the p65/RelA:p65/RelA homodimer relative to the other two. Taken together, these observations help to explain why the p65/RelA:p65/RelA homodimer forms with lower stability than does the p50:p50 homodimer and why the p50:p65/RelA heterodimer is more stable than both homodimers.

However, direct contact between complementary amino acids at the interface fails to completely explain the observed trends of NF- κ B subunit dimerization. Mutation of non-interfacial residues contained within the dimerization domain has been shown to modulate dimerization. For example, changing p65/RelA Cys-216 to Ala affects homodimer formation (Ganchi et al. 1993). The role of noninterfacial amino acid residues in dimerization is most strikingly illustrated in the case of RelB. All

interfacial residues in RelB are either identical or homologous to those of other NF- κ B subunits. And yet, RelB assembles into a completely unique domain-swapped homodimer (Huang et al. 2005). Domain swapping occurs as a consequence of the destabilization of the folded RelB dimerization domain, suggesting that domain stability is an important determinant of protein–protein interaction. In cells, decreased folding stability in both the amino-terminal and dimerization domains contributes to its degradation by the proteasome, which explains why the RelB homodimer does not exist in vivo (Marienfeld et al. 2001).

Post-translational modification may also play a role in modulating dimerization propensity. One study has shown that RelB forms a dimer with a p65/RelA that is phosphorylated at position Ser-276 (Jacque et al. 2005). This serine is located in a loop within the dimerization domain, projected away from the dimer interface. It is unclear as to how phosphorylation of this serine positively impacts p65/RelA:RelB heterodimer formation. One explanation could be that the increased negative charge may indirectly modulate dimer-forming residues of both p65/RelA and RelB.

NF- κ B Recognition of κ B DNA

X-ray structures of NF- κ B in complex with κ B DNA revealed a new mode of DNA recognition wherein a dimer composed of the RHR of two NF- κ B subunits intimately contacts double-stranded DNA within the major groove through one complete turn (Ghosh et al. 1995; Müller et al. 1995). NF- κ B employs both its amino-terminal and dimerization domains to encircle its target DNA. DNA contacts are mediated by amino acids emanating from loops that connect β -strand elements of secondary protein structure. The p50:p50 homodimer structures revealed that this NF- κ B subunit employs its amino acids His-66, Arg-58, and Arg-56 to contact three guanine nucleotide bases at the extreme 5' ends of its consensus κ B DNA. However, X-ray analyses of NF- κ B p65/RelA:p65/RelA homodimers bound to κ B DNA revealed that, when bound to a canonical

10-base-pair κ B DNA, one p65/RelA subunit contacts DNA in an analogous manner as observed in the p50 homodimer structures, whereas the second p65/RelA subunit significantly repositions its entire amino-terminal domain to mediate interactions with the DNA backbone (Chen et al. 1998b). Such a binding mode, which is afforded by flexibility in the short linker region that connects the two structured domains of the RHR, preserves binding affinity at the cost of fewer contacts to DNA bases. A close inspection of the homodimer:DNA complex structures leads to the suggestion that a homodimer of p50 might optimally bind to an 11-base-pair sequence composed of two 5'-GGGPyN half sites bracketing a central A:T base pair. In contrast, the NF- κ B p65/RelA homodimer optimally recognizes a nine-base-pair target sequence containing two 5'-GGPyN half sites and a central A:T base pair. Determination of a second X-ray structure of NF- κ B p65/RelA homodimer in complex with the κ B DNA sequence from the promoter of IL-8 (5'-GGAA T TTCC-3') confirmed this hypothesis (Chen et al. 2000).

The lessons learned from structural analyses of p50 and p65/RelA homodimers bound to diverse κ B DNA sequences suggested that the canonical NF- κ B p50:p65/RelA heterodimer might recognize its 10-base-pair target sequence with the p50 subunit binding specifically to a 5'-GGGPyN half site, whereas the p65/RelA subunit binds to a 5'-GGPyN half site separated from the p50 site by one A:T base pair. X-ray structure determination of a p50:p65/RelA RHR heterodimer bound to κ B DNA from the original κ light chain gene promoter, which is identical to a κ B sequence that is present in the promoter of HIV genome (5'-GGGAC T TTCC-3'), served to confirm this speculation (Chen et al. 1998a). Additional crystal structures of NF- κ B p50:p65/RelA heterodimer bound to different κ B DNA further support the basic rules of DNA half-site recognition developed from the homodimer:DNA structures (Berkowitz et al. 2002; Escalante et al. 2002).

X-ray crystal structures of several additional NF- κ B:DNA complexes have now been

determined (Cramer et al. 1997; Cramer et al. 1999; Moorthy et al. 2007; Panne et al. 2007; Fusco et al. 2009). Taken together, these structures suggest a model for how NF- κ B dimers recognize κ B DNA that contain significant deviations from the consensus sequence. Changes of κ B DNA sequence can be of two types. In the first, alteration occurs within the G:C base pairs that occupy positions on the outside of the κ B DNA and that are directly contacted by the amino-terminal domain. An example is the altered κ B sequence 5'-GGGAC T TTTC-3' (change from immunoglobulin κ B sequence is underlined). Because of the loss of an important C:G base pair, the NF- κ B p65/RelA subunit conformation when bound to TTCC-3' is drastically different than to TTTC-3'. The flexible linker region and modular domain architecture within the RHR allows the amino-terminal domain to reposition itself and bind the DNA backbone to accommodate such variations in κ B DNA sequence. A second type of κ B DNA sequence alteration involves changes within the central five base pairs. Of these, the central A:T is not directly contacted by the protein and the others are recognized nonspecifically through van der Waals contacts by p50 Tyr-59 (Tyr-36 in p65/RelA). The third base pair from the 5'-end or its symmetric pair are less sensitive to change, as illustrated by a comparison of immunoglobulin κ B and IFN- β κ B sequences (5'-GGGAC T TTCC-3' and 5'-GGGAA T TTCC-3', respectively). The C:G to A:T base-pair change does not affect DNA recognition by the protein but may influence stability of binding because of the more rigid DNA structure of IFN- β κ B sites. In this second case, overall conformations of the protein:DNA complexes are similar, but binding affinity could differ significantly.

Molecular dynamics simulations have revealed intriguing structural transitions of a 20-base-pair DNA containing the κ B site of IL-2 promoter (AGAA A TTCC). The central A:T base pair (underlined) undergoes cross-stand stacking and the central A:T base pair flips out of the DNA axis (Mura and McCammon 2008). This dynamic behavior of the DNA suggests a highly complex mechanism of DNA

T. Huxford and G. Ghosh

recognition by the NF- κ B dimers, in which DNA sequence variations may play a significant role in the recognition process. In general, one can confidently say that the sequences at the center of a κ B sequence may profoundly affect the binding affinity and specificity.

I κ B

Introduction to I κ B

Almost immediately on detecting NF- κ B in immune cells, researchers discovered that a latent κ B DNA binding activity was present in the cytoplasm of all resting cells and that this pool of NF- κ B could be activated by treatment of cell lysates with the weak detergent deoxycholate (Baeuerle and Baltimore 1988a). This suggested that noncovalent interaction with an inhibitor protein was responsible for maintaining NF- κ B in an inactive state. Purification of the inhibitor activity led to the cloning of the inhibitory proteins I κ B α and I κ B β (Baeuerle and Baltimore 1988b; Thompson et al. 1995). A third I κ B gene was later identified by sequence homology in an EST database and was called I κ B ϵ (Li and Nabel 1997; Simeonidis et al.

1997; Whiteside et al. 1997). Subsequent experiments demonstrated that it also exhibits NF- κ B inhibitory activity.

Classical I κ B Sequence and Structure

Both I κ B α and I κ B β , as well as the more recently discovered I κ B ϵ , contain a central ankyrin repeat domain (ARD) that contains six ankyrin repeats (Fig. 3A). The ankyrin repeat (ANK) is a roughly 33-amino-acid consensus amino acid sequence that appears in multiple copies in numerous proteins (Fig. 3B) (Sedgwick and Smerdon 1999). Ankyrin repeats are part of a greater superfamily of helical repeat motifs, which include HEAT repeats, armadillo repeats, and leucine-rich repeats, and are common to proteins involved in protein–protein interactions (Groves and Barford 1999). At their amino-terminal ends, classical I κ B proteins contain a sequence of amino acids that do not adopt a folded structure in solution (Jaffray et al. 1995). Contained within this signal response region are the conserved serine sites of phosphorylation by IKK. Roughly 10 amino acids amino-terminal to this pair of serines

Figure 3. The family of human I κ B proteins. (A) I κ B proteins are classified as in text. Classical I κ B proteins possess ankyrin repeats (ANK) flanked by an amino-terminal signal response region and carboxy terminal PEST region. The signal response regions contain sites of phosphorylation by IKK (S), ubiquitination (K), and nuclear export (E). The NF- κ B precursors serve as I κ B proteins as well as the source of the mature p50 and p52 NF- κ B subunits. (B) Ribbon diagram of the I κ B α structure from the NF- κ B:I κ B α complex crystal structure. Individual ankyrin repeats are numbered, ANK 4 is colored magenta, and the PEST region is labeled. (C) Ribbon diagram of the NF- κ B:I κ B α complex. (D) Another view of the complex.

reside conserved lysine amino acid sites of poly-ubiquitination. This amino-terminal region of I κ B α also contains a functional nuclear export sequence (Johnson et al. 1999; Huang et al. 2000). It is not masked on binding to NF- κ B and contributes to the observed cytoplasmic localization of NF- κ B:I κ B α complexes. Neither I κ B β nor I κ B ϵ possess this inherent nuclear export potential. Consequently, stable NF- κ B:I κ B β complexes reside stably either in the cytoplasm or the nucleus, whereas I κ B ϵ appears to function as a negative feedback regulator of cytoplasmic NF- κ B (Malek et al. 2001; Tam and Sen 2001; Kearns et al. 2006). At their carboxy-terminal ends, the three classical I κ B proteins contain a short sequence rich in the amino acids proline, glutamic acid, serine, and threonine. This so-called PEST region is common to many proteins that, like I κ B, display rapid turnover in cells (Rogers et al. 1986; Pando and Verma 2000). The PEST region of I κ B α , however, is also required for its ability to disrupt preformed NF- κ B:DNA complexes (Ernst et al. 1995).

I κ B Interactions with NF- κ B

The X-ray structure of I κ B α in complex with the NF- κ B p50:p65/RelA heterodimer was determined independently by two separate laboratories in 1998 (Huxford et al. 1998; Jacobs and Harrison 1998). Both groups relied on a similar strategy of removing the signal response region of I κ B α and the amino-terminal domain of the p50 subunit to stabilize the conformationally dynamic complex for cocrystallization (Huxford et al. 2000). The structure reveals how I κ B α uses its entire ankyrin repeat-containing domain as well as its carboxy-terminal PEST sequence to mediate an extensive protein–protein interface of roughly 4300 Å² (Fig. 3C,D). The carboxy-terminal 30 amino acids from the NF- κ B p65/RelA subunit RHR, which were disordered in NF- κ B:DNA complex structures, adopt an ordered helical structure that contacts the first two ankyrin repeats and forms significant hydrophobic contacts with the amino-terminal face of the I κ B α ankyrin repeat stack. This interaction

masks the p65/RelA nuclear localization signal. Ankyrin repeats three through five participate in multiple van der Waals contacts with one surface of the p50:p65/RelA heterodimer dimerization domains. The sixth ankyrin repeat and PEST region of I κ B α present a vast acidic patch, which opposes the largely positively charged DNA binding surfaces of the p65/RelA amino-terminal domain. As a consequence of this electrostatic interaction, the p65/RelA amino-terminal domain occupies a position relative to the dimerization domain that is rotated roughly 180° and translated 40 Å when compared with its DNA bound structures. The transition of p65/RelA to the conformation observed in the NF- κ B:I κ B complex does not disrupt the amino-terminal domain structure and is afforded entirely by the flexible linker region that connects the amino-terminal and dimerization domains. The structure of a similar construct of I κ B β bound to the dimerization domain from the NF- κ B p65/RelA:p65/RelA homodimer suggests that I κ B β uses a similar strategy in binding to NF- κ B, although it relies less on interactions with the p65/RelA amino-terminal domain for complex stability (Malek et al. 2003).

I κ B α Dynamics

Protein dynamics, or the rates with which a protein exchanges between quasi-stable folded states, is an extremely important aspect of protein structure and function. Several independent lines of investigation, including thermal and chemical denaturation, computer simulations, NMR spectroscopy, and failed crystallization attempts, have led to the conclusion that the I κ B α protein exhibits a high degree of structural dynamics in solution (Huxford et al. 2000; Pando and Verma 2000; Croy et al. 2004; Bergqvist et al. 2006). This runs counter to the data that have emerged from protein engineering studies that clearly show that ankyrin repeat proteins designed after consensus sequences or those that appear in nature are extremely stably folded (Binz et al. 2003; Binz et al. 2004). I κ B α has, therefore, evolved as an inherently unstable ankyrin repeat-containing

T. Huxford and G. Ghosh

protein. The consequences of this are twofold. First of all, free I κ B α is easily degraded in cells. This signal-independent degradation involves the 20S proteasome and the carboxy-terminal PEST of I κ B α (Mathes et al. 2008). Moreover, recent computational modeling of the NF- κ B pathway through a systems biology approach has confirmed that regulation of NF- κ B activation can be controlled by small changes in the rate of degradation of a constitutively expressed free cytoplasmic I κ B α (O'Dea et al. 2007). On binding to NF- κ B, the dynamic I κ B α ankyrin repeat fold becomes stable and the PEST region protein turnover signal sequence is adopted as a DNA-inhibitory functional element (Sue et al. 2008). Degradation of I κ B α is shifted from the steady state to a signal-dependent pathway that requires phosphorylation within the flexible amino-terminal signal response region. It is likely that the inherent folding instability of I κ B α contributes to it being targeted by the proteasome, whereas NF- κ B, which is composed of two stably folded domains and with which I κ B α is associated in a complex at subnanomolar dissociation binding constant, remains intact.

Nonclassical I κ B Proteins

Through the efforts of investigators attempting to understand regulation of NF- κ B, a more diverse family of I κ B proteins has emerged. Proteins of the I κ B family are all linked by the fact that they contain ankyrin repeats and interact with NF- κ B subunits to affect gene expression. The classical I κ B proteins, I κ B α , I κ B β , and I κ B ϵ , have been described. A second class of I κ B proteins is represented by the NF- κ B precursor proteins p105 and p100 (Basak et al. 2007). These two proteins act both as precursors of NF- κ B p50 and p52 subunits, respectively, and as inhibitors of NF- κ B (Fig. 3A). The NF- κ B precursor proteins are responsible for inhibiting nearly half of the NF- κ B in resting cells. However, unlike classical I κ B proteins that inhibit NF- κ B by forming 1:1 complexes, these NF- κ B precursors participate in large multiprotein assemblies, wherein more than one NF- κ B dimer can bind to multiple

p100 and/or p105 subunits. The assembly of p100 and p105 into larger complexes is mediated by an oligomerization domain located immediately amino-terminal to their ankyrin-repeat domains. This assembly is heterogeneous, i.e., several different NF- κ B subunits can be inhibited in a single inhibitory complex (Savinova et al. 2009). Therefore, stimulus-specific degradation of p100 or p105 can in principle release different NF- κ B dimers. Together, these dimers exhibit a much broader spectrum of gene regulatory activities (Savinova et al. 2009, Shih et al. 2009). The large heterogeneous NF- κ B inhibitory complex assemblies have been dubbed NF- κ Bsomes to distinguish them from the smaller NF- κ B inhibitory complexes formed by I κ B α , - β , and - ϵ .

Together with I κ B α and I κ B ϵ , the NF- κ B precursors are targets of NF- κ B-driven transcription. Newly synthesized I κ B serve to block NF- κ B activity postinduction. This negative feedback regulation of NF- κ B by I κ B proteins is critical for control of inflammation and other diseases (Hoffmann et al. 2002).

Nuclear I κ B Proteins

A third, entirely different class of I κ B is represented by the proteins Bcl-3, I κ B ζ /MAIL, and I κ BNS (Fig. 3A). Bcl-3 was cloned as a consequence to its proximity to a breakpoint mutation in some leukemias (Ohno et al. 1990). I κ B ζ was discovered in a screen of genes that displayed increased expression after induction of immune cells with bacterial lipopolysaccharide (LPS) or the inflammatory cytokine interleukin-1 (IL-1) (Kitamura et al. 2000; Haruta et al. 2001; Yamazaki et al. 2001). I κ BNS was identified as a gene that is expressed in T cells during negative selection (Fiorini et al. 2002). Unlike classical I κ B, these proteins do not contain amino-terminal signal-dependent phosphorylation sites or carboxy-terminal PEST regions. Their classification as “nuclear I κ B” derives from the fact that they contain ankyrin repeats, bind NF- κ B subunits, and concentrate within the nucleus when expressed in cells (Michel et al. 2001).

Each of the three nuclear I κ B proteins are themselves the products of NF- κ B-dependent genes (Eto et al. 2003; Ge et al. 2003; Hirotani et al. 2005). They bind to NF- κ B, but, whereas the classical I κ B proteins prefer dimers that possess at least one p65/RelA or c-Rel subunit, or nonclassical p105 and p100 binds all NF- κ B subunits, nuclear I κ Bs bind specifically to homodimers of p50 (Hatada et al. 1992; Yamazaki et al. 2001; Trinh et al. 2008). Finally, association of nuclear I κ B proteins with nuclear NF- κ B can have diverse but important consequences on gene expression (Franzoso et al. 1992; Muta et al. 2003; Hirotani et al. 2005; Motoyama et al. 2005; Riemann et al. 2005). In peritoneal macrophages derived from mice lacking the gene encoding I κ B ζ , for example, a complete inability to produce the NF- κ B-dependent cytokine interleukin-6 (IL-6) in response to LPS treatment was observed (Yamamoto et al. 2004). As IL-6 is a gene that is expressed in a later phase of NF- κ B induction, it is apparent that the early induction and nuclear accumulation of I κ B ζ plays a vital role in the LPS-dependent expression of this pluripotent cytokine.

IKK

Introduction to IKK

In a thrilling conclusion to a search to identify an enzymatic activity that was capable of phosphorylating the two serine amino acids of the signal response region of I κ B α , researchers from three labs reported in 1997 the I κ B Kinase complex (IKK) (DiDonato et al. 1997; Mercurio et al. 1997; Regnier et al. 1997). This IKK was purified from cytokine-induced HeLa cells and exhibited an apparent molecular mass of 700–900 kDa. Microsequencing revealed two related kinase domain-containing subunits, referred to as IKK α and IKK β (or alternatively as IKK1 and IKK2, respectively) (DiDonato et al. 1997; Zandi et al. 1997). These exhibit molecular masses of 85 and 87 kDa, respectively, and display 50% identity at the amino-acid level. The IKK α subunit was recognized to be the

same protein that was originally cloned in 1995 as CHUK, a kinase with homology to the helix-loop-helix transcription factors, and IKK β was subsequently identified as a hit in a yeast two-hybrid screen with NIK, a kinase suspected to function upstream of IKK, as bait (Connelly and Marcu 1995; Regnier et al. 1997; Woronicz et al. 1997). A third subunit, known as IKK γ (also known as NEMO), was also identified as a 49 kDa member of the IKK complex (Rothwarf et al. 1998; Yamaoka et al. 1998; Li et al. 1999b; Mercurio et al. 1999).

The function of IKK1/IKK α in cells is somewhat exotic and continues to be elucidated (Hu et al. 2001; Senftleben et al. 2001; Anest et al. 2003; Yamamoto et al. 2003; Sil et al. 2004; Lawrence et al. 2005). However, multiple studies including mouse knockouts have revealed that IKK2/IKK β is responsible for phosphorylating I κ B in response to NF- κ B-inducing signals (Li et al. 1999a; Li et al. 1999c; Tanaka et al. 1999). Because of its role as the primary inducer of the NF- κ B transcription factor, IKK2/IKK β plays the critical role in promoting inflammation and cell survival in response to proinflammatory stimuli. The NEMO/IKK γ subunit does not possess any kinase domain or enzymatic activity and instead it acts as an adapter subunit that links the catalytic subunits to receptor proximal signaling molecules. Mouse knockout studies clearly reveal that proper NF- κ B signaling through IKK is not possible without this subunit (Makris et al. 2000; Rudolph et al. 2000; Schmidt-Supprian et al. 2000).

IKK Catalytic Subunit Domain Organization

The IKK1/IKK α and IKK2/IKK β subunits exhibit an uncommon domain organization (Fig. 4A). Their first roughly 300 amino acids contain a clearly recognizable kinase domain. This is followed by a short region that exhibits distant homology to ubiquitin (Ikeda et al. 2007). A central region contains a leucine zipper motif followed by a region with slight homology to the helix-loop-helix transcription factors. This is followed by a serine-rich region. Finally, the carboxy-terminal element of IKK β

T. Huxford and G. Ghosh

Figure 4. Subunits of the human IKK complex. (A) Domain organization of IKK subunits. Catalytic subunits contain a kinase domain (KD), ubiquitin-like domain (U), leucine zipper (L), helix-loop-helix (H), serine-rich (S), and NEMO-binding motif (N). The NEMO/IKKγ subunit contains two predicted coiled-coil motifs (CC1 and -2), a leucine zipper (L), and a carboxy-terminal zinc-finger (ZF). (B) Ribbon diagram of the IKK2/IKKβ:NEMO/IKKγ complex. Individual polypeptides are labeled as well as some of the conserved hydrophobic amino acid side chains from IKK2/IKKβ. (C) Ribbon diagram of the NEMO/IKKγ:di-ubiquitin complex. (D) The NEMO/IKKγ carboxy-terminal zinc-finger motif structure.

has been shown to interact directly with NEMO/IKKγ (May et al. 2000; May et al. 2002). A genome-wide analysis has identified a small clade of proteins with domain organization reminiscent of the catalytic IKK1/IKKα and IKK2/IKKβ subunits (Manning et al. 2002). This kinase subgroup includes the proteins IKKε and TBK1/NAK. Both have been characterized as upstream modulators of IKK activity and are, therefore, both structurally and functionally related to the catalytic IKK subunits (Tojima et al. 2000; Peters and Maniatis 2001).

With the exception of the extreme carboxy-terminal NEMO/IKKγ-interacting motif, the structures of IKK1/IKKα and IKK2/IKKβ are unknown. Although it is clear that the kinase domain of IKK2/IKKβ is necessary for catalyzing phospho-transfer, regions of the protein outside of this domain are necessary for directing specificity to the amino-terminal serines of IκBα. Deletion of the leucine zipper and helix-loop-helix has been shown to yield a mutant enzyme that, although it is capable of catalyzing phospho-transfer to IκBα, fails to recognize the amino-terminal serines required for NF-κB activation in response to inflammatory signaling (Shaul et al. 2008).

NEMO/IKKγ Domain Organization

Structural interest in NEMO/IKKγ arises from its lack of a kinase domain or, for that matter, homology to any other protein of known structure. Secondary-structure prediction methods suggest that NEMO/IKKγ is a mostly helical protein with two signature coiled coil (CC) elements and a leucine zipper motif in the middle are flanked by a helical dimerization domain near the amino-terminal end and a Zn-finger motif at the carboxyl terminus. Three-dimensional structures of several fragments of the NEMO/IKKγ subunit either as free polypeptides or bound to ligands have recently been elucidated. These NEMO/IKKγ structures allow one to envision that with the exception of the very ends, NEMO/IKKγ consists of long helices that are punctuated by short unstructured regions. These helical segments wrap around each other forming a long coiled-coil dimer with fraying of the monomers at each end. The carboxy-terminal end contains a Zn-finger motif and the 40-residue long amino terminus is likely to be flexible and unstructured.

The NEMO/IKKγ fragment structures suggest how IKKγ might be involved in

cellular signaling. NEMO/IKK γ was previously thought to form a multimer with different segments shown to form dimers, trimers, or tetramers (Agou et al. 2002; Tegethoff et al. 2003; Marienfeld et al. 2006; Drew et al. 2007; Herscovitch et al. 2008). With new structural information, it now appears that previous conclusions might not be accurate. It is not surprising as flexible coiled-coil motifs migrate through the gel filtration beads differently than the stably folded globular proteins of identical mass.

Two Distinct IKK Activation Pathways

The IKK1/IKK α and IKK2/IKK β subunits are activated through distinct signaling pathways (Senftleben et al. 2001). These pathways are activated by two distinct sets of stimuli. The canonical pathway is triggered by LPS or inflammatory cytokines TNF- α or IL-1 and signals through IKK2/IKK β to activate p65/RelA and c-Rel dimers through the degradation of I κ B α , I κ B β , I κ B ϵ , and p105. The noncanonical pathway results from BAFF, LT- β , and CD40 signaling through IKK α , leading to activation of the p52:RelB heterodimer through processing of p100 into p52 (Ghosh and Karin 2002). Activation of both pathways requires interactions of signaling molecules through specific poly-ubiquitin moieties that are covalently linked to some of these molecules. In addition, upstream protein kinases are also essential for activation of both kinases. The major distinction between these two pathways is the involvement of a single upstream kinase, known as NF- κ B inducing kinase (NIK), to activate IKK α , whereas multiple kinases can activate IKK β . IKK α and IKK β are both present in the same particle in vivo. One puzzling question, however, is whether the different pathways target these two distinct subunits in a single complex or if there is a separate pool of IKK α present in cells that is activated by the noncanonical pathway. For the canonical pathway, however, the IKK β subunit of the heterodimer must be activated. The role for IKK α in canonical signaling is unclear. The presence of the I κ B kinases in multiple signaling pathways suggests

that they are capable of participating in diverse signaling complexes throughout the cell.

IKK Complex Oligomerization

Although several combinations of the IKK1/IKK α , IKK2/IKK β , and NEMO/IKK γ subunits have been proposed to account for the original 700–900 kDa complex, no successful reconstitution from purified components has resulted in an active complex of this size. Therefore, 10 years after isolation of the complex, even the oligomerization state of the IKK complex remains unclear. Given the presence of multiple conserved elements that can potentially mediate homo- and heteromeric interactions between subunits, there exist many possibilities. Early attempts to determine the arrangement of subunits in the complex led to the model that, independent of its NEMO/IKK γ scaffolding protein, IKK1/IKK α and IKK2/IKK β are capable of forming stable homo- or heterodimers (Zandi et al. 1998). More recent work with human IKK2/IKK β purified in milligram quantities from recombinant baculovirus-infected sf9 insect cells has shown that the full length subunit purifies as a tetramer. Removal of the carboxy-terminal 100 amino acids containing the serine-rich and NEMO/IKK γ -binding regions results in a dimeric enzyme, whereas removal of the entire carboxy-terminal portion beginning at the leucine zipper renders the kinase monomeric. Interestingly, it was observed that although this monomeric IKK2/IKK β kinase domain remains catalytically active, it fails to specifically phosphorylate the signal response region of I κ B α and instead catalyzes phosphorylation of the PEST (Shaul et al. 2008). The presence of multiple domains linked to flexibility in all three subunits shows why it is difficult to structurally characterize functional IKK complexes.

The Emerging Structure of NEMO/IKK γ

A working structural model of the NEMO/IKK γ subunit has begun to take shape with the recent successful determination of several X-ray and NMR structures of discrete functional

T. Huxford and G. Ghosh

portions. The structure of an IKK2/IKK β polypeptide bound the amino-terminal helical region (residues 44 to 111) of NEMO/IKK γ revealed the complex forms a parallel four-helix bundle where two IKK2/IKK β peptides associate with the NEMO/IKK γ dimer (Rushe et al. 2008). The IKK2/IKK β peptides appear to fold on binding to the NEMO/IKK γ -dimer scaffold, making several contacts (Fig. 4B). The affinity of interaction between NEMO/IKK γ and IKK2/IKK β subcomplex is high (K_D is in low nanomolar range) and requires a large array of contacts. However, few of the hydrophobic residues in IKK2/IKK β that are observed to line the hydrophobic pocket formed by the NEMO/IKK γ dimer are required for complex formation. Although no clear experimental data is available, the IKK:IKK γ complexes might exist as a 2:2 complex, where a dimeric NEMO/IKK γ binds to an IKK dimer. The amino-terminal helical dimerization domain of NEMO/IKK γ interacts with both the IKK1/IKK α and IKK2/IKK β carboxy-terminal peptides, forming a 1:1:2 (IKK1/IKK α :IKK2/IKK β :NEMO/IKK γ) IKK complex. This is possibly the basal state of the IKK complex in cells.

Several reports demonstrated an interaction between poly-ubiquitin chains and the CC2-LZ region of NEMO/IKK γ (Lo et al. 2009). Although it was previously thought that the poly-ubiquitin chain is linked through lysine 63 and glycine 76 of ubiquitin, experiments have now shown that linear ubiquitin chains bind to NEMO/IKK γ with 100-fold tighter binding affinity than do K63-linked chains. The X-ray structure of the complex between the CC2-LZ of NEMO/IKK γ and a linear di-ubiquitin motif has been recently elucidated (Rahighi et al. 2009). Two di-ubiquitin motifs are bound to two chains of the LZ motif (Fig. 4C). It is striking that this central region of NEMO/IKK γ also exhibits the elongated coiled-coil motif that was observed in the previously determined IKK2/IKK β :NEMO/IKK γ complex as well as in the X-ray structure of a complex between the carboxy-terminal helical region of NEMO/IKK γ and the viral protein vFLIP from Kaposi's sarcoma virus (Bagnris et al. 2008).

The extreme carboxy-terminal region of NEMO/IKK γ adopts a CCHC-type zinc-finger motif. The solution structure of this region has been determined by multidimensional NMR spectroscopy (Cordier et al. 2008). Its structure adopts the familiar fold of a zinc-finger motif (Fig. 4d). Furthermore, this motif has been characterized as a ubiquitin-binding motif required for NF- κ B signaling in response to TNF- α (Cordier et al. 2009).

When taken together, the NEMO/IKK γ substructures suggest that this subunit adopts an elongated helical structure. Two long helices bind one another through extended coiled-coil interactions to form the docking site for a pair of kinase subunits at the amino terminus, binding sites for di-ubiquitin and other proteins throughout the central region, and a pair of zinc-finger ubiquitin binding sites at the carboxyl terminus. This modular arrangement of docking sites for diverse proteins seems appropriate for a subunit that is thought to function principally as a scaffold for inducible signaling. Although significant structural work on IKK complexes remains to be carried out, this oddly elongated NEMO/IKK γ model serves to explain some of the complications associated with early studies that relied primarily on size exclusion chromatography for characterization of complex size and subunit stoichiometry.

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Proteins of the NF- κ B Signaling Module

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T. Huxford and G. Ghosh

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