

Dual-specific autophosphorylation of kinase IKK2 enables phosphorylation of substrate I κ B α through a phosphoenzyme intermediate

Prateeka Borar, Tapan Biswas, Ankur Chaudhuri, Tom Huxford, Saikat Chakrabarti, Gourisankar Ghosh, Smarajit Polley 

Department of Biological Sciences, Bose Institute, Kolkata, India • Department of Chemistry and Biochemistry, University of California San Diego, La Jolla, USA • Structural Biology and Bioinformatics Division, CSIR-Indian Institute of Chemical Biology, Kolkata, India • Department of Chemistry and Biochemistry, San Diego State University, San Diego, USA

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Abstract

Rapid and high-fidelity phosphorylation of two serines (S32 and S36) of I κ B α by a prototype Ser/Thr kinase IKK2 is critical for fruitful canonical NF- κ B activation. Here, we report that IKK2 is a dual specificity Ser/Thr kinase that autophosphorylates itself at tyrosine residues in addition to its activation loop serines. Mutation of one such tyrosine, Y169, located in proximity to the active site, to phenylalanine, renders IKK2 inactive for phosphorylation of S32 of I κ B α . Surprisingly, auto-phosphorylated IKK2 relayed phosphate group(s) to I κ B α without ATP when ADP is present. We also observed that mutation of K44, an ATP-binding lysine conserved in all protein kinases, to methionine renders IKK2 inactive towards specific phosphorylation of S32 or S36 of I κ B α , but not non-specific substrates. These observations highlight an unusual evolution of IKK2, in which autophosphorylation of tyrosine(s) in the activation loop and the invariant ATP-binding K44 residue define its signal-responsive substrate specificity ensuring the fidelity of NF- κ B activation.

eLife assessment

This study presents **fundamental** findings that could redefine the specificity and mechanism of action of the well-studied Ser/Thr kinase IKK2 (a subunit of inhibitor of nuclear factor kappa-B kinase (IκB) that propagates cellular response to inflammation). **Solid** evidence supports the claim that IKK2 exhibits dual specificity that allows tyrosine autophosphorylation and the authors further show that auto-phosphorylated IKK2 is involved in an unanticipated relay mechanism that transfers phosphate from an IKK2 tyrosine onto the IκBα substrate. These are potentially provocative results but open questions remain due to the nature of the in vitro assays and questions about protein purity and identity. Nevertheless, the findings are a starting point for follow-up studies to confirm the unexpected mechanism and further pursue functional significance.

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Introduction

Protein kinases bestow novel identity to their substrates by adding phosphate moieties at the site of modification that often play key regulatory roles in a diverse array of signaling events. For example, inflammatory or pathogenic signals trigger precisely regulated induction of the transcription factor NF-κB in metazoans (Hoffmann and Baltimore, 2006) that depends upon the upstream activation of the Inhibitor of κB Kinase-complex (IKK-complex), comprising two catalytic subunits, IKK1 (also known as IKKα) and IKK2 (or IKKβ), and an essential regulatory scaffolding protein, NEMO (IKKγ) (DiDonato et al., 1997; Rothwarf et al., 1998; Zandi et al., 1997). IKK2 activity is maintained at a very low basal level in resting cells (Ghosh and Karin, 2002; Hacker and Karin, 2006; Hinz and Scheidereit, 2014; Liu et al., 2012). Upon activation, IKK2 phosphorylates the N-terminus of IκBα (Inhibitor of κB) protein specifically at two Serine residues (S32 and S36) marking it for ubiquitinylation-mediated proteasomal degradation (Figure 1A). Signal-responsive phosphorylation of IκBα refers to the phosphorylation of S32 and S36 by the IKK-activity. Degradation of IκBα liberates NF-κB to execute its gene expression program (Hayden and Ghosh, 2008; Hinz and Scheidereit, 2014; Karin and Ben-Neriah, 2000; Scheidereit, 2006). Mutation of both S32 and S36 to alanine residues converts IκBα into a non-phosphorylatable super-repressor of NF-κB (Brown et al., 1995; Lin et al., 1995). Furthermore, IKK possesses exquisite specificity for these two serines, and it fails to phosphorylate IκBα when S32 and S36 are mutated even to another phosphorylatable residue, threonine (DiDonato et al., 1997).

IKK2 is an essential Ser/Thr kinase that ensures the activation of a particular signaling cascade - NF-κB, with high fidelity despite its reported pleiotropy in other contexts (Antonia et al., 2021; Schröfelbauer et al., 2012; Schröfelbauer and Hoffmann, 2011). IKK2 is a multidomain protein consisting of a kinase domain (KD) followed by a Ubiquitin-like domain (ULD) and the scaffold dimerization domain (SDD) (Figure 1B). IKK2 toggles between inactive and active states based on the phosphorylation status of two serine residues (S177 and S181; S176 and S180 in the case of IKK1) within its activation loop (activation loop) in the KD (Huse and Kuriyan, 2002; Liu et al., 2013; Polley et al., 2013; Xu et al., 2011). Replacement of these two serines with phosphomimetic glutamate residues (S177E, S181E; henceforth EE) renders the kinase constitutively active. IKK2 is reported to be primed through phosphorylation at S177 by the upstream kinase TAK1 that leads to auto-phosphorylation at S181, thus rendering the kinase fully active in cells (Zhang et al., 2014). However, oligomerization upon association with NEMO, in the presence of linear or Lys-

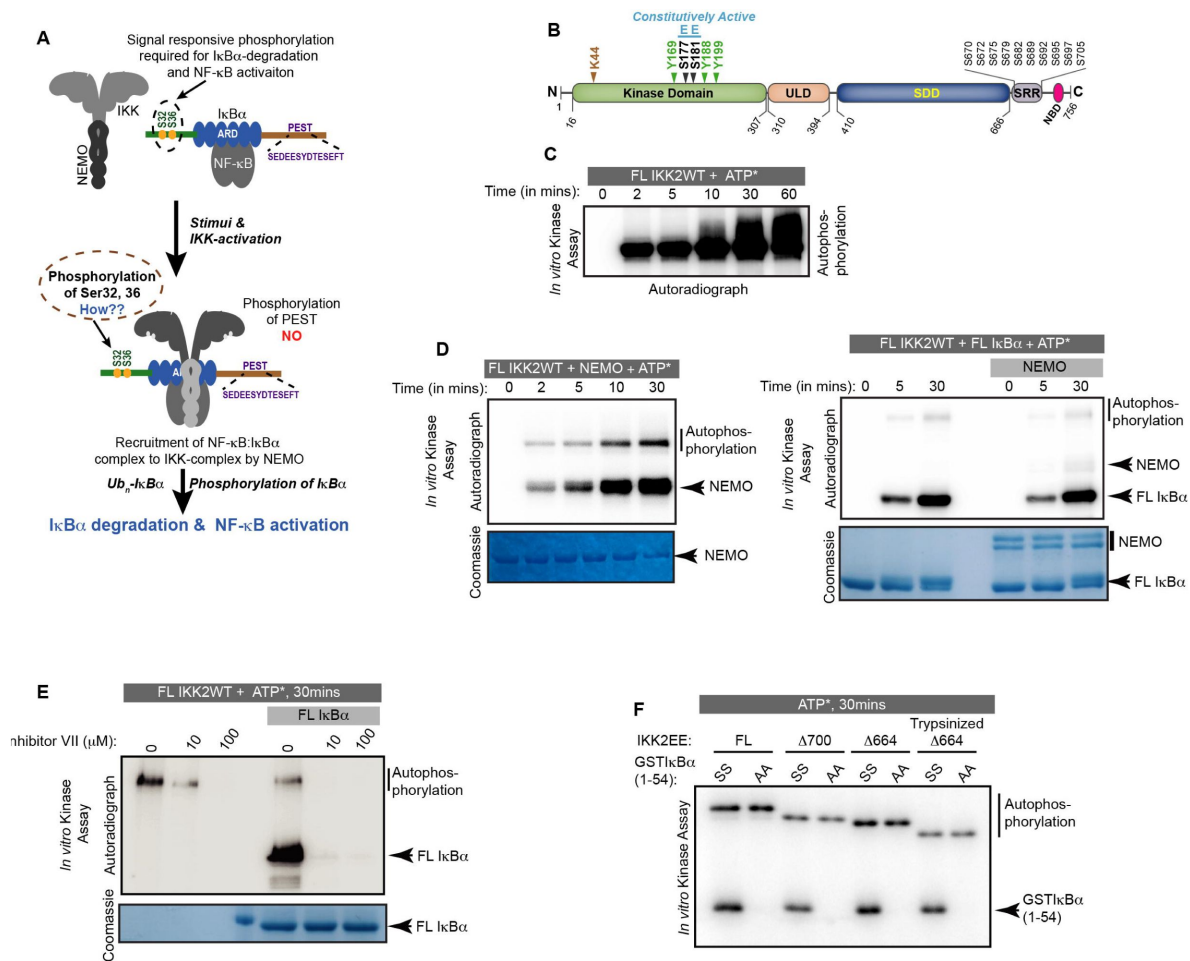


Figure 1

Autophosphorylation of IKK2 at hitherto uncharacterized sites.

(A) NEMO is known to remain bound to the IKK-complex pre- and post-stimulation, helps in activation as well as substrate recognition to channelize IKK-activity towards IκBα. However, it is not clear how IKK specifically phosphorylates the Ser32&36 of IκBα. (B) Domain organisation of FL IKK2 depicting functional kinase domain (KD), ubiquitin-like domain (ULD), and scaffold dimerization domain (SDD) based on the X-ray structures. Known sites of Ser phosphorylation and NEMO interacting regions at the extreme C-termini are highlighted. Serine and tyrosine residues in the activation loop are shown, mutation of which to Glu renders IKK2 constitutively active. The conserved ATP-interacting Lys (at position 44) is also shown. (C) Autophosphorylation of FL IKK2WT upon incubation with radioactive ATP for different time periods in an *in vitro* kinase assay. (D) *In vitro* radioactive kinase assays performed to check the effect of NEMO on autophosphorylation of FL IKK2WT (left panel), and that of NEMO and IκBα on autophosphorylation and substrate phosphorylation (right panel) activity of FL IKK2WT. (E) Effect of Inhibitor VII on FL IKK2WT autophosphorylation and IκBα substrate phosphorylation at different inhibitor concentrations in a radioactive *in vitro* kinase assay. A schematic representation of the assay procedure is shown in Figure S1B. This assay was performed twice. (F) Radioactive kinase assay conducted to check auto- and substrate phosphorylation of full length and deletion constructs of constitutively active form of IKK2 harbouring phosphomimetic Ser177Glu and Ser181Glu mutations.

63 linked poly-ubiquitin (Ub) chains or at a high concentration of IKK2, also enables *trans* auto-phosphorylation at S177 and S181 (Chen, 2012; Du et al., 2022; Ea et al., 2006; Polley et al., 2013). It was also shown that NEMO not only helps in the activation of the IKK-complex, but also ensures that the IKK-activity is well directed towards I κ B α in the inhibitor bound transcriptionally inactive NF- κ B (NF- κ B: I κ B α complex) (Schröfelbauer et al., 2012). Activated IKK2 phosphorylates I κ B α specifically at residues S32/S36, for which the C-terminal SDD of IKK2 (residue range 645-756) is critical – a shorter construct of IKK2 lacking the NBD and SDD displayed CK2-like activity that did not retain the exquisite specificity of IKK2 to phosphorylate the S32/S36 of I κ B α (Shaul et al., 2008). Other sites of I κ B α e.g., the residues in the C-terminal PEST region that contains proline (P), glutamic acid (E), serine (S) and threonine (T) residues, but not S32/S36, can be phosphorylated by various kinases (Barroga et al., 1995; Teragaonkar et al., 2003). The role of I κ B α -phosphorylation at sites other than the signal responsive S32/S36 phosphorylation in NF- κ B activation and signaling is elusive. These findings underscore the ability of IKK2 to function specifically in order to guarantee rapid NF- κ B activation in canonical signaling. It is however, still not fully understood how IKK2 achieves its exquisite specificity towards S32/S36 of I κ B α to mark it for proteasomal degradation that liberates active NF- κ B dimers.

Here we report an in-depth analysis of the catalytic features of IKK2 that revealed that IKK2 undergoes multisite autophosphorylation and can phosphorylate its own tyrosine residue(s) in addition to its own serines or the serines of its substrate I κ B α , i.e., IKK2 possesses dual specificity. This phosphorylation at tyrosine residue(s) is autocatalytic. This phosphorylation at tyrosine residues is dependent upon the prior activation of IKK2 via activation loop S177/S181 phosphorylation, and is critical to phosphorylation of signal responsive S32/S36 of I κ B α . Mutation of Y169 of IKK2 to phospho-ablative phenylalanine severely compromised phosphorylation of I κ B α at S32. Mutation of Y188 to phenylalanine rendered the kinase practically inactive *in vitro*. IKK2 activity towards signal responsive S32/S36 of I κ B α was severely compromised in MEF cells reconstituted with these two mutant IKK2s in response to treatment with TNF α . We also observed that mutation of the conserved ATP-interacting residue K44 in IKK2 to methionine leads to a loss of tyrosine as well as S177, S181 auto-phosphorylation activity of the kinase along with its ability to phosphorylate S32/S36 of I κ B α , thereby clearly suggesting a critical role of this Lysine residue in regulating IKK2's kinase activity for tyrosine as well as serine phosphorylations. Interestingly, IKK2 K44M retained non-specific kinase activity referring to phosphorylation of I κ B α at residues other than S32 and S36. Furthermore, we observed that, after auto-phosphorylation at tyrosine, IKK2 can phosphorylate I κ B α in the absence of exogenous supply of ATP, as long as ADP is present in the reaction. Overall, our results reveal unique, and likely transient, autophosphorylated intermediates of IKK2 that appear to be critical in relaying phosphoryl group(s) specifically to S32/S36 of I κ B α rather than following the conventional mode of phosphate group transfer directly from ATP to the substrate seen in eukaryotic protein kinases. IKK2 thus ensures fidelity of NF- κ B activation. Recent reports in other kinases indicate that multisite (auto)phosphorylation is a major determinant in regulating function of respective kinases and regulation exerted by those kinases. Our study adds another layer of regulation and critical nature of such autophosphorylation that plays a significant biological role in metazoans.

Results

IKK2 undergoes auto-phosphorylation at uncharacterized sites

Signal-induced phosphorylation of activation loop Ser177/Ser181 marks the general activation of IKK2 (Figure 1B) from an unphosphorylated inactive state. Phosphorylation of several other serines located within the flexible C-terminal region of IKK2, spanning amino acid 701 to the end, has also been reported (Figure 1B) (Delhase et al., 1999). Hyperphosphorylation of these C-terminal sites down-regulates IKK2 activity in cells. It was also reported earlier that, ectopically overexpressed IKK2 is capable of autophosphorylating these two serines in the activation loop in

trans even in the absence of an activating physiological signal. Indeed, recombinantly purified FL IKK2WT is capable of autophosphorylation *in vitro* when incubated with $\gamma^{32}\text{P}$ -ATP that increases with the time of incubation (**Figure 1C**). The spread-out of slower migrating phospho-IKK2 band likely indicates phosphorylation of the protein at multiple sites. We also observed that autophosphorylation of IKK2 occurred irrespective of the presence of its cognate binding partner NEMO and *bonafide* substrate I κ B α (**Figure 1D**). To confirm that this phosphorylation of IKK2 was indeed autophosphorylation and not due to an unrelated kinase copurified with IKK2, we generated a K44M mutant of full-length IKK2 where the ATP-binding invariant lysine residue conserved in Ser/Thr/Tyr kinases was mutated to methionine. This lysine residue anchors ATP in the respective binding pocket of the kinase by forming hydrogen bonds with the α - and β -phosphate of ATP that is critical for productive phosphorylation reaction in protein kinases. Mutating this K to M is expected to greatly reduce or eliminate the kinase activity. An *in vitro* kinase assay using this mutant along with the WT kinase using $\gamma^{32}\text{P}$ -ATP as the phosphor-donor confirmed that the IKK2 K44M lacked both I κ B α (substrate) and autophosphorylation activities (Figure S1A). That these phosphorylations were indeed autophosphorylation was further supported by another *in vitro* radioactive kinase assay performed in the presence or absence of an IKK-specific ATP-competitive inhibitor, Calbiochem Inhibitor VII (**Figure 1E**) where the kinase and kinase-substrate reactions were incubated for 30 minutes with the inhibitor prior to adding the phosphor-donor, $\gamma^{32}\text{P}$ -ATP. A similar assay was performed with cold ATP and specific phosphorylation of S32/36 on I κ B α was monitored by immunoblotting in presence and absence of the same inhibitor (Figure S1B). Purification status of recombinant FL IKK2 WT and FL IKK2 K44M proteins used in this study are shown in Figure S1C.

Next, we used different versions of the IKK2 protein where all previously known phosphorylation sites were lacking: Ser177 and 181 were mutated to phosphor-mimetic Glu (EE; this version of IKK2 is known to be constitutively active (Zandi et al., 1998)), and the C-terminal serine-rich region was truncated to different extents. The shortest construct,

Δ 664EE that entirely lacked the flexible C-terminus (670-756) was further subjected to limited proteolysis using trypsin to eliminate residues, if any, from the flexible ends of the enzymes and this trypsin-treated IKK2 was purified prior to *in vitro* kinase assays. Each of these constructs was active in phosphorylating itself and the substrate I κ B α (**Figure 1F**). The occurrence of autophosphorylation despite the absence of activation loop serines and serines within the C-terminus strongly suggested the presence of hitherto uncharacterized sites of autophosphorylation in IKK2.

Auto-phosphorylation of IKK2 reveals its dual specificity

IKK2 is known as a *bonafide* Ser/Thr kinase that undergoes autophosphorylation. In the previous section, we found that IKK2's autophosphorylation activity might extend beyond the previously characterized site(s). We asked if it is a dual specificity kinase. To test if Tyr residues can undergo auto-phosphorylation in addition to Ser, IKK2 incubated with Mg^{2+} -ATP for various time periods was probed with antibodies that specifically recognize activation loop serines of IKK2 (**Figure 2A**, upper panel) or phospho-tyrosine residues (**Figure 2A**, middle panel). Unexpectedly, we observed *de novo* auto-phosphorylation of IKK2 on tyrosine residues along with the anticipated auto-phosphorylation of activation loop serines. It may be noted that autophosphorylation activation loop-S177/181 also increased with time, indicating inhomogeneous and incomplete phosphorylation of activation loop-S177/181 in the recombinant IKK2 when purified from Sf9 cells. We tested another phospho-tyrosine specific antibody to further substantiate this *de novo* tyrosine autophosphorylation of IKK2. It confirmed that detection of *de novo* phosphotyrosine on IKK2 was independent of the source of the antibody used (**Figure 2B**). The tyrosine auto-phosphorylation was abolished in the presence of an IKK-specific inhibitor, Inhibitor VII (**Figure 2C**). A similar assay was performed to monitor the phospho-tyrosine level in IKK2 as a function of urea concentration. This revealed the detrimental effects of urea on tyrosine autophosphorylation of IKK2 at concentrations beyond 1M (Figure S2A). A more sensitive assay with $\gamma^{32}\text{P}$ -ATP as the

phospho-donor also indicated that urea concentrations beyond 1M inhibited both auto-phosphorylation (pan-phosphorylation) and substrate (I κ B α) phosphorylation (Figure S2B). Furthermore, the IKK2-K44M mutant (kinase inactivating) did not display any tyrosine autophosphorylation when incubated with ATP unlike WT IKK2 (Figure 2D). These results confirm the novel tyrosine phosphorylation ability of IKK2, demonstrating its dual specificity to both its own Ser and Tyr residues. We also observed that activity of IKK2 towards S32 and S36 of I κ B α and its own tyrosine residue(s) were affected similarly when the kinase assays were performed in the presence of various kinase inhibitors, both general (AMPPNP and Staurosporine) and IKK-specific (MLN-120B, TPCA, Inhibitor VII) (Figure S2C). These observations confirm that canonical activity of IKK2 towards S32 and S36 of I κ B α , and towards its own tyrosine residues are mediated by the same active site.

Next, we checked the requirement of the C-terminal NEMO binding domain (NBD) for autocatalytic dual specificity. Constructs of various truncations (lacking NBD) were capable of phosphorylating itself on tyrosine residue(s), and phosphorylating I κ B α at signal-responsive S32 and S36 (Figure 2E), indicating that the NBD domain is not required for either its ability to phosphorylate S32 and S36, as well as autophosphorylating itself on tyrosine. Next, we assessed the autocatalysis at Ser and Tyr by IKK2 using a constitutively active NBD-truncated version of IKK2 (IKK2 Δ 664EE). As anticipated, the protein was recognized by monoclonal phospho-serine antibody with or without fresh ATP-treatment; however, phospho-tyrosine antibody recognized it strongly only upon fresh ATP-treatment (Figure 2F) further confirming dispensability of the NBD in *de novo* tyrosine autophosphorylation of IKK2.

Next, we asked if phosphorylation of activation loop S177/S181 was necessary for this dual specificity of IKK2. To this end, we purified the S177A, S181A (AA; non-phosphorylatable activation loop) phospho-ablative mutant of IKK2, and checked its tyrosine autophosphorylation activity. We found that the IKK2AA mutant lacked tyrosine autophosphorylation activity (Figure 2G) in an *in vitro* kinase assay. It suggests that activation loop S177/S181 serine phosphorylation is essential in conducting dual specificity in IKK2. Indeed, the IKK2 AA mutant was incapable of phosphorylating S32/S36 of I κ B α as well (Figure S2D) in an *in vitro* kinase assay.

IKK2 autophosphorylation is critical to specificity of I κ B α phosphorylation

As mentioned earlier, it was long known that activation loop-serine phosphorylation was necessary for IKK-activation; we have now established that IKK2 phosphorylates itself on tyrosine residue(s) in addition to activation loop-serines. We wanted to further understand the significance of IKK2 autophosphorylation in the context of I κ B α S32/S36 phosphorylation. It is known that disruption of a universal salt bridge mediated by the ATP-binding Lys (K44 in IKK2) is detrimental for kinase activity. However, it is not known if disruption of this salt bridge has any bearing on substrate specificity of eukaryotic protein kinases (EPKs). During our investigation, we noted unusual behavior in the IKK2-K44M mutant. We had noticed earlier that IKK2-K44M was incapable of *de novo* autophosphorylation on tyrosine residues (Figure 2D). It also failed to phosphorylate itself on the activation loop Ser177 and 181 (Figure 3A). Interestingly, IKK2-K44M phosphorylated full-length WT I κ B α robustly albeit much less efficiency than WT IKK2 (Figure 3B) in a radioactive *in vitro* kinase assay. IKK2-K44M also showed a minor auto-phosphorylation activity compared to WT IKK2 in the absence of substrate (Figure 3B). We performed a similar kinase assay using full-length WT I κ B α where cold ATP was used as the phosphate donor, and phosphorylation status of I κ B α at S32, S36 was monitored by immunoblotting using an antibody that specifically recognizes I κ B α when phosphorylated at those two N-terminal serines. It revealed that IKK2-K44M is unable to phosphorylate S32/S36 of I κ B α (Figure 3C). Since radioactive kinase assays report pan-phosphorylation of a substrate (Figure 3B), it possibly reflects phosphorylation of the PEST region of FL I κ B α . We additionally compared IKK2-K44M with WT IKK2 using radioactive kinase assay with both FL I κ B α WT and FL I κ B α AA substrates in the

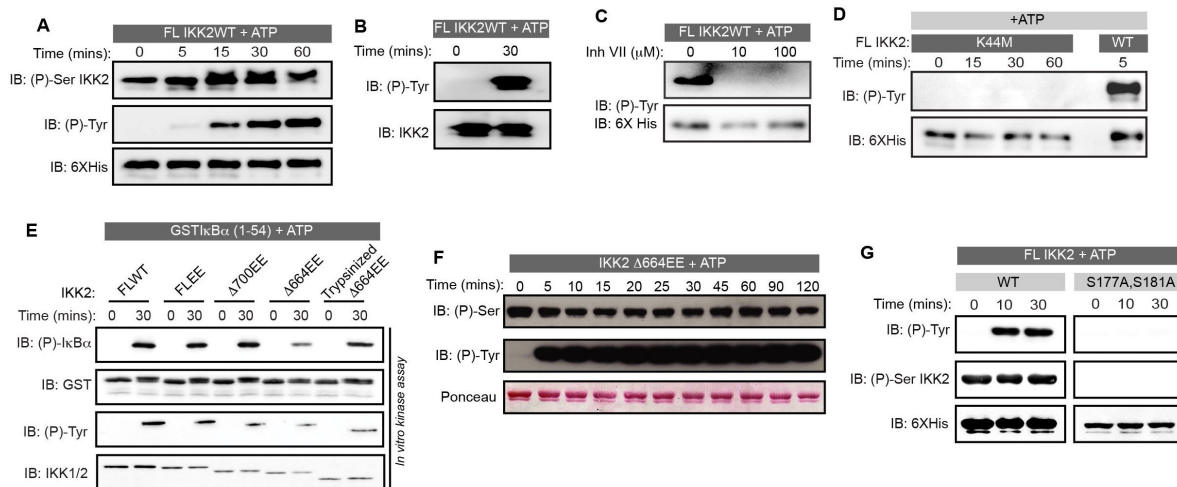


Figure 2

IKK2 possesses autocatalytic dual specificity.

(A) *In vitro* kinase assay, followed by immunoblotting with FL IKK2WT using antibodies against phosphor-Ser (177/181) and phospho-Tyr. **(B)** A different commercial source of phospho-Tyr antibody was used to detect the same. **(C)** Effect of Inhibitor VII on tyrosine autophosphorylation of FL IKK2WT. **(D)** Immunoblotting performed with phospho-Tyr antibody to check autophosphorylation of IKK2 mutant K44M at different time points compared to that of IKK2 WT. **(E)** Tyrosine autophosphorylation and substrate phosphorylation of GST (1-54) IκBα in presence of full length and deletion mutants of IKK2 harbouring phosphomimetic Ser177Glu and Ser181Glu mutations. **(F)** *De novo* auto-phosphorylation of IKK2 on tyrosine monitored by phospho-Ser and phospho-Tyr specific monoclonal antibodies. **(G)** Autophosphorylation and subsequent immunoblot analysis of FLIKK2 mutant S177A, S181A using antibodies against phospho-IKK2-Ser(177/181) and phospho-Tyr.

absence or presence of Inhibitor VII. As anticipated, FL I κ B α was phosphorylated much more efficiently than FL I κ B α AA by WT IKK2, and phosphorylation was practically abolished in the presence of 10 μ M Inhibitor VII (**Figure 3D** [↗](#)). In contrast, IKK2-K44M phosphorylated FL I κ B α AA more efficiently than FL I κ B α , and reduction in the presence of inhibitor VII was strikingly less (**Figure 3D** [↗](#)). It indicates that IKK2-K44M, despite retaining non-specific kinase activity towards the PEST region, lost phosphorylation activity towards S32, S36 of the *bona fide* substrate I κ B α . We subjected 100ug of FL IKK2 K44M protein used in the above studies to LC-MS/MS analysis, which did not detect any contaminating kinase from Sf9 cells. It is noteworthy that an equivalent K to M mutation in Erk2 also did not abolish the basal activity, but rather retained ~5 % of it (Robbins et al., 1993 [↗](#)). The above set of experiments confirm that K44 residue is not only crucial for both IKK2's well-established serine phosphorylation as well as newly-found tyrosine autophosphorylation activities, as well as its specificity for phosphorylating its *bona fide* substrate I κ B α at S32/36. Hence, these results strongly suggest that IKK2-autophosphorylation is critical for its substrate specificity.

Tyrosine phosphorylation is critically important for IKK2 activity

A number of earlier studies indicated that phosphorylation of IKK2 on Y169, Y188 and/or Y199 to be critical for its function (Darwech et al., 2010 [↗](#); Meyer et al., 2013 [↗](#)). Our mass spectrometric analyses identified multiple tyrosine residues to be phosphorylated, e.g., Y169, Y188, Y199, Y325 (data not shown). We compared available structural models of human IKK2 (PDB ID: 4KIK, 4E3C) to investigate whether tyrosine phosphorylation within the activation loop could be supported based on its structural features (Liu et al., 2013 [↗](#); Polley et al., 2013 [↗](#)). The structure of IKK2 dimer is asymmetric in that the activation loop of one monomer is largely disordered while that of the other monomer is mostly ordered (PDB ID: 4KIK) (Liu et al., 2013 [↗](#)). In the active monomers S177 and S181 are clearly observed. In these active state conformers, the position and conformation of Tyr169 appears to be well-poised for accepting a phosphate from ATP (Figure S4A) intramolecularly. A superposition of IKK2 with a pseudo-substrate-bound PKA shows that the hydroxyl of Tyr169 in IKK2 projects toward the γ -phosphate of ATP similarly to the serine hydroxyl on the PKA inhibitor pseudo-substrate (Knighton et al., 1991 [↗](#); Nolen et al., 2004 [↗](#)) (Figure S4A & B). Interestingly, Y169 is the DFG+1 (DLG in case of IKK2 and IKK1) residue while Y188 (equivalent to Y204 of PKA) is located at the end of activation loop prior to the conserved APE motif (**Figure 4A** [↗](#) and **4B** [↗](#)). It was previously established that the DFG+1 position is crucial for defining the substrate specificity of a kinase (Chen et al., 2014 [↗](#)). Y188 counterpart Y204 of PKA is reported to allosterically modulate PKA activity by altering its conformational entropy (Ahuja et al., 2017 [↗](#); Yang et al., 2005 [↗](#)). We further focused our attention to Y169 and Y188 as they are located in the activation loop of the kinase.

To test the effect of phosphorylation at Y169 and Y188, we generated Y169F and Y188F mutants of IKK2. Radioactive *in vitro* kinase assays using recombinant purified IKK2^{FL}-Y188F and IKK2^{FL}-Y169F mutants revealed a drastic reduction in the activity of IKK2 (**Figure 4C** [↗](#)). Since the DFG+1 residue in kinases (Y169 in IKK2) is known to dictate its specificity, we created different mutants of the I κ B α substrate where S32 and S36 were singly and independently altered to either alanine (phosphoablative) or glutamic acid (phosphomimetic) residues. Kinase activity of IKK2-Y169F was monitored in an *in vitro* radioactive kinase assay using WT as well as mutant I κ B α . Mutation of S36 to either alanine or glutamate drastically reduced the phosphorylation signal of I κ B α while that for the S32A/E mutants were minimally affected (**Figure 4D** [↗](#)). This suggests that the presence of Tyr (and possibly its phosphorylation) is critically important for signal-responsive phosphorylation of I κ B α .

Next, we tested the consequence of Y169F and Y188F mutants on the pro-inflammatory signal-induced IKK activity in cells. We reconstituted IKK2 knock-out mouse embryonic fibroblast (*ikk2*^{-/-} MEF) cells with WT, Y169F and Y188F mutants of IKK2, and measured signal-induced activation of IKK2 upon TNF- α treatment. IKK2 activity is significantly reduced in Y169F and nearly abolished

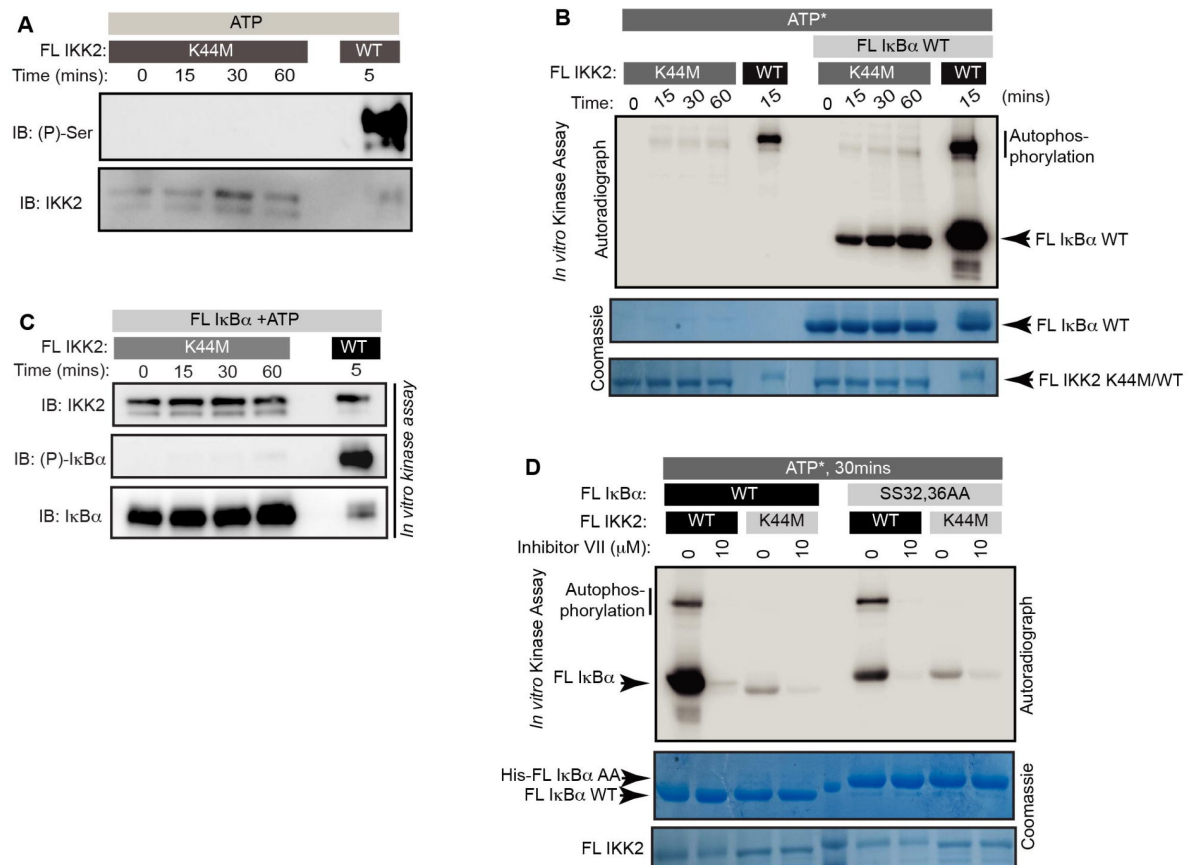


Figure 3

Autophosphorylation of IKK2 is critical to the specificity of IκBα phosphorylation.

(A) Immunoblotting performed with phospho-IKK2 Ser(177/181) antibody to check autophosphorylation of IKK2 mutant K44M at different time points compared to that of IKK2 WT. (B) Comparison of substrate and autophosphorylation activities of FL IKK2 WT and K44M mutant in an *in vitro* radioactive kinase assay performed with or without FL IκBα WT as the substrate. (C) Residue level specificity of phosphorylation by the FL IKK2 K44M was analysed using an antibody that specifically recognizes phospho-S32/36 of IκBα. (D) Differential sensitivity of specific vs. non-specific phosphorylation by IKK2 as assessed by the effect of Inhibitor VII on FL IKK2 WT and IKK2 K44M using IκBα WT and AA as substrates.

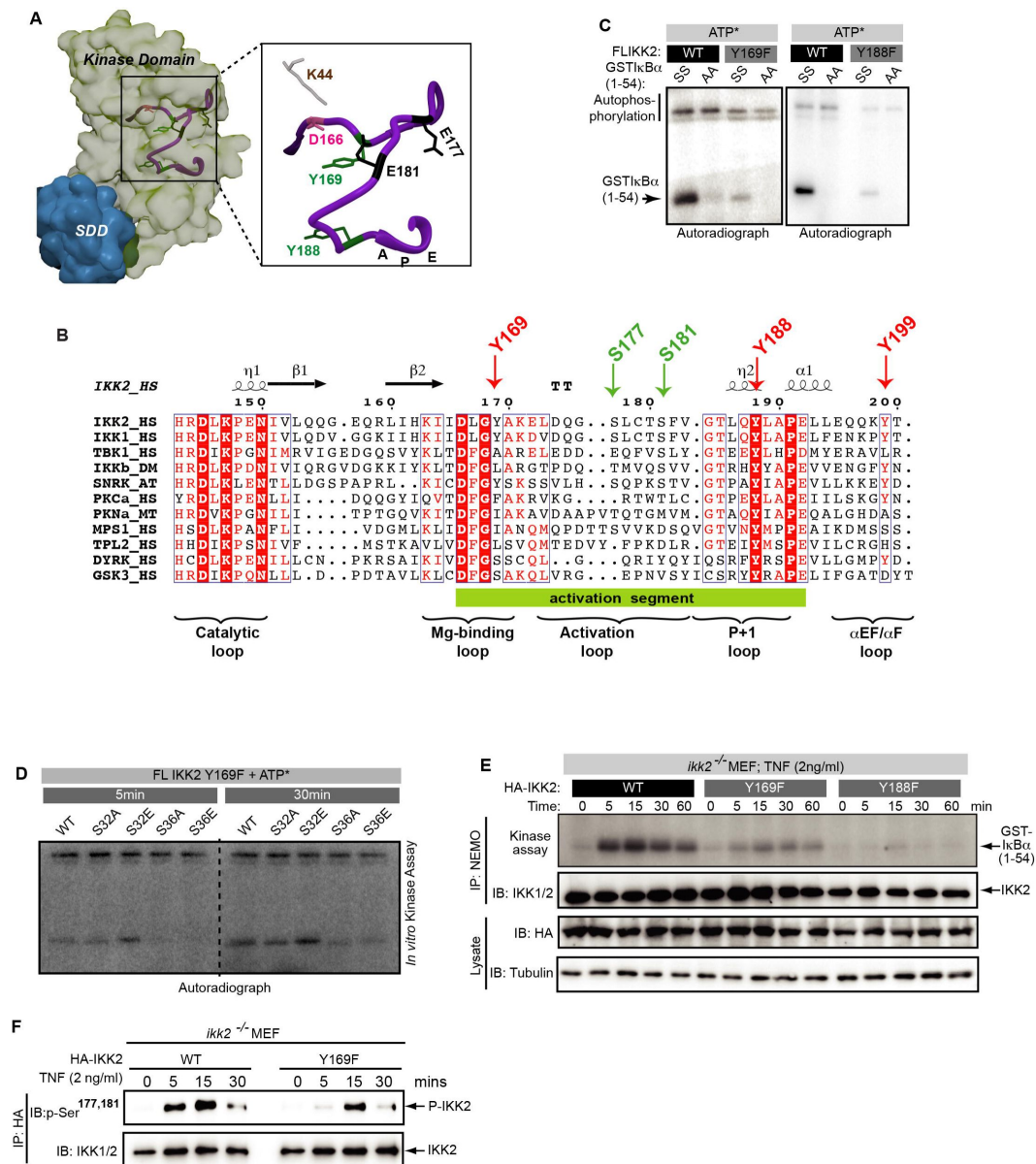


Figure 4

Tyrosine autophosphorylation is critical for IKK2 activity.

(A) Position of canonically important residues in the AL (purple ribbon) of IKK2-KD are shown in the context of its structure (PDB ID 4E3C; surface representation; KD is shown light green with respect to the SDD in teal). Autophosphorylated tyrosine residues identified in this study are shown in green. (B) Alignment of the activation segment sequences of different IKK family kinases. Conservation of Tyr at the DFG+1 (DLG in case of IKK1 and IKK2) is retained only in a stress response related plant kinase SnRK2, but not in other IKK homologues or any known dual specificity kinases, e.g., DYRK and GSK3 β (both contain Ser at that position). Tyr188 (204 in PKA) is conserved. (C) Radioactive *in vitro* kinase assay performed with IKK2 Y169F and Y188F mutants using GST (1-54) IkbA WT and AA as substrates. (D) Radioactive *in vitro* kinase assay performed with IKK2 Y169F mutant with a range of GST (1-54) IkbA mutants where the substrate phosphorylation signal is abolished in case of S36 A/E IkbA. (E) Severe reduction in TNF α -induced IKK activity in *ikk2*^{-/-} MEF-3T3 cells reconstituted with mutant (Y169F and Y188F) IKK2s compared to wild-type. Kinase assay was performed with IKK immunoprecipitated (IP-ed) with anti-NEMO antibody from whole cell extract. (F) AL-serine phosphorylation status of IKK2 WT and Y169F was assessed in reconstituted MEFs post-stimulation with TNF- α after immunoprecipitating the kinase with monoclonal anti-HA antibody.

in Y188F (**Figure 4E** [↗](#)). Interestingly, phosphorylation of activation loop serines of the less active Y169F mutant was only marginally defective (**Fig. 4F** [↗](#)). This suggests that phosphorylation of the activation loop serines at 177 and 181 is necessary but not sufficient for IKK2 to become fully active. Together, these results suggest an unreported communication between the phosphorylation of Y169 of IKK2 and that of S32 of IκBα.

Structural analyses of IKK2 autophosphorylation

To get a more comprehensive picture, it is necessary to analyze at least three phosphorylated states of IKK2: unphosphorylated (UnP-IKK2), phosphorylated at S177 and S181 (p-IKK2), and phosphorylated at Y169, S177 and S181 (P-IKK2). We took advantage of computational approaches using molecular dynamic (MD) simulations and flexible molecular docking (see methods).^{43–50} MD simulation was performed for 200ns for each of these three states of IKK2, unphosphorylated and phosphorylated. Phosphorylation at S177, S181 increased the stability of the kinase, which was further increased by the phosphorylation at Y169 as evidenced by a gradual decrease in the total energy of system (**Figure 5A** [↗](#)). A striking enhancement in stability of IKK2 upon this auto-phosphorylation is reflected in the change of T_m from ~ 40 °C to ~ 50 °C in differential scanning calorimetry (DSC) scans in the presence of ADP and ATP (**Figure S4A**). This behavior supports our observation in MD simulation, and tallies with increased solubility of IKK2 observed after auto-phosphorylation that was essential for its successful crystallization (Polley et al., 2013 [↗](#))

Interestingly, RMSD values increased with more phosphorylation (**Figure 5B** [↗](#)) despite phosphorylation-induced stabilization, P-IKK2 showed the highest RMSD followed by p-IKK2 and UnP-IKK2, indicating increased internal motion in the kinase as a result of phosphorylation. It may be noted that, the changes in total energy and RMSD are consistent throughout the course of simulation though the magnitude is small. We found that, phosphorylation at S177, S181 and S177, S181, Y169 induced distinct structural alteration in the KD when compared with the unphosphorylated version (**Figure 5C** [↗](#), models shown separately in **Figure S4B**). Careful analyses of the trajectory of RMSF values for these three states revealed that, different regions of the kinase domain had different RMSF values in different phosphorylated states (**Figure S4C**). We observed that, while the glycine-rich loop and the αC-helix in p-IKK2 were situated in a manner reminiscent of an active kinase (**Figure S4D**), those in UnP and P-IKK2 were placed very differently. In fact, helix αC was found to be distorted in P-IKK2, that is unlikely to support canonical phosphotransfer from ATP. Activation segment/loop (activation loop) in these three differently phosphorylated states adopted conformations distinct from each other. activation loop in the UnP model was displayed out and moved closer to the tip of the C-lobe, whereas activation loop in p-IKK2 and P-IKK2 moved inwards and adopted conformations closer to the N-lobe. The formation of the proper R- and C-spines in p-IKK2 confirmed its active form (**Figure 5D** [↗](#)). p-IKK2 KD also exhibited other canonical features of an active kinase, viz., a salt bridge between K44 and E61 (K72, E91 in PKA, respectively), ‘DFG in’ (DLG here) conformation. Next, we docked ATP onto the p-IKK2 structure (ΔG = -10.64 kcal/mol) using a flexible docking protocol (see methods). The ATP and its phosphate groups exhibited a pose (**Figure S4D**) very similar to that in the PKA structure (PDB ID: 1ATP), and the terminal phosphate was in close proximity to Y169-OH making autophosphorylation of Y169 highly plausible (**Figure 5E** [↗](#)). Further analyses of the ATP-bound p-IKK2 structure confirm that presence of ATP helped the R and C-spine form the desired continuum seen in the active form of a kinase (**Figure 5F** [↗](#)). Superposition of the ATP from the p-IKK2 on P-IKK2 indicated that the narrowing of ATP-binding pocket in P-IKK2 may lead to rejection of ATP emanating from severe clashes between the glycine-rich loop and ATP (**Figure 5G** [↗](#)). Furthermore, the R-spine in P-IKK2 was not continuous (**Figure 5H** [↗](#)), wherein M65 (L95 in PKA) of the four conserved residues were located at a position far away from the rest of three residues that constitute the R-spine, and from the glycine rich loop. In addition, dynamic cross-correlation matrix (DCCM) or contact map of each structure suggests that specific phosphorylation events render the kinase display distinct allosteric changes at locations far from the phosphorylation sites themselves (**Figure S4E**).

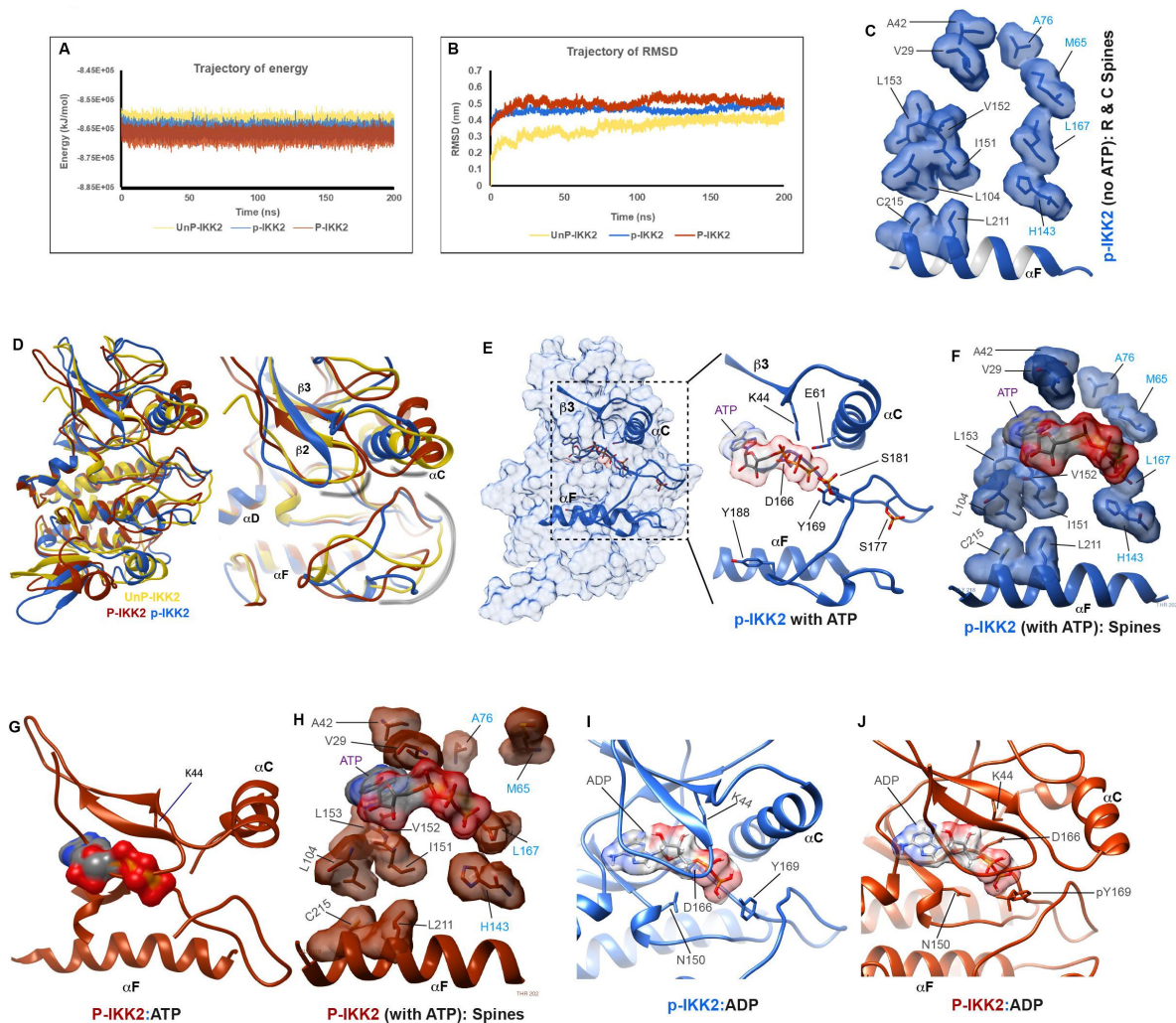


Figure 5

Structural analyses of IKK2 autophosphorylation.

(A) Molecular dynamics (MD) simulations of all three differently phosphorylated states of IKK2 was performed. 200ns trajectory of energy for each state is shown in golden yellow, blue and copper red representing UnP-IKK2, p-IKK2 and P-IKK2, respectively. Same color coding has been followed throughout this figure and the corresponding supplementary figure. (B) 200ns trajectory of RMSD for each state is shown. (C) Residues forming canonical R- (labelled in blue) & C-spines (labelled in dark gray) representative of an active conformation in p-IKK2 are shown. (D) Superposition of structures representing the differently phosphorylated states post 200ns MD simulation. Relevant sections, e.g., activation loop, αC -helix, Gly-rich loop areas are highlighted in the right panel. (E) ATP-bound p-IKK2 structure is depicted. In the right panel, relevant residues are highlighted. Y169 is found in a position poised for autophosphorylation. (F) ATP completes the desired continuum of R- & C-spine observed in active kinase conformations. (G) P-IKK2 cannot accommodate ATP in its cleft, and αC -helix is displaced. (H) M is moved away from the R-spine, failing to form canonically active conformation of R- & C-spines. (I & J) ADP-bound p-IKK2 and P-IKK2 states are shown, respectively. Both p-IKK2 and P-IKK2 states can accommodate ADP in their respective clefts.

ATP binding to p-IKK2 leads to autophosphorylation at Y169 generating P-IKK2 in an ADP bound state. Interestingly, the superposition of ADP (from the docked complex of p-IKK2 and ADP, $\Delta G = -9.88$ kCal/mol, **Figure 5I**) onto the P-IKK2 structure confirmed that while P-IKK2 was unable to accommodate ATP in a manner similar to that in other known experimentally derived kinase structures, it could comfortably accommodate ADP into the ATP-binding cleft (**Figure 5J**). It may be noted that, p-IKK2 has higher affinity for ATP than ADP as indicated by their respective binding energy values. We speculate that ADP-bound P-IKK2 is an intermediate step where presence of as many phosphate groups in the activation loop and ADP in such a close proximity persuades the kinase to adopt a conformation that can withstand this high density of negative charges albeit risking the integrity of an active fold, and making some of those *de novo* phosphate groups transient in nature to return back to the p-IKK2 state to readily bind to ATP that is abundant in the reaction.

These structural observations suggest that it is possible for the residues within the flexible activation segment to undergo auto-phosphorylation and possibly render the kinase with a novel phospho-transfer activity distinct from the conventional transfer of γ -phosphate from ATP to the substrate.

Freshly auto-phosphorylated IKK2 phosphorylates I κ B α even in absence of ATP

We observed phosphorylation of tyrosine residue(s) only upon fresh ATP-treatment, indicating that tyrosines of purified IKK2 loses its phosphate group during the course of purification (Eckhart et al., 1979). It may be noted that, we did not observe any density of phosphates on any of these residues in the human IKK2 crystal structure derived from ATP-treated protein sample (Polley et al., 2013). Of note, human IKK2 crystals grown with autophosphorylated IKK2 (PDB ID 4E3C) took about a month at 18-20°C before they were subjected to X-ray diffraction. This suggested an intriguing possibility that the phosphate linked to Tyr residue(s) may serve as a temporary phosphate-sink and could eventually be transferred to substrate I κ B α . To test this possibility, we first allowed purified IKK2 to undergo auto-phosphorylation by incubating the excess kinase with limiting amounts of radiolabeled γ - 32 P-ATP followed by two passes through desalting spin columns (40 kDa MW cut-off) to remove unreacted ATP. Nucleotide-free auto-phosphorylated IKK2 (P-IKK2) was incubated with I κ B α either in absence of nucleotide or with newly added unlabeled nucleotides, and phosphorylation status of the substrate was subsequently monitored by autoradiography after resolving the reaction mixture on an SDS-PAGE (**Figure 6A**). Transfer of radiolabeled phosphate to I κ B α was observed and the transfer was significantly enhanced in the presence of cold ADP (**Figure 6B**). Phosphorylation of I κ B α did not significantly decrease in the presence of a high concentration of cold ATP. That addition of excess cold ATP did not reduce the extent of I κ B α phosphorylation-signal on the autoradiograph strongly suggests the transfer of 32 P-phosphate from 32 P-IKK2 to I κ B α (**Figure 6B**) by a relay mechanism. This transfer perhaps occurs through a phosphoenzyme intermediate with or without involving ADP.

We undertook a different strategy to further confirm transfer of phosphate from the kinase (and not ATP) to the substrate. We auto-phosphorylated IKK2^{FL}-WT with excess cold ATP and removed the unreacted ATP from the reaction mixture using size-exclusion chromatography on a Superdex200 column (**Figure 6C**). P-IKK2 eluted at fractions corresponding to MW between 670 and 158 kDa, i.e., much larger than free ATP (**Figure 6D**). This P-IKK2 protein was incubated with FL I κ B α either in presence or absence of ADP for different time periods. A similar reaction in presence of ATP instead of ADP was performed as a positive control. Phosphorylation of S32 and S36 (detected by the phospho-serine antibody specific to S32 and S36 of I κ B α) was efficient in presence of 10 and 50 μ M ADP but absent when ADP was absent in the reaction mixture. Phosphorylation of I κ B α in the presence of 50 μ M ATP was significantly more robust as anticipated (**Figure 6E**). The minor detection of I κ B α phosphorylation in absence of ADP in the radioactive assay and lack of it in the cold immunoblotting assay (compare lanes marked with vertical arrows

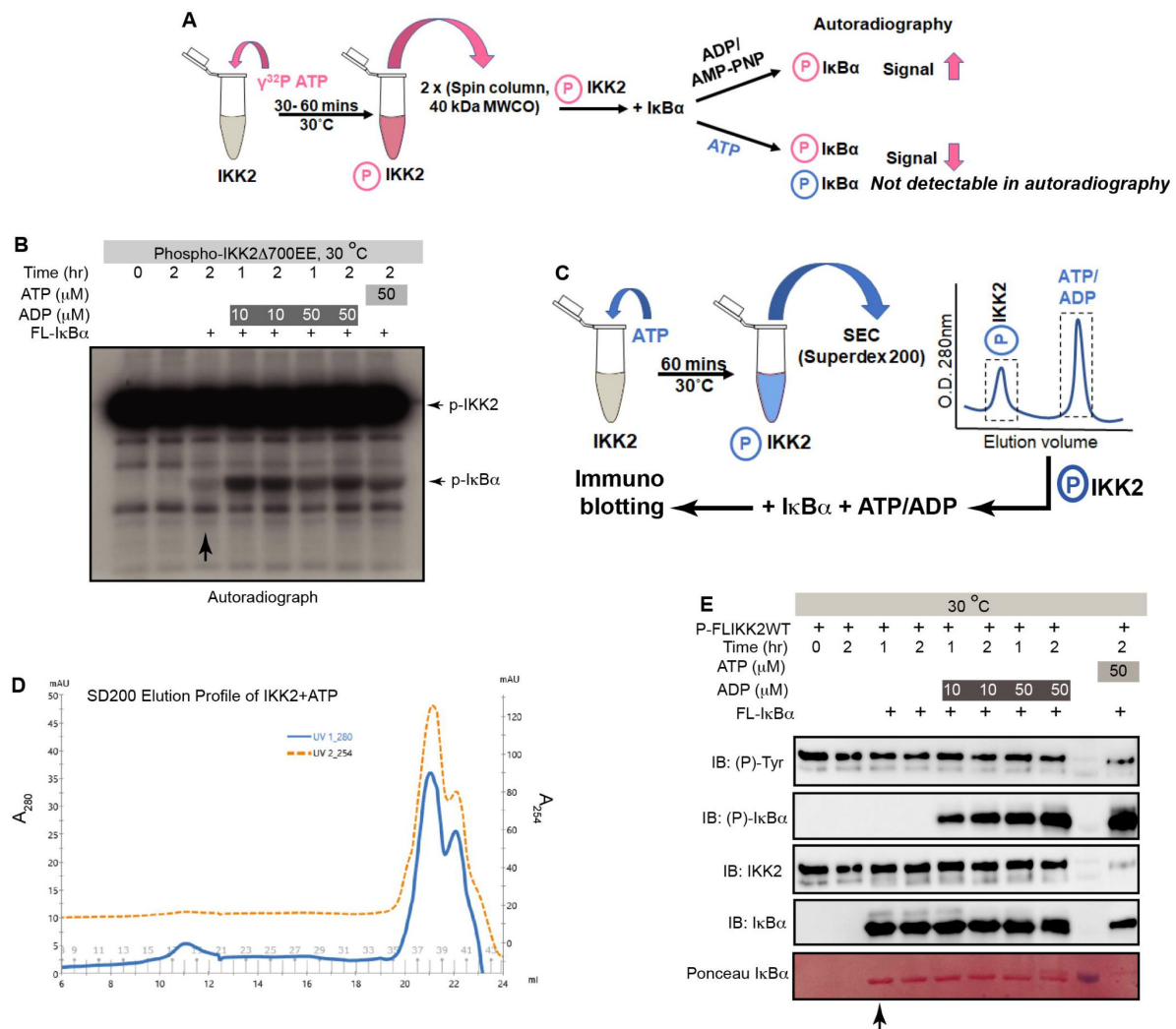


Figure 6

Freshly autophosphorylated IKK2 relays phosphates to IkBα.

(A) Schematic representation of the experiment performed to monitor the path traversed by phosphate(s) from phospho-IKK2 to substrate by autoradiography detection method. (B) Purified autophosphorylated (radiolabelled) IKK2 transfers its phosphate to IkBα substrate in absence of any nucleotide, and transfer efficiency is enhanced upon addition of ADP or ATP. (C) Schematic representation of the experiment performed to monitor the path traversed by phosphate(s) from phospho-IKK2 to substrate by immunoblotting detection method. (D) Elution profile of the size exclusion chromatography run of phospho-IKK2 to get rid of excess unlabelled ATP performed in Superdex200 10/30 increase. (E) Purified autophosphorylated (cold) IKK2 transfers its phosphate to IkBα substrate with enhanced transfer efficiency upon addition of ADP or ATP monitored by immunoblotting using specific antibodies as indicated in the figure.

in **Figures 6B** and **6F**) may indicate higher sensitivity of the radioactive assay. The need of ADP in phosphotransfer prompted us to consider whether the ADP was contaminated with a trace amount of ATP. ESI-MS analysis of 50 μ M ADP did not detect any peak corresponding to ATP (Figure S5A). Dependence of I κ B α phosphorylation on ADP could raise another possibility if microscopic reversibility generated ATP from ADP in the active site of the kinase. To this end, we employed TLC-based analysis to detect generation/presence of ATP in a similar setting as described in **Fig. 5A-B** which revealed that no detectable ATP was present in the system, i.e., no detectable ATP was retained in the protein after passing it through the spin column twice, nor did any γ P³²-ATP was generated when incubated with cold ADP (Figure S5B).

Phosphotransfer from P-IKK2 is critical to fidelity and specificity of I κ B α phosphorylation

We next tested if phosphotransfer is restricted only to the N-terminal signal responsive S32/S36 or to serines/threonines of the PEST domain as well. Radioactive kinase assay as described earlier was performed with full-length WT and mutant S32A, S36A (AA) I κ B α as the phosphate recipient. Phosphate group(s) of P-IKK2 was not transferred to full-length I κ B α AA which retains the serines and threonines within its PEST domain (**Figure 7A**), indicating phosphotransfer only to the N-terminal serines of I κ B α that are S32 and S36. We also tested the effect of AMPPNP, an unhydrolyzable analogue of ATP, on the phosphotransfer, and found no inhibitory effect irrespective of the presence of ADP. This suggests adenine nucleotides poisoning the kinase for phosphotransfer (**Figure 7A**). A recent report proposed that IKK2 phosphorylates I κ B α at S32 followed by S36 following a single binding event and that phosphorylation of S32 increases the phosphorylation rate of S36 (Stephenson et al., 2022). Based on our results and other evidence, we propose that IKK2 phosphorylated at S177/S181 undergoes autophosphorylation on tyrosine residues in its activation loop area that serves as a phosphoenzyme intermediate to transfer phosphate group from IKK2 to S32 that enhances the rate of second site (S36) phosphorylation (**Figure 7B**). For the second site phosphorylation ATP serves as the phosphate donor.

Discussion

Phosphorylation of substrates by protein kinases is regulated in various ways (Cullati et al., 2022; Sang et al., 2022). IKK2, a kinase central to inflammation, phosphorylates serine residues of its primary substrate I κ B α with γ -phosphate of ATP. Here, we report multiple intriguing properties of IKK2 that provide specificity to its substrate phosphorylation. These include: 1) dual specificity that entails autophosphorylation of tyrosine residues of IKK2, in addition to known autophosphorylation at activation loop serines; 2) relay of phosphate to substrate I κ B α from autophosphorylated IKK2 (P-IKK2) and 3) loss of specificity of the kinase due to disruption of a universally conserved salt bridge (mediated by K44).

Ser/Thr/Tyr family of eukaryotic protein kinases transfer γ -phosphate of ATP directly to substrate, in general to S, T and Y residues (Cohen, 2002; Fischer and Krebs, 1955; Hunter, 1991; Krebs and Fischer, 1955; Nolen et al., 2004). Histidine kinases (in prokaryotes and in some lower eukaryotes) however shuttles the γ -phosphate to Asp residue on a response regulator (RR) substrate through a phospho-His intermediate (Borkovich and Simon, 1990; Gao and Stock, 2009; Hunter, 2022; Kalagiri and Hunter, 2021; Laub et al., 2007; Robbins et al., 1993). Our present analyses of eukaryotic protein kinase IKK2 revealed its surprising property in which IKK2 phosphorylated at activation loop-serines (S177/S181) could be further auto-phosphorylated by ATP at multiple other residues yielding another form (P-IKK2) (**Figure 7B**). This marks IKK2 as an autocatalytic dual-specificity kinase rather than a prototype S/T kinase. This multisite phosphorylated form of IKK2 (P-IKK2) could transfer phosphate(s) to substrate I κ B α in presence of ADP without requiring exogenous supply of fresh ATP, hinting at an unusual property of IKK2 acting possibly as a phosphate-sink. It is possible that phospho-tyrosines residues of IKK2 act as

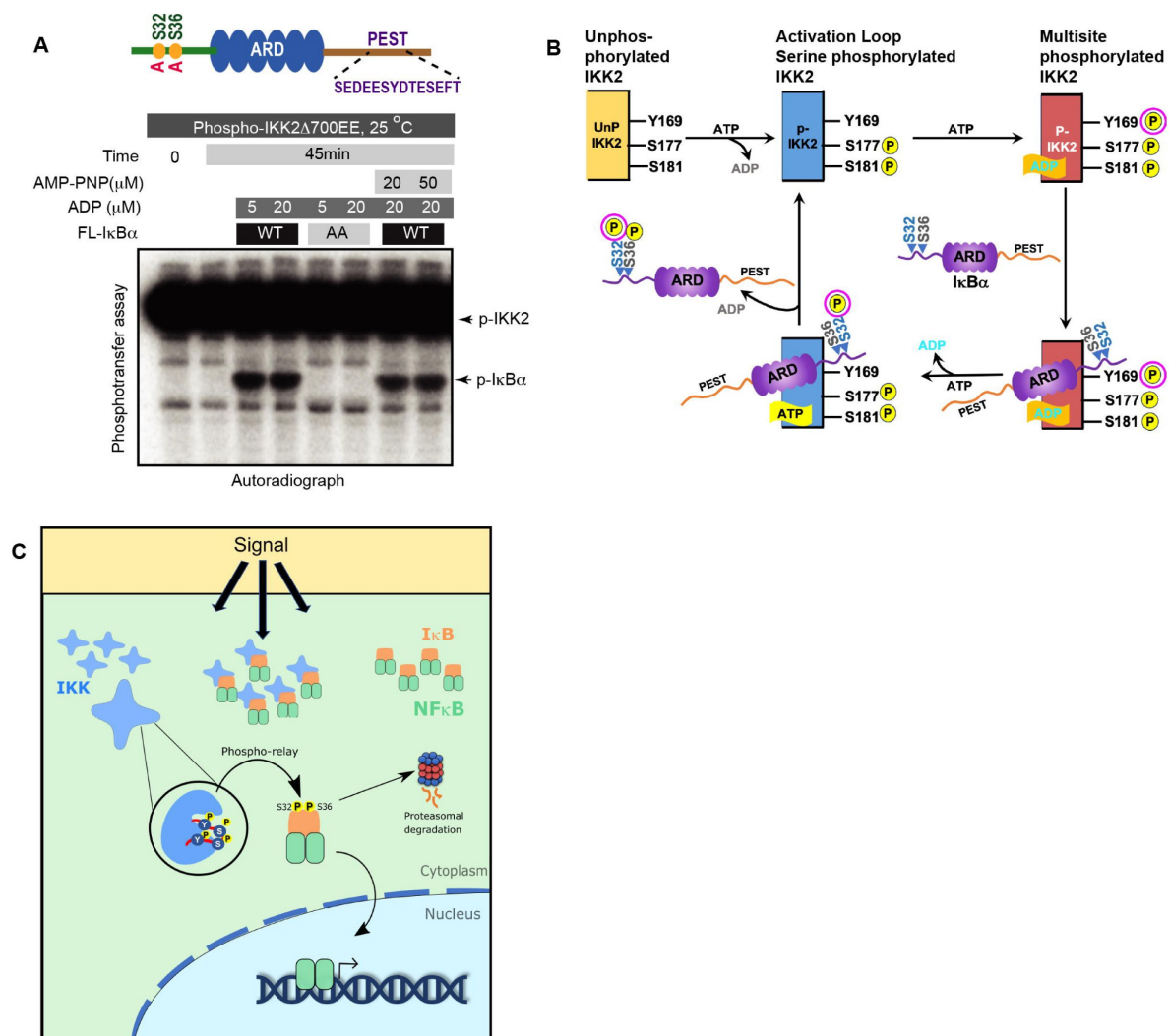


Figure 7

Specificity and fidelity of phosphotransfer by IKK2 to IκBα.

(A) AMP-PNP can support efficient phosphotransfer. Also, phosphotransfer is observed in IκBα WT but not in S32,36Ala double mutant. Domain organization and position of relevant S/T/Y residues of IκBα are shown above the autoradiograph. (B) A detailed scheme of reactions showing IKK2-activation and subsequent phosphorylation of the signal responsive S32,36 on IκBα by phosphotransfer in presence of ADP. (C) A cartoon representation of the proposed model of NF-κB activation by IKK2 employing phosphotransfer to IκBα.

this sink. The Y169 located at the DFG+1 (DLG in IKK2) position is unique only to IKK1 and IKK2, and is absent even in close mammalian homologues such as IKK ϵ and TBK1. Interestingly, neither IKK ϵ nor TBK1 can phosphorylate I κ B α at S32 and S36, *in vitro* or *in vivo* (Larabi et al., 2013; Tu et al., 2013). A recent report also indicated the role of DFG+1 residue in differentiating Ser versus Thr residues (Chen et al., 2014). These observations hinted to an involvement of Y169 in substrate selectivity. Indeed, our experimental data suggests a pivotal role of Y169 in phosphorylating S32 of I κ B α both *in vitro* and *ex vivo* (Figure 4D & 4E). It is worthwhile to note that, a phosphoablative mutation at S32 in I κ B α to Ile (S32I) is associated with ectodermal dysplasia and T-cell immunodeficiency in patients (Courtois et al., 2003; Mooster et al., 2015). More interestingly, the placement of Y169 in the kinase active site makes it amenable to a phosphorylation-dephosphorylation cycle, suggesting its possible role both in phosphotransfer and substrate specificity. Mutation of Y188, a conserved tyrosine, to non-phosphorylatable phenylalanine abolished the kinase activity of IKK2 *in vitro* as well as in MEFs upon TNF-treatment (Figure 4C & 4E). The placement of Y188 indicates that its mutation will be detrimental to the structural fold of kinase, thus it is unclear if or how Y188 plays a role in the phosphotransfer. Since phosphotransfer to S32, S36 of I κ B α is observed with auto-phosphorylated IKK2 EE, transferred phosphates are not from S177 or S181. Kinase domain autophosphorylation is known to regulate function of other kinases as well, e.g., autophosphorylation at T220 greatly influenced both the activity and substrate specificity of CK1 (Cullati et al., 2022).

The strength of input signal, level of I κ B α degradation, and NF- κ B activation correlate with the amplitude, duration, and kinetics of IKK activity (Behar and Hoffmann, 2013; Cheong et al., 2006). This digital (all-or-none) activation profile of NF- κ B appears to be due to rapid activation and inactivation of IKK2. IKK targets multiple substrates *in vivo*; however, the rapid activation of NF- κ B is regulated primarily by I κ B α . How IKK specifically selects I κ B α for phosphorylation is unclear, although NEMO is reported to direct the kinase activity of IKK2 specifically towards I κ B α (Schröfelbauer et al., 2012). We observe that P-IKK2 could transfer phosphate(s) specifically to the S32/S36 of I κ B α , and not to other sites (such as the PEST sequence of I κ B α) (Figure 6B, 6E and 7A). We surmise if transiency of multisite phosphorylated IKK2 (P-IKK2) underlie this activation-inactivation events during signal response. It is possible that regulatory system of IKK2 is somewhat analogous to the two-component histidine kinase-effector systems in prokaryotes that show a threshold activation response (Bhate et al., 2015; Lamarche et al., 2008; Laub and Goulian, 2007). It is noteworthy that, transfer of a phosphate moiety directly from pHis is energetically much more probable than that from a pTyr, unless it is an enzymatic process which is likely the case for IKK2:I κ B α system.

Tyrosines in activation loop of IKK2 are previously reported to undergo signal-induced phosphorylation in cells. These modifications were proposed to be mediated by a tyrosine kinase although there is no evidence of a specific tyrosine kinase (Meyer et al., 2013; Otero et al., 2008; Rieke et al., 2011). Several members of the Ser/Thr kinase family undergo tyrosine phosphorylation and these kinases employ diverse mechanisms for auto-phosphorylation (Bhattacharyya et al., 2006; Ge et al., 2002; Lochhead, 2009; Lochhead et al., 2006, 2005; Tigno-Aranjuez et al., 2010).

The phosphotransfer from P-IKK2 to I κ B α monitored here is detection of a single turnover event in absence of the phosphate donor, ATP in the bulk (Figure 6B). Phosphorylation on IKK2 occurs at multiple sites, and not every phosphorylated residue in P-IKK2 expected to transfer its phosphate to I κ B α . There is more than one type of phosphorylated residues in P-IKK2 distributed over the length of the kinase, arguably with differential stability wherein only a fraction of the total pool can be relayed. In addition, it is likely that the phosphorylated IKK2 in the bulk harbors a non-homogenous pool where differently phosphorylated species of the kinase are present, and are indistinguishable in our assay schemes. Another intriguing feature of phosphorelay in IKK2 is IKK2's dependence on ADP (Figure 6B & 6E). It was also observed that, the generic kinase inhibitor AMPPNP (unhydrolyzable ATP analogue) failed to inhibit phosphor-transfer from IKK2

(**Figure 7A**) and I κ B α phosphorylation (Figure S2C). Taken together, we surmise that presence of ADP or AMP-PNP help IKK2 attain a conformational state poised for phosphorylating S32/36 of I κ B α even in absence of fresh exogenous ATP. Similar phenomenon was reported for IRE1, where ADP was found to be a more potent stimulator of its ribonuclease activity and that AMP-PNP could also potentiate this activity in IRE1 (Lee et al., 2008; Sidrauski and Walter, 1997). The list of Ser/Thr kinase family proteins that undergo tyrosine phosphorylation is growing that employ diverse mechanisms for auto-phosphorylation (Bhattacharyya et al., 2006; Ge et al., 2002; Lochhead, 2009; Lochhead et al., 2006, 2005; Tigno-Aranjuez et al., 2010). The most notable example of this dual specificity are DYRK and GSK3 β . However, the role of Tyr-phosphorylation in DYRK/GSK3 β is not related to that in IKK2, nor does it render DYRKs/ GSK3 β capable of relaying phosphates to the substrates. To the best of our knowledge the only other reported example of an EPK, though atypical, transferring phosphate through a phosphoenzyme intermediate is MHCK of *Dictyostelium discoideum* (Ye et al., 2010).

Our study reveals tyrosine auto-phosphorylation in IKK2 and indicates that consequences of Ser and Tyr autophosphorylations in dual-specific IKK2 are distinct. Moreover, the crucial tyrosine residues are positioned in such a manner that some that are amenable to auto-phosphorylation are essential for substrate phosphorylation. Most surprisingly, the tyrosine-linked phosphate groups can be transferred by a relay mechanism to specific serines in I κ B α , an event reminiscing phospho-relay in two-component systems prevalent in prokaryotes and some lower eukaryotes. Our study also reveals the role of K44 of IKK2 in auto-phosphorylation and thus in specific phosphorylation of S32, S36 of I κ B α , although the mutant kinase retains its non-specific kinase activity. It may be noted that, while a monomeric version of IKK2 (devoid of NBD and large portion of the dimeric interface in the SDD) still retained its specificity for the N-terminal serines of I κ B α (Hauenstein et al., 2014), while a shorter construct containing only the KD and ULD lacked this specificity that phosphorylated the PEST domain like CK2 (Shaul et al., 2008). It requires further investigation to determine if a structural state of IKK2, perhaps through a different modification or through binding to another protein, allow it to phosphorylate substrates relatively non-specifically.

In all, it is intriguing how the gateway to the activation of NF- κ B is regulated through unique phosphorylation events in IKK2 (**Figure 7C**). It appears that a multi-layered fail-safe mechanism is devised to tightly control NF- κ B activation. Firstly, in absence of upstream signals IKK-complex is unable to phosphorylate Ser 32,36 of I κ B α efficiently thereby keeping check on the aberrant and untimely activation of NF- κ B. Upon encountering proinflammatory cues, cells need to activate NF- κ B immediately and efficiently, i.e., IKK needs to be activated and phosphorylate Ser 32,36 of I κ B α . NEMO warrants IKK-activation and ensures that the IKK-complex specifically chooses I κ B α from a large pool of substrates of IKK2. IKK2 is designed in such a manner that it phosphorylates itself at activation loop-tyrosines when activated, such that phosphate group(s) can be relayed directly to Ser32,36 of I κ B α with great fidelity, thus leaving little chance of a misconstrued signaling event thereby confirming NF- κ B activation (**Figure 7C**). Our discovery might also present the beginning of a new aspect in eukaryotic cell-signaling by EPKs and provide a foundation for mechanistic studies of an intriguing phospho-transfer reaction that helps accomplish the desired specificity. To the best of our knowledge, it is the first report, apart from MHCK of *Dictyostelium discoideum* (an atypical EPK), to suggest that a metazoan EPK confirms its specificity through the formation of a phosphoenzyme intermediate.

Methods summary

Protein purification, auto-phosphorylation assay, immunoprecipitation coupled kinase assay, phosphotransfer assay and other experiments have been described in details in the supplementary document.

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Author contribution

SP, GG and TB designed experiments; PB, SP, TB & GG performed experiments. Computational analyses were performed by AC & SC. SP, GG, TB, PB, TH, SC and AC analyzed the results. SP wrote the manuscript with the help of PB, TB and AC in preparing figures and images. GG, TB, PB and TH edited/modified the manuscript. All authors commented on the manuscript.

Ethics declarations

Competing interests: The authors declare no competing financial interests.

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Editors

Reviewing Editor

Amy Andreotti

Iowa State University, Ames, United States of America

Senior Editor

Amy Andreotti

Iowa State University, Ames, United States of America

Reviewer #1 (Public Review):

The model of phosphotransfer from Y169 IKK to S32 IkBa is compelling and an important new contribution to the field. In fact, this model will not be without controversy, and publishing the work will catalyze follow-up studies for this kinase and others as well. As such, I am supportive of this paper, though I do also suggest some shortening and modification.

Generally, the paper is well written, but several figures should be quantified, and experimental reproducibility is not always clear. The first 4 figures are slow-going and could be condensed to show the key points, so that the reader gets to Figures 6 and 7 which contain the "meat" of the paper.

<https://doi.org/10.7554/eLife.98009.1.sa2>

Reviewer #2 (Public Review):

Summary:

The authors investigate the phosphotransfer capacity of Ser/Thr kinase I κ B kinase (IKK), a mediator of cellular inflammation signaling. Canonically, IKK activity is promoted by

activation loop phosphorylation at Ser177/Ser181. Active IKK can then unleash NF- κ B signaling by phosphorylating repressor I κ B α at residues Ser32/Ser26. Noting the reports of other IKK phosphorylation sites, the authors explore the extent of autophosphorylation.

Semi-phosphorylated IKK purified from Sf9 cells, exhibits the capacity for further autophosphorylation. Anti-phosphotyrosine immunoblotting indicated unexpected tyrosine phosphorylation. Contaminating kinase activity was tested by generating a kinase-dead K44M variant, supporting the notion that the unexpected phosphorylation was IKK-dependent. In addition, the observed phosphotyrosine signal required phosphorylated IKK activation loop serines.

Two candidate IKK tyrosines were examined as the source of the phosphotyrosine immunoblotting signal. Activation loop residues Tyr169 and Tyr188 were each rendered non-phosphorylatable by mutation to Phe. The Tyr variants decreased both autophosphorylation and phosphotransfer to I κ B α . Likewise, Y169F and Y188F IKK2 variants immunoprecipitated from TNF α -stimulated cells also exhibited reduced activity in vitro.

The authors further focus on Tyr169 phosphorylation, proposing a role as a phospho-sink capable of phosphotransfer to I κ B α substrate. This model is reminiscent of the bacterial two-component signaling phosphotransfer from phosphohistidine to aspartate. Efforts are made to phosphorylate IKK2 and remove ATP to assess the capacity for phosphotransfer. Phosphorylation of I κ B α is observed after ATP removal, although there are ambiguous requirements for ADP.

Strengths:

Ultimately, the authors draw together the lines of evidence for IKK2 phosphotyrosine and ATP-independent phosphotransfer to develop a novel model for IKK2-mediated phosphorylation of I κ B α . The model suggests that IKK activation loop Ser phosphorylation primes the kinase for tyrosine autophosphorylation. With the assumption that IKK retains the bound ADP, the phosphotyrosine is conformationally available to relay the phosphate to I κ B α substrate. The authors are clearly aware of the high burden of evidence required for this unusual proposed mechanism. Indeed, many possible artifacts (e.g., contaminating kinases or ATP) are anticipated and control experiments are included to address many of these concerns. Taken together, the observations are thought-provoking, and I look forward to seeing this model tested in a cellular system.

Weaknesses:

It seems that the analysis hinges on the fidelity of pan-specific phosphotyrosine antibodies.

The analysis often returns to the notion that tyrosine phosphorylation(s) (and critical active site Lys44) dictate IKK2 substrate specificity, but evidence for this seems diffuse and indirect. This is an especially difficult claim to make with in vitro assays, omitting the context of other cellular specificity determinants (e.g., localization, scaffolding, phosphatases).

Multiple phosphorylated tyrosines in IKK2 were apparently identified by mass spectrometric analyses, but the data and methods are not described. It is common to find non-physiological post-translational modifications in over-expressed proteins from recombinant sources. Are these IKK2 phosphotyrosines evident by MS in IKK2 immunoprecipitated from TNF α -stimulated cells? Identifying IKK2 phosphotyrosine sites from cells would be especially helpful in supporting the proposed model.

<https://doi.org/10.7554/eLife.98009.1.sa1>

Reviewer #3 (Public Review):**Summary:**

The authors investigate the kinase activity of IKK2, a crucial regulator of inflammatory cell signaling. They describe a novel tyrosine kinase activity of this well-studied enzyme and a highly unusual phosphotransfer from phosphorylated IKK2 onto substrate proteins in the absence of ATP as a substrate.

Strengths:

The authors provide an extensive biochemical characterization of the processes with recombinant protein, western blot, autoradiography, and protein engineering.

Weaknesses:

The identity and purity of the used proteins is not clear. Since the findings are so unexpected and potentially of wide-reaching interest - this is a weakness. Similar specific detection of phospho-Ser/Thr vs phospho-Tyr relies largely on antibodies which can have varying degrees of specificity.

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