IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

Gillespie, ME., Shamovsky, V.

European Bioinformatics Institute, New York University Langone Medical Center, Ontario Institute for Cancer Research, Oregon Health and Science University.

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25/11/2019
Introduction

Reactome is open-source, open access, manually curated and peer-reviewed pathway database. Pathway annotations are authored by expert biologists, in collaboration with Reactome editorial staff and cross-referenced to many bioinformatics databases. A system of evidence tracking ensures that all assertions are backed up by the primary literature. Reactome is used by clinicians, geneticists, genomics researchers, and molecular biologists to interpret the results of high-throughput experimental studies, by bioinformaticians seeking to develop novel algorithms for mining knowledge from genomic studies, and by systems biologists building predictive models of normal and disease variant pathways.

The development of Reactome is supported by grants from the US National Institutes of Health (P41 HG003751), University of Toronto (CFREF Medicine by Design), European Union (EU STRP, EMI-CD), and the European Molecular Biology Laboratory (EBI Industry program).

Literature references


Reactome database release: 70

This document contains 1 pathway and 5 reactions (see Table of Contents)
IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

Stable identifier: R-HSA-975163

Compartments: cytosol, endosome membrane

Although IRAK-1 was originally thought to be a key mediator of TRAF6 activation in the IL1R/TLR signaling (Dong W et al. 2006), recent studies showed that IRAK-2, but not IRAK-1, led to TRAF6 polyubiquitination (Keating SE et al 2007). IRAK-2 loss-of-function mutants, with mutated TRAF6-binding motifs, could no longer activate NF-kB and could no longer stimulate TRAF-6 ubiquitination (Keating SE et al 2007). Furthermore, the proxyvirus protein A52 - an inhibitor of all IL-1R/TLR pathways to NF-kB activation, was found to interact with both IRAK-2 and TRAF6, but not IRAK-1. Further work showed that A52 inhibits IRAK-2 functions, whereas association with TRAF6 results in A52-induced MAPK activation. The strong inhibition effect of A52 was also observed on the TLR3-NFkB axis and this observation led to the discovery that IRAK-2 is recruited to TLR3 to activate NF-kB (Keating SE et al 2007). Thus, A52 possibly inhibits MyD88-independent TLR3 pathways to NF-kB via targeting IRAK-2 as it does for other IL-1R/TLR pathways, although it remains unclear how IRAK-2 is involved in TLR3 signaling.

IRAK-2 was shown to have two TRAF6 binding motifs that are responsible for initiating TRAF6 signaling transduction (Ye H et al 2002).

Literature references


## Editions

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IRAK2 induces TRAF6 oligomerization initiated from endosomal compartments

Location: IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

Stable identifier: R-HSA-975185

Type: binding

Compartments: endosome membrane, cytosol

The mechanism by which IRAK-2 induces TRAF6 E3 ligase activity remains to be deciphered, but one possibility is that IRAK-2 may direct TRAF6 oligomerization.

Followed by: Auto ubiquitination of oligo-TRAF6 bound to p-IRAK2 at endosome membrane

Literature references


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Auto ubiquitination of oligo-TRAF6 bound to p-IRAK2 at endosome membrane

Location: IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

Stable identifier: R-HSA-975147

Type: omitted

Compartments: endosome membrane, cytosol

TRAF6 possesses ubiquitin ligase activity and undergoes K-63-linked auto-ubiquitination after its oligomerization. In the first step, ubiquitin is activated by an E1 ubiquitin activating enzyme. The activated ubiquitin is transferred to a E2 conjugating enzyme (a heterodimer of proteins Ubc13 and Uev1A) forming the E2-Ub thioester. Finally, in the presence of ubiquitin-protein ligase E3 (TRAF6, a RING-domain E3), ubiquitin is attached to the target protein (TRAF6 on residue Lysine 124) through an isopeptide bond between the C-terminus of ubiquitin and the epsilon-amino group of a lysine residue in the target protein. In contrast to K-48-linked ubiquitination that leads to the proteosomal degradation of the target protein, K-63-linked polyubiquitin chains act as a scaffold to assemble protein kinase complexes and mediate their activation through proteosome-independent mechanisms. This K63 polyubiquitinated TRAF6 activates the TAK1 kinase complex.

Preceded by: IRAK2 induces TRAF6 oligomerization initiated from endosomal compartments

Followed by: Activated TRAF6 synthesizes unanchored polyubiquitin chains upon TLR stimulation, Activated TRAF6:p-IRAK2 interacts with TAK1 complex upon TLR7/8 or 9 stimulation

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Activated TRAF6 synthesizes unanchored polyubiquitin chains upon TLR stimulation

**Location:** IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

**Stable identifier:** R-HSA-450358

**Type:** transition

**Compartments:** cytosol, endosome membrane

Polyubiquitinated TRAF6 (as E3 ubiquitin ligase) generates free K63-linked polyubiquitin chains that non-covalently associate with ubiquitin receptors of TAB2/TAB3 regulatory proteins of the TAK1 complex, leading to the activation of the TAK1 kinase.

**Preceded by:** Auto ubiquitination of oligo-TRAF6 bound to p-IRAK2 at endosome membrane

**Followed by:** Activated TRAF6:p-IRAK2 interacts with TAK1 complex upon TLR7/8 or 9 stimulation

**Literature references**


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Activated TRAF6:p-IRAK2 interacts with TAK1 complex upon TLR7/8 or 9 stimulation

**Location:** IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

**Stable identifier:** R-HSA-975097

**Type:** binding

**Compartments:** endosome membrane, cytosol

TAK1-binding protein 2 (TAB2) and/or TAB3, as part of a complex that also contains TAK1 and TAB1, binds polyubiquitinated TRAF6. The TAB2 and TAB3 regulatory subunits of the TAK1 complex contain C-terminal Npl4 zinc finger (NZF) motifs that recognize with Lys63-pUb chains (Kanayama et al. 2004). The recognition mechanism is specific for Lys63-linked ubiquitin chains [Kulathu Y et al 2009]. TAK1 can be activated by unattached Lys63-polyubiquitinated chains when TRAF6 has no detectable polyubiquitination (Xia et al. 2009) and thus the synthesis of these chains by TRAF6 may be the signal transduction mechanism.

**Preceded by:** Activated TRAF6 synthesizes unanchored polyubiquitin chains upon TLR stimulation, Auto ubiquitination of oligo-TRAF6 bound to p-IRAK2 at endosome membrane

**Followed by:** Auto phosphorylation of TAK1 bound to p-IRAK2:pUb oligo-TRAF6: free K63 pUb: TAB1:TAB2/TAB3 upon TLR7/8 or 9 activation

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[https://reactome.org](https://reactome.org)
Auto phosphorylation of TAK1 bound to p-IRAK2:pUb oligo-TRAF6: free K63 pUb:TAB1:TAB2/TAB3 upon TLR7/8 or 9 activation

**Location:** IRAK2 mediated activation of TAK1 complex upon TLR7/8 or 9 stimulation

**Stable identifier:** R-HSA-975103

**Type:** transition

**Compartments:** endosome membrane, cytosol

The TAK1 complex consists of Transforming growth factor-beta (TGFβ)-activated kinase (TAK1) and TAK1-binding protein 1 (TAB1), TAB2 and TAB3. TAK1 requires TAB1 for its kinase activity (Shibuya et al. 1996, Sakurai et al. 2000). TAB1 promotes TAK1 autophosphorylation at the kinase activation lobe, probably through an allosteric mechanism (Brown et al. 2005, Ono et al. 2001). The TAK1 complex is regulated by polyubiquitination. Binding of TAB2 and TAB3 to Lys63-linked polyubiquitin chains leads to the activation of TAK1 by an uncertain mechanism. Binding of multiple TAK1 complexes to the same polyubiquitin chain may promote oligomerization of TAK1, facilitating TAK1 autophosphorylation and subsequent activation of its kinase activity (Kishimoto et al. 2000). The binding of TAB2/3 to polyubiquitinated TRAF6 may facilitate polyubiquitination of TAB2/3 by TRAF6 (Ishitani et al. 2003), which might result in conformational changes within the TAK1 complex that lead to TAK1 activation. Another possibility is that TAB2/3 may recruit the IKK complex by binding to ubiquitinated NEMO; polyubiquitin chains may function as a scaffold for higher order signaling complexes that allow interaction between TAK1 and IKK (Kanayama et al. 2004).

**Preceded by:** Activated TRAF6:p-IRAK2 interacts with TAK1 complex upon TLR7/8 or 9 stimulation

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