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Phospho-TAK1 (S439) Antibody

Rabbit mAb Catalog # AP92813

Product Information

Application WB, IP Primary Accession 043318

Reactivity Rat, Human, Mouse

Clonality Monoclonal

Other Names MAP3K 7; MEKK7; Mitogen activated protein kinase kinase kinase 7; TAK1; TGF

beta activated kinase 1; TGF1a;

IsotypeRabbit IgGHostRabbitCalculated MW67196

Additional Information

Dilution WB 1:500~1:2000 IP 1:50 **Purification** Affinity-chromatography

ImmunogenA synthesized peptide derived from human Phospho-TAK1 (S439) **Description**Component of a protein kinase signal transduction cascade. Mediator of

TRAF6 and TGF-beta signal transduction. Activates IKBKB and MAPK8 in response to TRAF6 signaling. Stimulates NF-kappa-B activation and the p38 MAPK pathway. In osmotic stress signaling, plays a major role in the

activation of MAPK8/JNK, but not that of NF-kappa-B.

Storage Condition and Buffer Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium

azide and 50% glycerol. Store at +4°C short term. Store at -20°C long term.

Avoid freeze / thaw cycle.

Protein Information

Name MAP3K7 {ECO:0000303 | PubMed:28397838,

ECO:0000312 | HGNC:HGNC:6859}

Function Serine/threonine kinase which acts as an essential component of the MAP

kinase signal transduction pathway (PubMed: 10094049, PubMed: 11460167,

PubMed: 12589052, PubMed: 16845370, PubMed: 16893890,

PubMed:<u>21512573</u>, PubMed:<u>8663074</u>, PubMed:<u>9079627</u>). Plays an important

role in the cascades of cellular responses evoked by changes in the environment (PubMed:10094049, PubMed:11460167, PubMed:12589052, PubMed:16845370, PubMed:16893890, PubMed:21512573, PubMed:8663074, PubMed:9079627). Mediates signal transduction of TRAF6, various cytokines including interleukin-1 (IL-1), transforming growth factor- beta (TGFB), TGFB-related factors like BMP2 and BMP4, toll-like receptors (TLR), tumor necrosis factor receptor CD40 and B-cell receptor (BCR) (PubMed:16893890, PubMed:9079627). Once activated, acts as an upstream activator of the

MKK/JNK signal transduction cascade and the p38 MAPK signal transduction cascade through the phosphorylation and activation of several MAP kinase kinases like MAP2K1/MEK1, MAP2K3/MKK3, MAP2K6/MKK6 and MAP2K7/MKK7 (PubMed:11460167, PubMed:8663074). These MAP2Ks in turn activate p38 MAPKs and c-jun N- terminal kinases (INKs); both p38 MAPK and JNK pathways control the transcription factors activator protein-1 (AP-1) (PubMed: 11460167, PubMed: 12589052, PubMed: 8663074). Independently of MAP2Ks and p38 MAPKs, acts as a key activator of NF-kappa-B by promoting activation of the I-kappa-B-kinase (IKK) core complex (PubMed: 12589052, PubMed:8663074). Mechanistically, recruited to polyubiquitin chains of RIPK2 and IKBKG/NEMO via TAB2/MAP3K7IP2 and TAB3/MAP3K7IP3, and catalyzes phosphorylation and activation of IKBKB/IKKB component of the IKK complex, leading to NF-kappa-B activation (PubMed: 10094049, PubMed: 11460167). In osmotic stress signaling, plays a major role in the activation of MAPK8/JNK1, but not that of NF-kappa-B (PubMed: 16893890). Promotes TRIM5 capsid-specific restriction activity (PubMed: 21512573). Phosphorylates RIPK1 at 'Ser-321' which positively regulates RIPK1 interaction with RIPK3 to promote necroptosis but negatively regulates RIPK1 kinase activity and its interaction with FADD to mediate apoptosis (By similarity). Phosphorylates STING1 in response to cGAMP-activation, promoting association between STEEP1 and STING1 and STING1 translocation to COPII vesicles (PubMed:37832545).

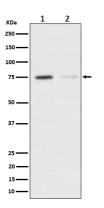
Cellular Location

Cytoplasm. Cell membrane; Peripheral membrane protein; Cytoplasmic side. Note=Although the majority of MAP3K7/TAK1 is found in the cytosol, when complexed with TAB1/MAP3K7IP1 and TAB2/MAP3K7IP2, it is also localized at the cell membrane

Tissue Location

Isoform 1A is the most abundant in ovary, skeletal muscle, spleen and blood mononuclear cells. Isoform 1B is highly expressed in brain, kidney and small intestine. Isoform 1C is the major form in prostate. Isoform 1D is the less abundant form

Images



Western blot analysis of Phospho-TAK1 (S439) expression in (1) HeLa treated with Calyculin A + IL-1 beta cell lysate; (2) Untreated.

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