

TAK1 Antibody (C-term)
Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP7953A**Specification**

TAK1 Antibody (C-term) - Product Information

Application	WB, IHC-P,E
Primary Accession	O43318
Other Accession	P0C8E4 , Q62073 , A2VDU3
Reactivity	Human
Predicted	Bovine, Mouse, Rat
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	67196
Antigen Region	574-606

TAK1 Antibody (C-term) - Additional Information**Gene ID** 6885**Other Names**

Mitogen-activated protein kinase kinase kinase 7, Transforming growth factor-beta-activated kinase 1, TGF-beta-activated kinase 1, MAP3K7, TAK1

Target/Specificity

This TAK1 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 574-606 amino acids from the C-terminal region of human TAK1.

Dilution

WB~~1:1000
IHC-P~~1:50~100

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

TAK1 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

TAK1 Antibody (C-term) - 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:[21512573](#), PubMed:[8663074](#), PubMed:[9079627](#)). 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 (JNKs); 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

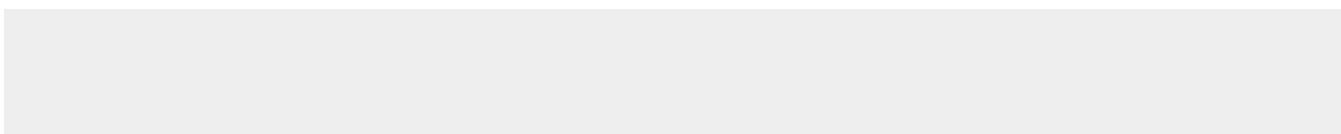
Isoform 1A is the most abundant in ovary, skeletal muscle, spleen and blood mononuclear cells. Isoform 1B is highly expressed in brain, kidney and small intestine. Isoform 1C is the major form in prostate. Isoform 1D is the less abundant form

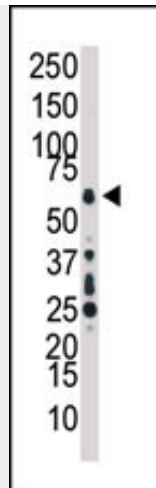
TAK1 Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

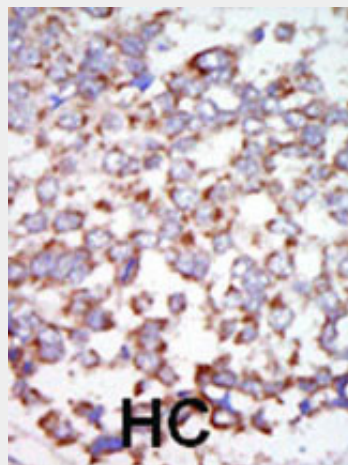
- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

TAK1 Antibody (C-term) - Images

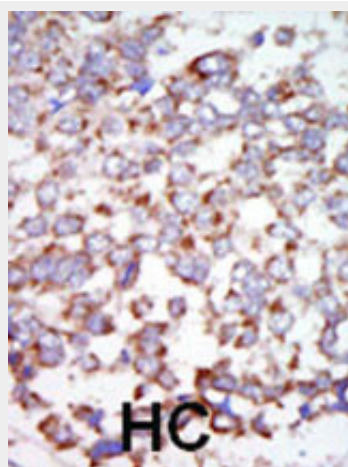




Western blot analysis of anti-TAK1 Pab (Cat. #AP7953a) in HL60 cell lysate. TAK1 (arrow) was detected using purified Pab. Secondary HRP-anti-rabbit was used for signal visualization with chemiluminescence.



Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by AEC staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated. BC = breast carcinoma; HC = hepatocarcinoma.



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TAK1 Antibody (C-term) - Background

TAK1 is a member of the serine/threonine protein kinase family. This kinase mediates the signaling transduction induced by TGF beta and morphogenetic protein (BMP), and controls a variety of cell functions including transcription regulation and apoptosis. In response to IL-1, this protein forms a kinase complex including TRAF6, MAP3K7P1/TAB1 and MAP3K7P2/TAB2; this complex is required for the activation of nuclear factor kappa B. This kinase can also activate MAPK8/JNK, MAP2K4/MKK4, and thus plays a role in the cell response to environmental stresses.

TAK1 Antibody (C-term) - References

- Cheung, P.C., et al., EMBO J. 22(21):5793-5805 (2003).
Sakurai, H., et al., J. Biol. Chem. 278(38):36916-36923 (2003).
Ono, K., et al., Biochem. Biophys. Res. Commun. 307(2):332-337 (2003).
Edlund, S., et al., Mol. Biol. Cell 14(2):529-544 (2003).
Takaesu, G., et al., J. Mol. Biol. 326(1):105-115 (2003).