

Division of Signal Tranduction Therapy

Standard Operating Procedure

Preparation of active TAK1 [1 - 303] / TAB1 [437 – 504] Fusion

Enzyme description:- TAK1 [1 – 303] / TAB1 [437 – 504] Fusion

Clone number:- DU 753

Source:- Recombinant

Expression system:- Baculovirus expression vector system

Tag:- N-terminal His(6) tag

Purification method:- Ni²⁺-NTA agarose

Calculated molecular mass:-

Monoisotopic 44,488.72 daltons

Average Mass 44,517.52 daltons

[cysteines reduced, methionines have not been oxidised]

Theoretical pI:- 5.93

Purity:- 80 %

Activation protocol:- Constitutively active

Enzyme storage buffer:-

50 mM Tris-HCl pH 7.5, 270 mM sucrose, 150 mM NaCl, 0.1 mM EGTA,
0.1 % 2-mercaptoethanol, 0.02 % Brij-35, 0.2 mM PMSF, 1 mM Benzamidine.

Storage temperature:- -70 °C

Assay buffer:-

50 mM Tris-HCl pH 7.5, 0.1mM EGTA, 0.1 % 2-mercaptoethanol, 10 mM MgAc

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Clone Data Sheet

TAK1 [1 - 303] / TAB1 [437 – 504] Fusion

<u>Protein</u>	TAK1 [1 – 303] / TAB1 [437 – 504] Fusion
<u>Clone Number</u>	DU 753
<u>Species</u>	Human
<u>Accession number</u>	NM_003188 (TAK1) and NM_006116 (TAB1)
<u>Tags</u>	N-terminal His(6)
<u>Baculovirus expressed protein</u>	MSYYHHHHHDYDIPPTENLYFQGAMGS <u>MSTASAASSSSSSAGEMIEA PSQVLNFEEIDYKEIEVEEVVGRGAFGVVCKAKWRAKDVAIKQIESESE RKA</u> FIVE <u>LQLSRVNHPNIVKLYGACLNPVCLVM</u> EYAEGGS <u>LYNV</u> LHG <u>A EPLPYYTAAHAMSWCLOCSQGVAYLHSMQPKALIHRDLKPPNLLVAGG TVLKICDFGTACDIQTHMTNNKGSAAWMAPEVFE</u> GSNY <u>SEKCDVF</u> SWGI ILWEVITRRKPFDEIGGP <u>AFRIMWAHNGTRP</u> LIKNL <u>PKPIESLMTRC WSKDPSQRPSMEEIVKIMTHLMRYFPGADEPLQYPCQ</u> QSPTLT <u>Q</u> STNT HT <u>QSSSSSSDGGLFRSRPAHSLPPGEDGRVEPYVDFAEFYRLWSVDHGE QSVVTAP</u>
<u>Native sequence</u>	Amino acids M1 – Q303 of human TAK1 [full length protein ends at residue S579] followed by amino acids Q437 – P504 (end) of human TAB1. Residue M29 of the fusion protein is equilivalent to M1 of TAK1 and residue Q332 of the fusion protein is equilivalent to Q332 of TAB1. The His(6) tag is located at residues 5 – 10.
<u>Protease cleavage</u>	rTEV (<u>ENLYFQG</u>) residues 18 - 24
<u>Cloning sites</u>	<i>Bam</i> H1 site of pFastBAC HTb

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Nucleotide
sequence of
insert

ATGTCTACAGCCTCTGCCGCCTCCTCCTCGTCTCGGCCGGTG
AGATGATCGAAGCCCTTCCAGTCCTCAACTTGAAAGAGATCGACTA
CAAGGAGATCGAGGTGGAAGAGGTTGGAAGAGGAGCCTTGGAGTT
GTTTGCAAAGCTAAGTGGAGAGCAAAAGATGTTGCTATTAAACAAATAG
AAAGTGAATCTGAGAGGAAAGCGTTATTGTTAGAGCTTCGGCAGTTATC
CCGTGTGAACCATCCTAATATTGTAAGCTTATGGAGCCTGCTGAAT
CCAGTGTGTCTTGTGATGGAATATGCTGAAGGGGGCTTTATATAATG
TGCTGCATGGTGTGAACCATTGCCATATTATACTGCTGCCACGCAAT
GAGTTGGTGTACAGTGTCCAAGGAGTGGCTATCTTCACAGCATG
CAACCCAAAGCGCTAATTACAGGGACCTGAAACCACAAACTACTGC
TGGTTGCAGGGGGGACAGTTCTAAAAATTGTGATTGGTACAGCCTG
TGACATTCAACACATGACCAATAACAAGGGAGTGCTGCTTGGATG
GCACCTGAAGTTTGAAAGGTAGTAATTACAGTAAAAATGTGACGTCT
TCAGCTGGGTATTATTCTTGGAAAGTGATAACCGCTGGAAACCCTT
TGATGAGATTGGTGGCCCAGCTTCCGAATCATGTGGCTGTTACATAAT
GGTACTCGACCACCACTGATAAAAATTACCTAACGCCATTGAGAGCC
TGATGACTCGTTGGTCTAAAGATCCTCCAGCGCCCTCAATGGA
GGAAATTGTAAAATAATGACTCACTGATGCGGTACTTCCAGGAGCA
GATGAGCCATTACAGTATCCTGTCAGCAAAGCCGACCTAACCCCTGC
AGTCCACCAACACGCACACGCAGAGCAGCAGCTCCAGCTGACGGAGG
CCTCTTCGCTCCGGCCCGCCACTCGCTCCGCCCTGGCGAGGACGGT
CGTGTGAGCCCTATGTGGACTTGCTGAGTTACCGCTCTGGAGCG
TGGACCATGGCGAGCAGAGCGTGGTACAGCACCGtag