

Division of Signal Transduction Therapy

Standard Operating Procedure

Preparation of active Sphingosine Kinase 1 [1 - 384]

<u>Enzyme description:-</u>	SPHK1 [1 - 384]
<u>Clone number:-</u>	DU 12303
<u>Source:-</u>	Recombinant
<u>Expression system:-</u>	Baculovirus expression vector system
<u>Tag:-</u>	N-terminal His(6)
<u>Purification method:-</u>	Ni ²⁺ -NTA agarose
<u>Expression level:-</u>	5 mg/L

Calculated molecular mass:-

Monoisotopic 47, 299.12 daltons
Average Mass 47, 329.83 daltons
[cysteines reduced, methionines have not been oxidised]

<u>Theoretical pI:-</u>	6.52
<u>Purity:-</u>	>80 %
<u>Activation protocol:-</u>	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, 0.2 mM PMSF, 1 mM Benzamidine

<u>Storage temperature:-</u>	-70 °C
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<u>Assay:-</u>	Kinase Glo
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Assay buffer:-

50 mM Tris-HCl pH 7.5, 150 mM KCl, 1 mM DTT, 1 mM EGTA, 5 mM MgCl₂

Substrate:-

Sphingosine Final concentration: 10 μM

<u>Specific activity range:-</u>	To be determined
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Clone Data Sheet

Sphingosine Kinase 1 [1 – 384]

<u>Protein</u>	SPHK1 [1 - 384]
<u>Clone number</u>	DU 12303
<u>Species</u>	Human
<u>Accession number</u>	AF200328.1
<u>Tags</u>	N-terminal His(6)
<u>Baculovirus Expressed protein</u>	MSYYHHHHHDYDIPTTENLYFQGAMGSGIQRPSTSTSSLVAAAMPAGG PRGVLPRPCRVLVLLNPRGGKQALQFRSHVQPLLAEEAISFTLMLTE RRNHARELVRSEELGRWDALVVMSGDGLMHEVVNGLMERPDWETAIQP LCSLPAGSGNALAASLNHYAGYEQVTNEDLLTNCTLLLCRLLSPMNL SLHTASGLRFLFSVLSLAWGFIADVLESEKYRRLGEMRFTLGTFRLAA LRTYRGLAYLPVGRVGSKTPASPVVVQGPVDAHLVPLEEPVPSHWT VPDEDFVLVLLALLHSHLGSEMFAAPMGRCAAGVMHLFYVRAGVSRAMLL RLFLAMEKGRHMEYECPYLVYVPVVAFRLEPKDGKGVFAVDGELMVSEA VQGQVHPNYFWMVSGCVEPPPSWKPOQMPPEEPL
<u>Native sequence</u>	Amino acids M1 – L384 (end) of human SPHK1. Residue M44 of the fusion protein is equivalent to M1 of the native enzyme. 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>Not1</i> of pFastBac HTb

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**Nucleotide
of sequence of
insert**

gcggccgcgATGGATCCAGCGGGCGGCCCCCGGGGCGTGCTCCCGCGGC
CCTGCCGCGTGCTGGTGCTGCTGAACCCGCGCGGCGGCAAGGGCAAGGC
CTTGACAGCTCTTCCGGAGTCACGTGCAGCCCCTTTTGGCTGAGGCTGAA
ATCTCCTTCACGCTGATGCTCACTGAGCGGCGGAACCACGCGGGGAGC
TGGTGCGGTTCGGAGGAGCTGGGCCGCTGGGACGCTCTGGTGGTCATGTC
TGGAGACGGGCTGATGCACGAGGTGGTGAACGGGCTCATGGAGCGGCCT
GACTGGGAGACCGCCATCCAGAAGCCCCTGTGTAGCCTCCAGCAGGCT
CTGGCAACGCGCTGGCAGCTTCCTTGAACCATTATGCTGGCTATGAGCA
GGTCACCAATGAAGACCTCCTGACCAACTGCACGCTATTGCTGTGCCG
CGGCTGCTGTACCCATGAACCTGCTGTCTCTGCACACGGCTTCGGGGC
TGCGCCTCTTCTCTGTGCTCAGCCTGGCCTGGGGCTTCATTGCTGATGT
GGACCTAGAGAGTGAGAAGTATCGGCGTCTGGGGGAGATGCGCTTCACT
CTGGGCACCTTCCTGCGTCTGGCAGCCCTGCGCACCTACCGCGGCCGAC
TGGCCTACCTCCCTGTAGGAAGAGTGGGTTCCAAGACACCTGCCTCCCC
CGTTGTGGTCCAGCAGGGCCCCGGTAGATGCACACCTTGTGCCACTGGAG
GAGCCAGTGCCCTCTCACTGGACAGTGGTGCCCGACGAGGACTTTGTGC
TAGTCCTGGCACTGCTGCACTCGCACCTGGGCAGTGAGATGTTTGCTGC
ACCCATGGGCCGCTGTGCAGCTGGCGTCATGCATCTGTTCTACGTGCGG
GCGGGAGTGTCTCGTGCCATGCTGCTGCGCCTCTTCTGGCCATGGAGA
AGGGCAGGCATATGGAGTATGAATGCCCTACTTGGTATATGTGCCCGT
GGTCGCCTTCCGTTTGGAGCCCAAGGATGGGAAAGGTGTGTTTGCAGTG
GATGGGGAATTGATGGTTAGCGAGGCCGTGCAGGGCCAGGTGCACCCAA
ACTACTTCTGGATGGTCAGCGGTTGCGTGGAGCCCCCGCCAGCTGGAA
GCCCCAGCAGATGCCACCGCCAGAAGAGCCCTTAtgagcggccgc