

Division of Signal Tranduction Therapy

Standard Operating Procedure

Preparation of active Choline Kinase Beta [1 – 395]

Enzyme description:- CHKB [1 - 395]

Clone number:- DU 19590

Source:- Recombinant

Expression system:- *E.coli*

Tag:- N-terminal GST

Purification method:- GSH Sepharose

Calculated molecular mass:-

Monoisotopic 74, 139.13 daltons

Average Mass 74, 186.83 daltons

[cysteines reduced, methionines have not been oxidised

Theoretical pI:- 5.48

Purity:- >75 %

Activation protocol:- Constitutively active

Enzyme storage buffer:-

25 mM glycine-NaOH pH 8.5, 67 mM KCl, 2 mM EDTA, 270 mM Sucrose, 0.1 % B-mercaptoethanol 1 mM benzamidine, 0.2 mM PMSF

Storage temperature:- -70 °C

Assay:- ADP Glo

Assay buffer:-

12.5mM Glycine-NaOH (pH 8.5), 50mM KCl, 2.5mM MgCl₂

Substrate:-

Choline chloride Final concentration: 0.2 mM

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Clone Data Sheet

Choline Kinase Beta [1 - 395]

<u>Protein</u>	CHKB [1 - 395]
<u>Clone number</u>	DU 19590
<u>Species</u>	Human
<u>Accession number</u>	NM_005198
<u>Tags</u>	N-terminal GST
<u>Bacterially expressed protein</u>	MSPILGYWKIKGLVQPTRLLEEKYEEHLYERDEGDKWRNKKFELG LEFPNLPPYYIDGDVKLTQSMAIIRYIADKHNLGGCPKERAETSMLEGA VLDIERYGVSRIAYSKDFETLKVDFLSKLPEMLKMFEDRLCHKTYLNGDH VTHPDFMLYDALDVLYMDPMCLDAFPKLVCFKKRIEAIPQIDKYLKSS KYIAWPLQGWQATFGGGDHPPKSDLEVLFQGPLGSMEPSPSEGSGAQPG LGPGRARAMAAEATAVAGSGAVGGCLAKDGLQOSKCPDTTPKRRASSL SRDAERRAYQWCREYLGGAWRRVQPEELRVYPVSGGLSNLLFRCSLPDH LPSVGEEPREVLLRLYGAILQGVDSLVLSEVMFAILAERSLGPOLYGVF PEGRLEQYIPSRPLKTQELREPVLSSAAIATKMAQFHGMEMPFTEPHWL FGTMERYLKQIQDLPPTGLPEMNLEMYSLKDEMGNLRKLLESTPSPVV FCHNDIQEGNILLSEHENADSMLVDFEYSSSYNRGFDIGNHFCEWVY DYTHEEWPFYKARPTDYPTQEQLHFIRHYLAEAKKGETLSQEEQRKLE EDLLVEVSRYALASHFFWGLWSILQASMSTIEFGYLDYAQSRFQFYFQQ KGQLTSHSSS
<u>Native sequence</u>	Amino acids M1 – 395 (end) of human Choline Kinase Beta. Residue M232 of the fusion protein is equivalent to M1 of the native enzyme. The GST tag is located at residues 1 – 220.
<u>Protease cleavage</u>	PreScission (<u>LEVLFQGP</u>) residues 221 - 228
<u>Cloning sites</u>	<i>Bam</i> H1 and <i>Not</i> 1 site of pGEX 6P-1

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<u>Nucleotide</u>	
<u>Sequence of insert</u>	
	ggatccATGGAACCGAGCCGTCCGAAGGGAGCGGAGCGCAGCCTGGCC TGGGGCCCGTCGAGCCGCCATGGCGGCCAGGGCAGACAGCTGTGGC CGGAAGCAGGGCTGTTGGCGGCTGCCTGGCAAAGACGGCTTGAGCAG TCTAAGTGCCCGAACACTACCCCCAAACGGCGCGCCCTCGCTGCTGT CGCGTGACGCCGAGCGCCGAGCCTACCAATGGTGCAGGGAGTACTTGGG CGGGGCCTGGCGCCGAGTGCAGCCCAGGGAGCTGAGGGTTACCCGTG AGCGGAGGCCTCAGCAACCTGCTCTCCGCTGCTCGCTCCGGACCAC TGCCCAGCGTTGGCGAGGGAGCCCGGGAGGTGCTTCTGCGGCTGTACGG AGCCATCTTGCAAGGGCGTGGACTCCCTGGTCTAGAAAGCGTGTACGG GCCATACTTGCGGAGCGGTGGCTGGGGCCCCAGCTGTACGGAGTCTTCC CAGAGGGCCGGCTGGAACAGTACATCCAAGTCGGCATTGAAAGAACTCA AGAGCTTCGAGAGCCAGTGTGTCAGCAGCCATTGCCACGAAGATGGCG CAATTCATGGCATGGAGATGCCTTCACCAAGGGAGCCCCACTGGCTGT TTGGGACCATGGAGCGGTACCTAAAACAGATCCAGGACCTGCCCCAAC TGGCCTCCCTGAGATGAACCTGCTGGAGATGTACAGCCTGAAGGATGAG ATGGGCAACCTCAGGAAGTTACTAGAGTCTACCCATGCCAGTCGTCT TCTGCCACAATGACATCCAGGAAGGAAACATCTTGCTGCTCTCAGAGCC AGAAAATGCTGACAGCCTCATGCTGGACTTCGAGTACAGCAGTTAT AACTATAAGGGCTTGACATTGGGAACCATTGTGAGTGGGTTATG ATTATACTCACGAGGAATGGCCTTCTACAAAGCAAGGCCACAGACTA CCCCACTCAAGAACAGCAGTTGCATTTCATTCTGTCATTACCTGGCAGAG GCAAAGAAAGGTGAGACCCCTCTCCAAGAGGAGCAGAGAAAATGGAAG AAGATTTGCTGGTAGAAGTCAGTCGGTATGCTCTGGCATCCATTCTT CTGGGGTCTGTGGTCCATCCTCAGGCATCCATGTCCACCATAAGAATT GGTTACTTGGACTATGCCAGTCTCGGTTCCAGTTCTACTTCAGCAGA AGGGGCAGCTGACCAGTGTCCACTCCTCATCCTGACTCCACCCCTCCCAC TCCTTGGATTCTCCTGGAGCCTCCAGGGCAGGACCTGGAGGGAGGAA CAACGAGCAGAAGGCCCTGGCGACTgagcggccgc