Certificate of Analysis KIT V559D, 10 µg

Recombinant human KIT V559D (544-end) expressed in insect cells

Part Number: A30497 Lot Number: 1861431 Immediate Storage: -80°C Shipping Conditions: dry ice



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Description:

Recombinant human KIT V559D (544-end) was expressed in insect cells using a N-terminal GST tag.

KIT is a proto-oncogene and a type 3 transmembrane receptor for MGF (mast cell growth factor, also known as stem cell factor). KIT was first identified as the cellular homolog of the feline sarcoma viral oncogene v-kit. KIT together with its ligand regulates growth and activation of a variety of hemopoietic and non-hemopoietic cells. Mutations in KIT are associated with gastrointestinal stromal tumors, mast cell disease, acute myelogenous lukemia, and piebaldism. Recently, deregulation of the KIT receptor TK by the prevalent activation loop mutation D816V has served as a focal point in therapeutic strategies aimed at curbing neoplastic mast cell growth.

Accession Number:

The gene accession number for KIT V559D is NP_000213.1.

Specific Activity:

52 nmoles of phosphate transferred to poly [Glu, Tyr] 4:1 substrate per minute per mg of total protein at 30°C.

Concentration:

0.1 mg/mL total protein as measured using the Bradford protein assay with BSA as a standard.

Calculated 1,330 nM.

Aliases:

PBT, SCFR, CD117

Storage and Handling:

For maximum recovery please spin prior to use. Unless noted below, aliquots of the 5 ug, 10ug and 20ug sizes of kinase are not recommended as materials can be used in original packaging until exhausted. For larger sizes, the number of freeze/thaws may be reduced by preparing aliquots, aliquots below 20 μ L are not recommended. **Please never store a kinase diluted**. If properly stored at -80°C, this product is guaranteed for 6 months from date of purchase.

Storage Buffer:

50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 10 mM glutathione, 0.1 mM EDTA, 0.25 mM DTT, 0.1mM PMSF and 25% Glycerol.

QUALITY ASSURANCE

KIT V559D Activity Graph



Dilution Buffer:

5 mM MOPS (pH 7.2), 2.5 mM β -glycerol-phosphate, 4 mM MgCl_2, 2.5 mM MnCl_2, 1 mM EGTA and 0.4 mM EDTA, 0.05 mM DTT and 50 ng/µL BSA.

Assay Conditions:

KIT V559D was pre-diluted in enzyme dilution buffer and assayed in 5 mM MOPS (pH 7.2), 2.5 mM β-glycerol-phosphate, 4 mM MgCl₂, 2.5 mM MnCl₂, 0.4 mM EDTA, 1 mM EGTA, 0.05 mM DTT, with 50 μM ATP, trace [³³P]-γ-ATP and 200 μg/mL poly [Glu, Tyr] 4:1 substrate for 15 minutes at 30°C.

Gel Information for KIT V559D

Page Description: Run on an SDS-PAGE gel and stained with Coomassie[®].

Lane 1: Molecular Weight markers as labeled.

Lane 2: KIT V559D



Purity:

> 70% as determined by a Coomassie[®] blue stained SDS-PAGE gel.

Molecular Weight:

75.3 kDa. Calculated from the protein sequence(s).

5-6268 OF INTE PH#. 760-603-7200 Select option 5, ext. 40266 Email: drugdiscoverytech@intetech.com

Certificate of Analysis

Protein sequence alignment with reference sequence(s)

GenBank Accession Number: NP_000213.1

	MSPILGYWKIKGLVQPTRLLLEYLEEKYEEHLYERDEGDKWRNKKFELGLEFPNLPYYIDGDVKLTQSMAIIRYIADKHNMLGGCPKERAEISMLEGAVL MSPILGYWKIKGLVQPTRLLLEYLEEKYEEHLYERDEGDKWRNKKFELGLEFPNLPYYIDGDVKLTQSMAIIRYIADKHNMLGGCPKERAEISMLEGAVL	
	DIRYGVSRIAYSKDFETLKVDFLSKLPEMLKMFEDRLCHKTYLNGDHVTHPDFMLYDALDVVLYMDPMCLDAFPKLVCFKKRIEAIPQIDKYLKSSKYIA DIRYGVSRIAYSKDFETLKVDFLSKLPEMLKMFEDRLCHKTYLNGDHVTHPDFMLYDALDVVLYMDPMCLDAFPKLVCFKKRIEAIPQIDKYLKSSKYIA	
	WPLQGWQATFGGGDHPPKSD WPLQGWQATFGGGDHPPKSDLVPRGSTYKYLQKPMYEVQWKDVEEINGNNYVYIDPTQLPYDHKWEFPRNRLSFGKTLGAGAFGKVVEATAYGLIKSDAA TYKYLQKPMYEVQWKVVEEINGNNYVYIDPTQLPYDHKWEFPRNRLSFGKTLGAGAFGKVVEATAYGLIKSDAA	
	MTVAVKMLKPSAHLTEREALMSELKVLSYLGNHMNIVNLLGACTIGGPTLVITEYCCYGDLLNFLRRKRDSFICSKQEDHAEAALYKNLLHSKESSCSDS MTVAVKMLKPSAHLTEREALMSELKVLSYLGNHMNIVNLLGACTIGGPTLVITEYCCYGDLLNFLRRKRDSFICSKQEDHAEAALYKNLLHSKESSCSDS	
220 401 175	TNEYMDMKPGVSYVVPTKADKRRSVRIGSYIERDVTPAIMEDDELALDLEDLLSFSYQVAKGMAFLASKNCIHRDLAARNILLTHGRITKICDFGLARDI TNEYMDMKPGVSYVVPTKADKRRSVRIGSYIERDVTPAIMEDDELALDLEDLLSFSYQVAKGMAFLASKNCIHRDLAARNILLTHGRITKICDFGLARDI	
	KNDSNYVVKGNARLPVKWMAPESIFNCVYTFESDVWSYGIFLWELFSLGSSPYPGMPVDSKFYKMIKEGFRMLSPEHAPAEMYDIMKTCWDADPLKRPTF KNDSNYVVKGNARLPVKWMAPESIFNCVYTFESDVWSYGIFLWELFSLGSSPYPGMPVDSKFYKMIKEGFRMLSPEHAPAEMYDIMKTCWDADPLKRPTF	
	KQIVQLIEKQISESTNHIYSNLANCSPNRQKPVVDHSVRINSVGSTASSSQPLLVHDDV KQIVQLIEKQISESTNHIYSNLANCSPNRQKPVVDHSVRINSVGSTASSSQPLLVHDDV	

* highlighted residues denote differences from the reference protein sequence(s).

Michol Heaksedier

Nichole Reaksecker

Date: 05/Jan/2017

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