Recombinant Human SRC/Proto-oncogene c-Src Protein (His Tag)

Catalog No. PDEH100026

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description		
Synonyms	Proto-oncogene tyrosine-protein kinase Src;Proto-oncogene c-Src;pp60c- src;p60-Src;SRC1	
Species	Human	
Expression Host	E.coli	
Sequence	Gly2-Gly287	
Accession	P12931-1	
Calculated Molecular Weight	31.2 kDa	
Observed molecular weight	32.4 kDa	
Tag	N-His	
Bioactivity	Not validated for activity	
Properties		
Purity	> 90 % as determined by reducing SDS-PAGE.	
Endotoxin	Please contact us for more information.	
Storage	Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.	
Shipping	This product is provided as lyophilized powder which is shipped with ice packs.	
Formulation	Lyophilized from sterile PBS, pH 7.4. Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as protectants before lyophilization. Please refer to the specific buffer information in the printed manual.	
Reconstitution	It is recommended that sterile water be added to the vial to prepare a stock solution of 0.5 mg/mL. Concentration is measured by UV-Vis	
TD 4		

Data

kDa 80 60	мк	R
40	-	
30	-	-
20	-	
12	-	

> 90 % as determined by reducing SDS-PAGE.

Background

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Proto-oncogene tyrosine-protein kinase SRC is a hydrophobic protein belonging to the SRC family kinase including nine members that is a family of non-receptor tyrosine kinases. SRC protein may exist in different forms: C-SRC and V-SRC. C-SRC is only activated under certain circumstances where it is required such as growth factor signaling, while V-SRC is constitutively active as opposed to normal SRC (C-SRC). Thus, V-SRC is an instructive example of an oncogene protein kinase whereas C-SRC is a proto-oncogene protein kinase. Inhibition of SRC with NR2A tyrosine phosphorylation mediated by PSD-95 may contribute to the lithium-induced downregulation of NMDA receptor function and provide neuroprotection against excitotoxicity.

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