Proteins

Product Data Sheet

PCSK9 Protein, Human (D374Y, HEK293, His)

Cat. No.: HY-P78663

Synonyms: PCSK9; FH3; HCHOLA3; LDLCQ1; NARC1; PC9

Species: Human Source: HEK293

Accession: Q8NBP7 (Q31-Q692, D374Y)

Gene ID: 255738

Molecular Weight: Approximately 17 & 70 kDa

PROPERTIES

PROPERTIES				
AA Sequence				
	QEDEDGDYEE	LVLALRSEED	GLAEAPEHGT	TATFHRCAKD
	PWRLPGTYVV	VLKEETHLSQ	SERTARRLQA	QAARRGYLTK
	ILHVFHGLLP	GFLVKMSGDL	LELALKLPHV	DYIEEDSSVF
	AQSIPWNLER	ITPPRYRADE	YQPPDGGSLV	EVYLLDTSIQ
	SDHREIEGRV	MVTDFENVPE	EDGTRFHRQA	SKCDSHGTHL
	AGVVSGRDAG	VAKGASMRSL	RVLNCQGKGT	VSGTLIGLEF
	IRKSQLVQPV	GPLVVLLPLA	GGYSRVLNAA	CQRLARAGVV
	LVTAAGNFRD	DACLYSPASA	PEVITVGATN	AQDQPVTLGT
	LGTNFGRCVD	LFAPGEDIIG	ASSYCSTCFV	SQSGTSQAAA
	HVAGIAAMML	SAEPELTLAE	LRQRLIHFSA	KDVINEAWFP
	EDQRVLTPNL	VAALPPSTHG	AGWQLFCRTV	WSAHSGPTRM
	ATAVARCAPD	EELLSCSSFS	RSGKRRGERM	EAQGGKLVCR
	AHNAFGGEGV	YAIARCCLLP	QANCSVHTAP	PAEASMGTRV
	H C H Q Q G H V L T	GCSSHWEVED	LGTHKPPVLR	PRGQPNQCVG
	HREASIHASC	CHAPGLECKV	KEHGIPAPQE	QVTVACEEGW
	TLTGCSALPG	$T\;S\;H\;V\;L\;G\;A\;Y\;A\;V$	DNTCVVRSRD	VSTTGSTSEG
	AVTAVAICCR	SRHLAQASQE	L Q	
Biological Activity	Measured by its ability to inhibit the proliferation of HT-29 human coloncancer cells. The ED ₅₀ for this effect is 16.92 ng/mL, corresponding to a specificactivity is 5.91×10^4 Unit/mg.			
Appearance	Lyophilized powder.			
Formulation	Lyophilized from a 0.2 μm filtered solution of PBS, pH 7.4.			
Endotoxin Level	<1 FILl/ug determined by L	Al mathod		
Elidotoxiii Levet	<1 EU/μg, determined by LAL method.			
Reconsititution	It is not recommended to reconstitute to a concentration less than 100 μ g/mL in ddH ₂ O. For long term storage it is recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).			
Storage & Stability	Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is			

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	recommended to freeze aliquots at -20°C or -80°C for extended storage.
Shipping	Room temperature in continental US; may vary elsewhere.

DESCRIPTION

Background

PCSK9 protein emerges as a pivotal regulator in the intricate orchestration of plasma cholesterol homeostasis. Demonstrating its influence on low-density lipid receptor family members, including the low-density lipoprotein receptor (LDLR), very low-density lipoprotein receptor (VLDLR), apolipoprotein E receptor (LRP1/APOER), and apolipoprotein receptor 2 (LRP8/APOER2), PCSK9 facilitates their degradation within intracellular acidic compartments. Employing a non-proteolytic mechanism, it enhances the hepatic LDLR degradation through a clathrin LDLRAP1/ARH-mediated pathway, possibly impeding LDLR recycling and directing it toward lysosomal degradation. Moreover, PCSK9 exhibits LDLR-independent inhibition of APOB intracellular degradation via the autophagosome/lysosome pathway and plays a role in the disposal of non-acetylated BACE1 intermediates in the early secretory pathway. Notably, it regulates epithelial Na(+) channel (ENaC)-mediated Na(+) absorption by augmenting ENaC proteasomal degradation, and influences neuronal apoptosis through the modulation of LRP8/APOER2 levels and associated anti-apoptotic signaling pathways.

Caution: Product has not been fully validated for medical applications. For research use only.

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