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KNG1 Protein (His tag)





Overview

Quantity:	50 μg
Target:	KNG1
Origin:	Mouse
Source:	HEK-293 Cells
Protein Type:	Recombinant
Biological Activity:	Active
Purification tag / Conjugate:	This KNG1 protein is labelled with His tag.

Product Details

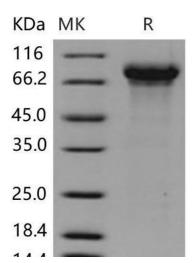
Purpose:	Recombinant Mouse Kininogen-1/KNG1 Protein (His Tag)(Active)
Sequence:	Glu 21-Ser 480
Characteristics:	A DNA sequence encoding the mouse KNG1 isoform 3 (NP_001095882.1) (Glu 21-Ser 480) was expressed, with a polyhistidine tag at the C-terminus and a signal peptide at the N-terminus.
Purity:	> 95 % as determined by SDS-PAGE
Endotoxin Level:	< 1.0 EU per µg of the protein as determined by the LAL method.
Biological Activity Comment:	Measured by its ability to inhibit papain cleavage of a fluorogenic peptide substrate Z-FR-AMC (R&D Systems, Catalog # ES009). The IC50 value is < 5 nM.

Target Details

Target: KNG1

Target Details

Alternative Name:	Kininogen-1/KNG1 (KNG1 Products)
Background:	Background: Kininogen-1, also known as high molecular weight kininogen, williams-Fitzgerald-
	Flaujeac factor, Alpha-2-thiol proteinase inhibitor, Fitzgerald factor, KNG1 and BDK, is a
	secreted protein which contains three cystatin domains. Kininogen-1 / KNG1 is a protein from
	the blood coagulation system as well as the kinin-kallikrein system. It is a protein that adsorbs
	to the surface of biomaterials that come in contact with blood. Kininogen-1 / KNG1 circulates
	throughout the blood and quickly adsorbs to the material surfaces. Kininogen-1 / KNG1 is one
	of the early participants of the intrinsic pathway of coagulation, together with Factor XII
	(Hageman factor) and prekallikrein. Kininogen-1 / KNG1 is one of the kininogens, a class of
	proteins. As with many other coagulation proteins, the protein was initially named after the
	patients in whom deficiency was first observed. When the clinical data were combined, it turned
	out that all patients, in fact, had a deficiency of the same protein. Defects in KNG1 are the
	cause of high molecular weight kininogen deficiency (HMWK deficiency) which is an autosoma
	recessive coagulation defect. Patients with HWMK deficiency do not have a hemorrhagic
	tendency, but they exhibit abnormal surface-mediated activation of fibrinolysis.
	Synonym: Kng
Molecular Weight:	52.5 kDa
NCBI Accession:	NP_001095882
Pathways:	ACE Inhibitor Pathway, Glycosaminoglycan Metabolic Process
Application Details	
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Please refer to the printed manual for detailed information.
Buffer:	Lyophilized from sterile 20 mM Tris, 100 mM NaCl, pH 7.5
Storage:	4 °C,-20 °C,-80 °C
Storage Comment:	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.



Western Blotting

Image 1.