Recombinant Rat ALK-1/ACVRL1 Protein (His &Fc Tag)

Catalog Number: PKSR030416

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

Description		
Species	Rat	
Source	HEK293 Cells-derived Rat ALK-1/ACVRL1 protein Met 1-Ala 118, with an C-terminal	
	His & Fc	
Calculated MW	38.7 kDa	
Observed MW	50-55 kDa	
Accession	NP_071886.1	
Bio-activity	Measured by its ability to inhibit BMP9-induced alkaline phosphatase production by	
	MC3T3E1 mouse chondrogenic cells. David, L. et al. (2007) Blood 109:1953. The ED	
	50 for this effect is typically 40-200 ng/mL in the presence of 2 ng/mL of recombiant	
	human BMP9.	
Properties		
Purity	>90 % as determined by reducing SDS-PAGE.	
Endotoxin	< 1.0 EU per µg of the protein as determined by the LAL method.	
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 $< -20^{\circ}$ 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% Tween 80 are added as protectants	
	before lyophilization.	
	Please refer to the specific buffer information in the printed manual.	
Reconstitution	Please refer to the printed manual for detailed information.	
Data		

KDa	М
116	- 10
66.2	
45.0	- 11
35.0	- 11
25.0	-
18.4 14.4	-

> 90 % as determined by reducing SDS-PAGE.

Background

Elabscience®

Activin A receptor, type II-like 1 (ACVRL1), also known as ALK-1 (activin receptor-like kinase 1), is an endothelialspecific type I receptor of the TGF-beta (transforming growth factor beta) receptor family of ligands. On ligand binding, a heteromeric receptor complex forms consisting of two type II and two type I transmembrane serine/threonine kinases. ACVRL1 protein is expressed in certain blood vessels of kidney, spleen, heart and intestine, serving as an important role during vascular development. Mutations in ACVRL1 gene are associated with hemorrhagic telangiectasia type 2, also known as Rendu-Osler-Weber syndrome 2 and vascular disease.