A Reliable Research Partner in Life Science and Medicine

# Recombinant Human ACVRL1 Protein(His Tag)

Catalog Number: PDMH100250

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

## Description

Species Human

Source Mammalian-derived Human ACVRL1 protein Asp22-Gln118, with an C-terminal His

Calculated MW10.5 kDaObserved MW25-30 kDaAccessionP37023

**Bio-activity** Not validated for activity

### **Properties**

**Purity** > 90% as determined by reducing SDS-PAGE.

Endotoxin <1.0 EU/mg 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°C for 3 months.

ShippingThis product is provided as lyophilized powder which is shipped with ice packs.FormulationLyophilized from a 0.2 μm filtered solution in PBS with 5% Trehalose and 5%

Mannitol.

**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.

#### Data



SDS-PAGE analysis of Human ACVRL1 proteins, 2  $\mu$ g/lane of Recombinant Human ACVRL1 proteins was resolved with an SDS-PAGE under reducing conditions, showing bands at 10.5KD

# Background

This gene encodes a type I cell-surface receptor for the TGF-beta superfamily of ligands. It shares with an other type I receptors a high degree of similarity in serine-threonine kinase subdomains, a glycine-and serine-rich region (called the GS domain) preceding the kinase domain, and a short C-terminal tail. The encoded protein, sometimes termed ALK1, shares similar domain structures with an other closely related ALK or activin receptor-like kinase proteins that form a subfamily of receptor serine/threonine kinases. Mutations in this gene are associated with an hemorrhagic telangiectasia type 2, also known as Rendu-Osler-Weber syndrome 2.

## For Research Use Only

Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017