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

Catalog Number: PKSR030416



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

## **Description**

Synonyms R3;R-3;SETHKIR;MGC91691;Acvrl1

**Species** Rat

Expression Host

Sequence

Met 1-Ala 118

Accession

NP\_071886.1

Calculated Molecular Weight

Observed molecular weight

Tag

HEK293 Cells

Met 1-Ala 118

NP\_071886.1

50-55 kDa

C-His-Fc

**Bioactivity** 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 ED50 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°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

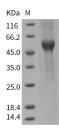
Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as

protectants before lyophilization.

Please refer to the specific buffer information in the printed manual.

**Reconstitution** Please refer to the printed manual for detailed information.

#### Data



> 90 % as determined by reducing SDS-PAGE.

### **Background**

Activin A receptor, type II-like 1 (ACVRL1), also known as ALK-1 (activin receptor-like kinase 1), is an endothelial-specific 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

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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.

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