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## Recombinant Mouse CXADR/CAR Protein (His Tag)

Catalog Number: PKSM040910

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

### **Description**

**Species** Mouse

Source HEK293 Cells-derived Mouse CXADR/CAR protein Met 1-Gly 237, with an C-terminal

His

Calculated MW 25.7 kDa
Observed MW 35 kDa

**Accession** NP\_001020363.1

**Bio-activity** Measured by the ability of the immobilized protein to support the adhesion of mouse

neutrophils. When 5 x  $10^4$  cells/well are added to CXADR-coated plates (4  $\mu$ g/ml and  $100 \mu$ l/well), approximately 20%-40% will adhere specifically after 60 minutes at 37°C.

## **Properties**

**Purity** > 95 % 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

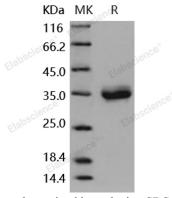
Normally 5% - 8% trehalose, mannitol and 0.01% Tween 80 are added as protectants

before lyophilization.

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

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

## Data



> 95 % as determined by reducing SDS-PAGE.

## Background

## For Research Use Only

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CXADR (coxsackie virus and adenovirus receptor), also known as CAR, is a type I transmembrane glycoprotein belonging to the CTX family of the Ig superfamily, and is essential for normal cardiac development in the mouse. Proposed as a homophilic cell adhesion molecule, CXADR is a component of the epithelial apical junction complex that is essential for the tight junction integrity, and probably involved in transepithelial migration of polymorphonuclear leukocytes (PMN). Mature mouse CXADR structrually comprises a 218 aa extracellular domain (ECD) with a V-type (D1) and a C2-type (D2) Ig-like domain, a 21 aa transmembrane segment and a 107 aa intracellular domain, among which,D1 is thought to be responsible for homodimer formation in trans within tight junctions. The ECD of mouse CXADR shares 9 7%, 90% sequence identity with the corresponding regions of rat, human CXADR.

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