# Recombinant Human CD155 / PVR / NECL5 Protein

Catalog Number: PKSH031867



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

## **Description**

Synonyms CD155;HVED;Necl-5;NECL5;PVS;TAGE4

Species Human

Expression Host HEK293 Cells
Sequence Met 1-Asn 343
Accession NP\_006496.3
Calculated Molecular Weight 35.7 kDa
Observed molecular weight 55-60 kDa

**Bioactivity** Measure by its ability to bind with recombinant human DNAM1 / CD226.

## **Properties**

Purity > 95 % as determined by reducing SDS-PAGE.
 Endotoxin < 1.0 EU per μg as determined by the LAL method.</li>

**Storage** Samples are stable for up to twelve months from date of receipt at -70°C. Store it

under sterile conditions at -20°C to -80°C. It is recommended that the protein be

aliquoted for optimal storage. Avoid repeated freeze-thaw cycles.

**Shipping** In general, recombinant proteins are provided as lyophilized powder which are

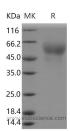
shipped at ambient temperature. Bulk packages of recombinant proteins are provided

as frozen liquid. They are shipped out with blue ice unless customers require

otherwise.

**Formulation** Lyophilized from sterile 50mM Tris, 100mM NaCl, pH 8.0 **Reconstitution** Please refer to the printed manual for detailed information.

#### Data



### **Background**

CD155, commonly known as PVR (poliovirus receptor) and Necl-5 (nectin-like molecule-5), is a type I transmembrane single-span glycoprotein, and belongs to the nectins and nectin-like (Necl) subfamily. CD155 was originally identified based on its ability to mediate the cell attachment and entry of poliovirus (PV), an etiologic agent of the central nervous system disease poliomyelitis. The normal cellular function is in the establishment of intercellular adherens junctions between epithelial cells. CD155 may assist in an efficient humoral immune response generated within the intestinal immune system. It has been demonstrated that CD155 can be recognized and bond by DNAM-1 and CD96 which promote the adhension, migration and NK-cell killing, and thus efficiently prime cell-mediated tumor-specific immunity.

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