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# Recombinant Human SUSD4/Sushi domain-containing protein 4 Protein (Fc Tag)

Catalog No. PKSH030623

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

#### **Description**

Synonyms PRO222
Species Human

Expression Host HEK293 Cells
Sequence Met 1-Phe290
Accession Q5VX71-3
Calculated Molecular Weight 53.8 kDa
Observed molecular weight 67 kDa
Tag C-mFc

**Bioactivity** Not validated for activity

## **Properties**

**Purity** > 85 % as determined by reducing SDS-PAGE.

**Endotoxin**  $< 1.0 \text{ EU per } \mu \text{g of the protein as determined by the LAL method.}$ 

**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

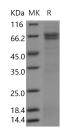
Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as

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



> 85 % as determined by reducing SDS-PAGE.

# **Background**

#### For Research Use Only

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#### **Elabscience Bionovation Inc.**



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SUSD4, also known as sushi domain-containing protein 4, is a hypothetical cell surface protein whose tissue distribution and function are completely unknown. SUSD4 is detectable in murine brains, eyes, spinal cords, and testis but not other tissues. In brains, SUSD4 is highly expressed in the white matter on oligodendrocytes/axons, and in eyes, it is exclusively expressed on the photoreceptor outer segments. In in vitro complement assays, SUSD4 augments the alternative but not the classical pathway of complement activation at the C3 convertase step. SUSD4 deficiency may cause autism or Fryns syndrome, both of which are genetic diseases with severe abnormal neurological development and/or functions.

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