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Recombinant Human MASP1 Protein (His Tag)

Catalog No. PKSH032735

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

Description

Synonyms Mannan-Binding Lectin Serine Protease 1; Complement Factor MASP-3; Mannose-

Binding Lectin-Associated Serine Protease 1;MASP-1;RaRF;Serine Protease

5;MASP1;CRARF;CRARF1;PRSS5

Species Human

HEK293 Cells **Expression Host** His20-Arg728 Sequence P48740-2 Accession Calculated Molecular Weight 80.7 kDa Observed molecular weight 95-120 kDa C-His Tag

Bioactivity Not validated for activity

Properties

Purity > 95 % as determined by reducing SDS-PAGE.

Endotoxin < 1.0 EU per µg of the protein as determined by the LAL method.

Store at < -20°C, stable for 6 months. Please minimize freeze-thaw cycles. **Storage**

Shipping This product is provided as liquid. It is shipped at frozen temperature with blue

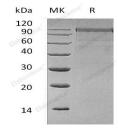
ice/gel packs. Upon receipt, store it immediately at < - 20°C.

Formulation Supplied as a 0.2 µm filtered solution of 20mM Tris-HCl, 200mM NaCl, 10%

Glycerol, pH 8.0.

Reconstitution Not Applicable

Data



> 95 % as determined by reducing SDS-PAGE.

Background

Mannan-Binding Lectin Serine Protease 1 (MASP-1) belongs to the peptidase S1 family. MASP1 contains two CUB domains, one EGF-like domain, one peptidase S1 domain and two Sushi (CCP/SCR) domains. MASP1 is primarily expressed in liver. MASP1 involved in the lectin pathway of the complement, performs a key role in innate immunity by

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recognizing pathogens through patterns of sugar moieties and neutralizing them. MASP1 is synthesized as a zymogen and activated when it complexes with the pathogen recognition molecules of lectin pathway, the mannose-binding lectin and the ficolins. MASP1 is not directly involved in complement activation but may act as an amplifier of complement activation by cleaving complement C2 or by activating another complement serine protease, MASP1 is also able to cleave fibrinogen and factor XIII and may may be involved in coagulation. MASP1 is inhibited by SERPING1 and A2M.

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