

## Product Information

### MemDX™ Antibody Discovery - Mouse Properdin (23-464) Membrane Protein, Partial, -His tag

Cat. No.: **MP0754F**

This product is for research use only and is not intended for diagnostic use.

This membrane protein is Mouse Properdin (23-464). It has been tested in SDS-PAGE. We provide this protein to facilitate your membrane protein antibody discovery and development.

#### Product Specifications

##### Host Species

Mouse

##### Target Protein

Properdin

##### Protein Length

ECD

##### Molecular Weight

The protein has a calculated MW of 49.8 kDa. The protein migrates as 55-60 kDa under reducing (R) condition (SDS-PAGE) due to glycosylation.

##### Sequence

AA Ser 23 - Pro 464 (Accession # P11680-1).

#### Product Description

##### Application

SDS-PAGE

##### Expression Systems

HEK293

##### Tag

His tag at the C-terminus

##### Protein Format

Soluble

##### Form

LYOPH

##### Reconstitution

Please see Certificate of Analysis for specific instructions.

**Endotoxin**

<1.0 EU/μg by the LAL method

**Conjugation**

PE

**Purity**

>95% as determined by SDS-PAGE.

**Buffer**

Lyophilized from 0.22 μm filtered solution in PBS, pH7.4. Normally trehalose is added as protectant before lyophilization.

**Storage**

Stored at lyophilized form at -20°C or lower. Avoid repeated freeze-thaw cycles.

The antigen can be stable for 12 months in lyophilized form after storage at -20°C to -80°C, 3 months under sterile conditions after reconstitution after storage at -80°C.

**Target****Target Protein**

Properdin

**Full Name**

complement factor properdin

**Introduction**

This gene encodes a plasma glycoprotein that positively regulates the alternative complement pathway of the innate immune system. This protein binds to many microbial surfaces and apoptotic cells and stabilizes the C3- and C5-convertase enzyme complexes in a feedback loop that ultimately leads to formation of the membrane attack complex and lysis of the target cell. Mutations in this gene result in two forms of properdin deficiency, which results in high susceptibility to meningococcal infections. Multiple alternatively spliced variants, encoding the same protein, have been identified

**Alternative Names**

Pf; Pfc; BCFG; properdin; complement factor P; properdin factor, complement

**Gene ID**

[18636](#)

**UniProt ID**

[P11680](#)