Proteins



Product Data Sheet

PFDN1 Protein, Mouse (His)

Cat. No.: HY-P76541

Synonyms: Prefoldin subunit 1; PFDN1; PFD1

Species: Mouse Source: E. coli

Q9CQF7 (M1-Q122) Accession:

Gene ID: 67199

Molecular Weight: Approximately 16 kDa

PROPERTIES

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AA	~	മവ	11	Δ	n	~	Δ

MAASVDLELK KAFTELQAKV IDTQQKVKLA DIQIEQLNRT KKHAHLTDTE IMTLVDETNM YEGVGRMFIL QSKEVIHNQL LEKQKIAEEK IKELEQKKSY LERSVKEAED NIREMLMARR

ΑQ

Appearance Lyophilized powder.

Formulation Lyophilized from a 0.2 μm filtered solution of PBS, pH 7.4.

Endotoxin Level <1 EU/µg, determined by LAL method.

Reconsititution It is not recommended to reconstitute to a concentration less than 100 μg/mL in ddH₂O. For long term storage it is

recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).

Storage & Stability Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is

recommended to freeze aliquots at -20°C or -80°C for extended storage.

Shipping Room temperature in continental US; may vary elsewhere.

DESCRIPTION

Background

HLA-DPB1, a pivotal player in the immune response, is instrumental in presenting peptides derived from antigens to CD4 Tcells. Operating within the endocytic route of antigen-presenting cells (APCs), HLA-DPB1's peptide binding cleft accommodates a range of 10-30 residue peptides. This MHC class II molecule is particularly involved in the exogenous antigen presentation pathway, where peptides from degraded proteins are presented on the cell surface. The complex process involves the association of three MHC class II molecules with a CD74 trimer in the endoplasmic reticulum (ER), forming a heterononamer. Upon entry into the endosomal/lysosomal system, CD74 undergoes sequential degradation, culminating in the release of a small fragment known as CLIP (class-II-associated invariant chain peptide). The removal of CLIP is facilitated by HLA-DM, which stabilizes MHC class II molecules until high-affinity antigenic peptides are bound. The

resultant MHC II molecule-peptide complex is then transported to the cell membrane surface for recognition. Notably, HLA-DO in B-cells and primary dendritic cells (DCs) regulate the interaction between HLA-DM and MHC class II molecules. The lysosomal microenvironment, characterized by increased acidification, plays a crucial role in regulating antigen loading into MHC II molecules, impacting proteolysis and efficient peptide loading. The heterodimeric structure of HLA-DPB1, composed of an alpha and a beta subunit, contributes to its multifaceted role in antigen presentation.

Caution: Product has not been fully validated for medical applications. For research use only.

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Page 2 of 2 www.MedChemExpress.com