

# **Product datasheet for TP720726M**

### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

# MAN1B1 (NM\_016219) Human Recombinant Protein

#### **Product data:**

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human mannosidase, alpha, class 1B, member 1 (MAN1B1)

Species: Human Expression Host: HEK293

**Expression cDNA Clone** 

Asp106-Ala699

or AA Sequence:

Tag: C-His

**Predicted MW:** 68.7 kDa

**Purity:** >95% as determined by SDS-PAGE and Coomassie blue staining

Buffer: Provided lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl

**Endotoxin:** Endotoxin level is < 0.1 ng/µg of protein (< 1 EU/µg)

Storage: Store at -80°C.

**Stability:** Stable for at least 3 months from date of receipt under proper storage and handling

conditions.

**RefSeq:** NP 057303

 Locus ID:
 11253

 UniProt ID:
 Q9UKM7

RefSeq Size: 2787
Cytogenetics: 9q34.3
RefSeq ORF: 2097

**Synonyms:** ERMAN1; ERManl; MANA-ER; MRT15





## MAN1B1 (NM\_016219) Human Recombinant Protein - TP720726M

Summary: This gene encodes an enzyme belonging to the glycosyl hydrolase 47 family. This enzyme

functions in N-glycan biosynthesis, and is a class I alpha-1,2-mannosidase that specifically converts Man9GlcNAc to Man8GlcNAc isomer B. It is required for N-glycan trimming to Man5-6GlcNAc2 in the endoplasmic-reticulum-associated degradation pathway. Mutations in this gene cause autosomal-recessive intellectual disability. Alternative splicing results in multiple transcript variants. A related pseudogene has been identified on chromosome 11. [provided

by RefSeq, Dec 2011]

**Protein Families:** Transmembrane

**Protein Pathways:** Metabolic pathways, N-Glycan biosynthesis