

# Product datasheet for TP720714XL

### 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

# MGAT2 (NM 001015883) Human Recombinant Protein

#### **Product data:**

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human mannosyl (alpha-1,6-)-glycoprotein beta-1,2-N-

acetylglucosaminyltransferase (MGAT2), transcript variant 2

Species: Human Expression Host: HEK293

Expression cDNA Clone

or AA Sequence:

Arg30-Gln447

Tag: C-His

Predicted MW: 49.3 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 001015883

**Locus ID:** 4247

UniProt ID: Q10469

RefSeq Size: 2531

Cytogenetics: 14q21.3

RefSeq ORF: 1341

**Synonyms:** GNT2, CDGS2, GNT-II, GLCNACTII



## MGAT2 (NM\_001015883) Human Recombinant Protein - TP720714XL

**Summary:** The product of this gene is a Golgi enzyme catalyzing an essential step in the conversion of

oligomannose to complex N-glycans. The enzyme has the typical glycosyltransferase

domains: a short N-terminal cytoplasmic domain, a hydrophobic non-cleavable signal-anchor domain, and a C-terminal catalytic domain. Mutations in this gene may lead to carbohydrate-

deficient glycoprotein syndrome, type II. The coding region of this gene is intronless.

Transcript variants with a spliced 5' UTR may exist, but their biological validity has not been

determined. [provided by RefSeq, Jul 2008]

**Protein Families:** Transmembrane

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