

# Recombinant Protein Technical Manual Recombinant Human MGAT2/GlcNAc-TII Protein (His Tag)

### **Product Data:**

**Product SKU:** RPES4280 **Size:** 10μg

Species: Human Cells

**RPES4280** 

Uniprot: Q10469

## **Protein Information:**

Molecular Mass: 49.3 kDa

AP Molecular Mass: 50 kDa

**Tag:** C-6His

**Bio-activity:** 

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

**Endotoxin:**  $< 1.0 \text{ EU per } \mu\text{g}$  as determined by the LAL method.

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

**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 TrisHCl,150mM NaCl, pH8.0 .

**Reconstitution:** Please refer to the printed manual for detailed information.

**Application:** 

**Synonyms:** Alpha;6-Mannosyl-Glycoprotein 2-Beta-N-Acetylglucosaminyltransferase; Beta;2-

N-acetylglucosaminyltransferase II; GlcNAc-T II; NT-II; Mannoside

Acetylglucosaminyltransferase 2; N-Glycosyl-Oligosaccharide-Glycoprotein N-

Acetylglucosaminyltransferase II; MGAT2

# Immunogen Information:

Sequence: Arg30-Gln447

# **Background:**

Mannoside Acetylglucosaminyltransferase 2 (MGAT2) is a single-pass type II membrane protein that contains the typical glycosyltransferase domains: a short N-terminal cytoplasmic domain, a hydrophobic non-cleavable signal-anchor domain and a C-terminal catalytic domain. MGAT2 catalyzes an essential step in the conversion of oligo-mannose to complex N-glycans. Defects in MGAT2 are the cause of congenital disorder of glycosylation type 2A.