

## **GPT** rabbit pAb

Catalog No: YN4210

**Reactivity:** Human; Mouse

**Applications:** WB

Target: GPT

**Fields:** >>N-Glycan biosynthesis;>>Metabolic pathways

Gene Name: DPAGT1 DPAGT2

Q9H3H5

P42867

Protein Name: GPT

Human Gene Id: 1798

**Human Swiss Prot** 

iuman Swiss Fit

No:

Mouse Gene Id: 13478

**Mouse Swiss Prot** 

No:

\_\_\_\_

Immunogen: Synthesized peptide derived from human GPT AA range: 209-259

**Specificity:** This antibody detects endogenous levels of GPT at Human/Mouse

**Formulation:** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

**Dilution:** WB 1 ? 500-2000

**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

1/2



Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight: 45kD

**Background:** 

The protein encoded by this gene is an enzyme that catalyzes the first step in the dolichol-linked oligosaccharide pathway for glycoprotein biosynthesis. This enzyme belongs to the glycosyltransferase family 4. This protein is an integral membrane protein of the endoplasmic reticulum. The congenital disorder of glycosylation type Ij is caused by mutation in the gene encoding this enzyme. [provided by RefSeq, Jul 2008],

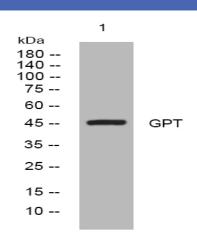
**Function:** 

catalytic activity:UDP-N-acetyl-D-glucosamine + dolichyl phosphate = UMP + N-acetyl-D-glucosaminyl-diphosphodolichol.,disease:Defects in DPAGT1 are the cause of congenital disorder of glycosylation type 1J (CDG1J) [MIM:608093]. CDGs are a family of severe inherited diseases caused by a defect in protein N-glycosylation. They are characterized by under-glycosylated serum proteins. These multisystem disorders present with a wide variety of clinical features, such as disorders of the nervous system development, psychomotor retardation, dysmorphic features, hypotonia, coagulation disorders, and immunodeficiency. The broad spectrum of features reflects the critical role of N-glycoproteins during embryonic development, differentiation, and maintenance of cell functions.,function:Catalyzes the initial step in the synthesis of dolichol-P-P-oligosaccharides.,online information:GlycoGene database,

Subcellular Location:

Endoplasmic reticulum membrane; Multi-pass membrane protein.

## **Products Images**



Western blot analysis of lysates from HuvEc cells, primary antibody was diluted at 1:1000, 4° over night