

Product datasheet for AP51767PU-N

GALNT2 (N-term) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

FC, IHC, WB **Applications:**

Recommended Dilution: ELISA: 1/1000.

Western Blot: 1/100-1/500. **Flow Cytometry:** 1/10-1/50.

Immunohistochemistry on Paraffin Sections: 1/50-1/100.

Reactivity: Human Host: Rabbit

Isotype: lg

Clonality: Polyclonal

KLH conjugated synthetic peptide between 32~62 amino acids from the N-terminal region of Immunogen:

human GALNT2

Specificity: This antibody recognizes Human GALNT2 (N-term).

Formulation: PBS containing 0.09% (W/V) Sodium Azide as preservative

State: Aff - Purified

State: Liquid purified Ig fraction

Concentration: lot specific

Purification: Protein A column, followed by peptide affinity purification

Conjugation: Unconjugated

Storage: Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: polypeptide N-acetylgalactosaminyltransferase 2

Database Link: Entrez Gene 2590 Human

Q10471



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Background: GALNT2 encodes polypeptide N-acetylgalactosaminyltransferase 2, a member of the GalNAc-

transferases family. This family transfers an N-acetyl galactosamine to the hydroxyl group of a serine or threonine residue in the first step of O-linked oligosaccharide biosynthesis.

Individual GalNAc-transferases have distinct activities and initiation of O-glycosylation in a cell

is regulated by a repertoire of GalNAc-transferases.

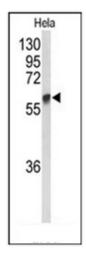
Synonyms: GalNAc transferase 2, pp-GaNTase 2, GalNAc-T2

Note: Molecular Weight: 64733 Da

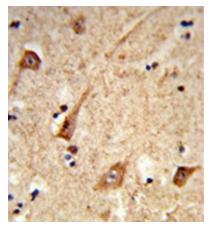
Protein Families: Secreted Protein, Transmembrane

Protein Pathways: Metabolic pathways, O-Glycan biosynthesis

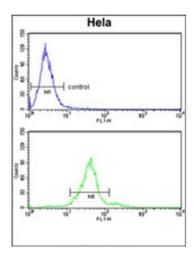
Product images:



Western blot analysis of GALNT2 Antibody (Nterm) in Hela cell line lysates (35ug/lane). GALNT2 (arrow) was detected using the purified Pab.



Immunohistochemistry analysis in formalin-fixed and paraffin-embedded human brain tissue reacted with GALNT2 Antibody (N-term), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining.



Flow cytometry analysis of Hela cells using GALNT2 Antibody (N-term) (bottom histogram) compared to a negative control cell (top histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.