

Product datasheet for TA302556

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OGT Goat Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: WB

Recommended Dilution: ELISA: 1:64,000. WB: 0.05-0.2µg/ml.An

Reactivity: Mouse, Rat (Expected from sequence similarity: Human, Dog)

Host: Goat Isotype: IgG

Clonality: Polyclonal

Immunogen: Peptide with sequence C-YEHPKDLKLSDGR, from the internal region of the protein sequence

according to NP_858058.1; NP_858059.1.

Formulation: Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum

albumin.

Concentration: lot specific

Purification: Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity

chromatography using the immunizing peptide. Supplied at 0.5 mg/ml in Tris saline, 0.02%

sodium azide, pH7.3 with 0.5% bovine serum albumin.

Conjugation: Unconjugated

Storage: Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Predicted Protein Size: 120561 Da

Gene Name: O-linked N-acetylglucosamine (GlcNAc) transferase

Database Link: NP 858058

Entrez Gene 26295 RatEntrez Gene 108155 MouseEntrez Gene 480955 DogEntrez Gene 8473

<u>Human</u> O15294





Background: O-linked N-acetylglucosamine (O-GlcNAc) transferase (OGT) catalyzes the addition of a single

N-acetylglucosamine in O-glycosidic linkage to serine or threonine residues. Since both phosphorylation and glycosylation compete for similar serine or threonine residues, the two processes may compete for sites, or they may alter the substrate specificity of nearby sites by steric or electrostatic effects. The protein contains nine tetratricopeptide repeats and a putative bipartite nuclear localization signal. Two alternatively spliced transcript variants

encoding distinct isoforms have been found for this gene. [provided by RefSeq]

Synonyms: HRNT1; O-GLCNAC **Protein Families:** Druggable Genome

Protein Pathways: Metabolic pathways, O-Glycan biosynthesis

Product images:



TA302556 (0.05ug/ml) staining of Rat Pancreas lysate (35ug protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.