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## Datasheet for ABIN1700952 anti-GCS1 antibody (AA 51-150) (Biotin)



Overview

| Quantity:            | 100 µL   |  |
|----------------------|--|--|
| Target:              | GCS1 (MOGS)  |  |
| Binding Specificity: | AA 51-150  |  |
| Reactivity:          | Human  |  |
| Host:                | Rabbit   |  |
| Clonality:           | Polyclonal   |  |
| Conjugate:           | This GCS1 antibody is conjugated to Biotin   |  |
| Application:         | Western Blotting (WB), ELISA, Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)),<br>Immunohistochemistry (Frozen Sections) (IHC (fro)) |  |

## Product Details

| Immunogen:            | KLH conjugated synthetic peptide derived from human GCS1 |  |
|-----------------------|--|--|
| Isotype:              | IgG  |  |
| Predicted Reactivity: | Human,Mouse,Rat,Dog                                      |  |
| Purification:         | Purified by Protein A.                                   |  |
| Target Details        |  |  |
| Target:               | GCS1 (MOGS)  |  |
| Alternative Name:     | GCS1 (MOGS Products)                                     |  |

Background: Synonyms: EC 3.2.1.106, glucosidase I, Mannosyl oligosaccharide glucosidase, Mannosyl-

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|                     | oligosaccharide glucosidase, Mogs, MOGS_HUMAN, Processing A glucosidase I, Processing A-   |  |  |
|---------------------|--|--|--|
|                     | glucosidase I.   |  |  |
|                     | Background: Glycosylation of asparagine residues in Asn-X-Ser/Thr motifs in proteins   |  |  |
|                     | commonly occur in the lumen of the endoplasmic reticulum (ER). Glucosidase I catalyzes the   |  |  |
|                     | first step in the N-linked oligosaccharide processing pathway. It specifically removes the distal<br>alpha 1,2-linked glucose residue from the Glc3-Man9-GlcNAc2 oligosaccharide precursor.<br>Glucosidase I contains a short cytosolic tail, a single pass transmembrane domain and a large |  |  |
|                     |  |  |  |
|                     |  |  |  |
|                     | C-terminal catalytic domain located on the luminal side of the ER. Mutations in the gene   |  |  |
|                     | encoding Glucosidase I result in the congenital disorder glycosylation (CDG-IIb), which is   |  |  |
|                     | characterized by generalized hypotonia, dysmorphic features, hepatomegaly, hypoventilation,  |  |  |
|                     | feeding problems, seizures and death. Two point mutations in the Glucosidase I gene have   |  |  |
|                     | been identified and result in amino acid substitutions, namely Arg486Thr and Phe652Leu, that   |  |  |
|                     | affect polypeptide folding and active site formation.  |  |  |
| Gene ID:            | 7841   |  |  |
| Pathways:           | SARS-CoV-2 Protein Interactome   |  |  |
| Application Details |  |  |  |
| Application Notes:  | WB 1:300-5000  |  |  |
|                     | IHC-P 1:200-400  |  |  |
|                     | IHC-F 1:100-500  |  |  |
| Restrictions:       | For Research Use only  |  |  |
| Handling            |  |  |  |
| Format:             | Liquid   |  |  |
| Concentration:      | 1 μg/μL  |  |  |
| Buffer:             | Aqueous buffered solution containing 0.01M TBS ( pH 7.4) with 1 % BSA, 0.03 % Proclin300 and   |  |  |
|                     | 50 % Glycerol.   |  |  |
| Preservative:       | ProClin  |  |  |
| Precaution of Use:  | This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be  |  |  |
|                     | handled by trained staff only.   |  |  |
| Storage:            | -20 °C   |  |  |
| Storage Comment:    | Store at -20°C for 12 months.  |  |  |
|                     |  |  |  |

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Expiry Date:

12 months

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