

**AGL Antibody (N-term) Blocking Peptide**

Synthetic peptide  
Catalog # BP2402c

**Specification**

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**AGL Antibody (N-term) Blocking Peptide - Product Information**

Primary Accession [P35573](#)

**AGL Antibody (N-term) Blocking Peptide - Additional Information**

Gene ID 178

**Other Names**

Glycogen debranching enzyme, Glycogen debrancher, 4-alpha-glucanotransferase, Oligo-1, 4-1, 4-glucantransferase, Amylo-alpha-1, 6-glucosidase, Amylo-1, 6-glucosidase, Dextrin 6-alpha-D-glucosidase, AGL, GDE

**Target/Specificity**

The synthetic peptide sequence used to generate the antibody [AP2402c](/product/products/AP2402c) was selected from the N-term region of human AGL. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**AGL Antibody (N-term) Blocking Peptide - Protein Information**

Name AGL

Synonyms GDE

**Function**

Multifunctional enzyme acting as 1,4-alpha-D-glucan:1,4- alpha-D-glucan 4-alpha-D-glycosyltransferase and amylo-1,6-glucosidase in glycogen degradation.

**Cellular Location**

Cytoplasm. Note=Under glycogenolytic conditions localizes to the nucleus

**Tissue Location**

Liver, kidney and lymphoblastoid cells express predominantly isoform 1; whereas muscle and

heart express not only isoform 1, but also muscle-specific isoform mRNAs (isoforms 2, 3 and 4). Isoforms 5 and 6 are present in both liver and muscle

### **AGL Antibody (N-term) Blocking Peptide - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

### **AGL Antibody (N-term) Blocking Peptide - Images**

### **AGL Antibody (N-term) Blocking Peptide - Background**

AGL is a glycogen debrancher enzyme which is involved in glycogen degradation. This enzyme has two independent catalytic activities which occur at different sites on the protein: a 4-alpha-glucotransferase activity and a amylo-1,6-glucosidase activity. Mutations in the AGL gene are associated with glycogen storage disease although a wide range of enzymatic and clinical variability occurs which may be due to tissue-specific alternative splicing.

### **AGL Antibody (N-term) Blocking Peptide - References**

Horinishi, A., et al., J. Hum. Genet. 47(2):55-59 (2002). Shen, J., et al., Hum. Mutat. 9(1):37-40 (1997). Bao, Y., et al., Genomics 38(2):155-165 (1996). Shen, J., et al., J. Clin. Invest. 98(2):352-357 (1996). Yang, B.Z., et al., J. Biol. Chem. 267(13):9294-9299 (1992).