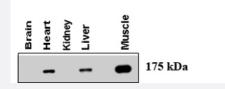


AGL polyclonal antibody

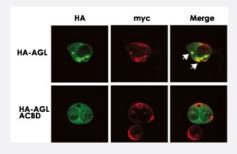
Catalog # PAB4298 Size 400 uL

Applications



Western Blot (Tissue lysate)

Western blot using AGL polyclonal antibody (Cat # PAB4298) at 1 : 500 dilution . A total of 20 ug of lysates was loaded for each tissue . Data courtesy of Dr . Alan Cheng, Department of Internal Medicine, Life Sciences Institute, University of Michigan Medical Center, Ann Arbor, Michigan .



Immunofluorescence

Expression of myc-GS causes wild type but not the CBD mutant of AGL to aggregate around the PAS-stain-positive inclusions. HepG2 cells were transfected with either HA-tagged wild-type AGL (HA-AGL) or HA-AGL CBD. Cells were fixed in formalin and processed for IF using anti-HA (green) and anti-myc (red) antibodies. White arrows indicate colocalization of HA-AGL and myc-GS.

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of AGL.
Immunogen	A synthetic peptide (conjugated with KLH) corresponding to amino acids 1479-1510 at the C-termin us of human AGL.
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Protein A purification



Product Information

Recommend Usage	ELISA Immunofluorescence (1:10-50) Western Blot (1:8000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.09% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

Western Blot (Tissue lysate)

Western blot using AGL polyclonal antibody (Cat # PAB4298) at 1:500 dilution. A total of 20 ug of lysates was loaded for each tissue. Data courtesy of Dr. Alan Cheng, Department of Internal Medicine, Life Sciences Institute, University of Michigan Medical Center, Ann Arbor, Michigan.

Immunofluorescence

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Enzyme-linked Immunoabsorbent Assay



Product Information

Gene	Summary
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This gene encodes the 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 this 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. Alternatively spliced transcripts encoding different isoforms have been described. [provided by RefSeq

Other Designations

OTTHUMP00000012500|OTTHUMP00000012501|OTTHUMP00000012502|OTTHUMP00000012503|OTTHUMP00000012504|OTTHUMP00000012505|glycogen debranching enzyme

Publication Reference

A role for AGL ubiquitination in the glycogen storage disorders of Lafora and Cori's disease.

Cheng A, Zhang M, Gentry MS, Worby CA, Dixon JE, Saltiel AR.

Genes & Development 2007 Oct; 21(19):2399.

Pathway

- Metabolic pathways
- Starch and sucrose metabolism

Disease

- Glycogen Storage Disease Type III
- Tobacco Use Disorder