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anti-AGL antibody (AA 1479-1510)

2 Images



Publication



Go to Product page

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Quantity:	400 μL
Target:	AGL
Binding Specificity:	AA 1479-1510
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This AGL antibody is un-conjugated
Application:	ELISA, Western Blotting (WB), Immunofluorescence (IF)

Product Details

Alternative Name:

Gene ID:

Purpose:	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-	
	terminus of human AGL.	
Cross-Reactivity:	Human	
Target Details		
Target:	AGL	

Pathways: Cellular Glucan Metabolic Process

178

AGL / GDE (AGL Products)

Application Details

Application Notes:	ELISA (1:1000)	
	Western Blot (1:100-500)	
	The optimal working dilution should be determined by the end user.	
Restrictions:	For Research Use only	

Handling

Format:	Liquid	
Buffer:	In PBS (0.09 % sodium azide)	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Storage:	4 °C,-20 °C	
Storage Comment:	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.	

Publications

Product cited in:

Cheng, Zhang, Gentry, Worby, Dixon, Saltiel: "A role for AGL ubiquitination in the glycogen storage disorders of Lafora and Cori's disease." in: **Genes & development**, Vol. 21, Issue 19, pp. 2399-409, (2007) (PubMed).

Images

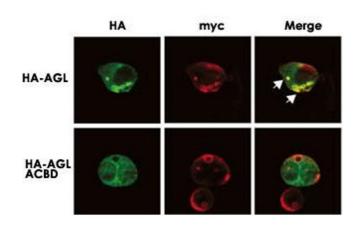
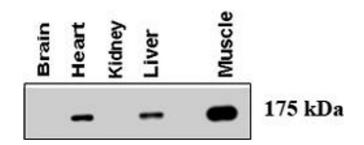


Image 1. Expression of myc-GS causes wild type but not the CBD mutant of AGL to aggregate around the PAS-stain-positive inclusions. HepG2 cells were transfected with either HA-tagged wild-type AGL (HA-AGL) or HA-AGL CBD. Cells were fixed in formalin and processed for IF using anti-HA (green) and anti-myc (red) antibodies. White arrows indicate colocalization of HA-AGL and myc-GS.



Western Blotting

Image 2. Western blot using AGL polyclonal antibody 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.