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Datasheet for ABIN6262045 anti-GPLD1 antibody (Internal Region)

2 Images



Overview

Quantity:	100 µL
Target:	GPLD1
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GPLD1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC)

Product Details

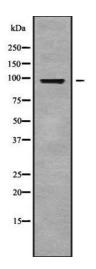
Immunogen:	A synthesized peptide derived from human GPLD1, corresponding to a region within the internal amino acids.
Isotype:	lgG
Specificity:	GPLD1 Antibody detects endogenous levels of total GPLD1.
Predicted Reactivity:	Pig,Bovine,Horse,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	

Target: GPLD1

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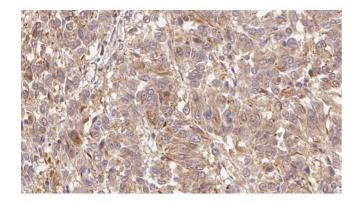
Target Details	
Alternative Name:	GPLD1 (GPLD1 Products)
Background:	Description: This protein hydrolyzes the inositol phosphate linkage in proteins anchored by phosphatidylinositol glycans (GPI-anchor) thus releasing these proteins from the membrane. Gene: GPLD1
Molecular Weight:	92 kDa
Gene ID:	2822
UniProt:	P80108
Pathways:	Inositol Metabolic Process, Regulation of Carbohydrate Metabolic Process
Application Details	
Application Notes:	WB 1:1000-3000, IHC 1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
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:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months

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Western Blotting

Image 1. Western blot analysis GPLD1 using K562 whole cell lysates



Immunohistochemistry

Image 2. ABIN6279662 at 1/100 staining Human Melanoma tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22_jãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary