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Datasheet for ABIN7127651 Recombinant anti-OGT antibody

2 Images



Overview

Quantity:	100 µL
Target:	OGT
Reactivity:	Human
Host:	Rabbit
Antibody Type:	Recombinant Antibody
Clonality:	Monoclonal
Conjugate:	This OGT antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC)

Product Details

Immunogen:	A synthesized peptide derived from human OGT / O-Linked N-Acetylglucosamine Transferase
Clone:	8G7
Isotype:	lgG
Cross-Reactivity:	Human, Mouse, Rat
Purification:	Affinity-chromatography

Target Details

Target:	OGT
Alternative Name:	OGT (OGT Products)
Background:	Background: Catalyzes the transfer of a single N-acetylglucosamine from UDP-GlcNAc to a

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	serine or threonine residue in cytoplasmic and nuclear proteins resulting in their modification with a beta-linked N-acetylglucosamine (O-GlcNAc). Glycosylates a large and diverse number of proteins including histone H2B, AKT1, EZH2, PFKL, KMT2E/MLL5, MAPT/TAU and HCFC1. Can regulate their cellular processes via cross-talk between glycosylation and phosphorylation or by affecting proteolytic processing. Involved in insulin resistance in muscle and adipocyte cells via glycosylating insulin signaling components and inhibiting the 'Thr-308' phosphorylation of AKT1, enhancing IRS1 phosphorylation and attenuating insulin signaling. Involved in glycolysis
	proteins including histone H2B, AKT1, EZH2, PFKL, KMT2E/MLL5, MAPT/TAU and HCFC1. Can regulate their cellular processes via cross-talk between glycosylation and phosphorylation or by affecting proteolytic processing. Involved in insulin resistance in muscle and adipocyte cells via glycosylating insulin signaling components and inhibiting the 'Thr-308' phosphorylation of
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	AKTE ENDANCING IRST DIOSDIOMATION AND ALIENDATION INSUIN SIONATION INVOIVED IN OVEROVSIS
	regulation by mediating glycosylation of 6-phosphofructokinase PFKL, inhibiting its activity
	(PubMed:22923583). Component of a THAP1/THAP3-HCFC1-OGT complex that is required for
	the regulation of the transcriptional activity of RRM1. Plays a key role in chromatin structure by
	mediating O-GlcNAcylation of 'Ser-112' of histone H2B: recruited to CpG-rich transcription start
	sites of active genes via its interaction with TET proteins (TET1, TET2 or TET3)
	(PubMed:22121020, PubMed:23353889). As part of the NSL complex indirectly involved in
	acetylation of nucleosomal histone H4 on several lysine residues (PubMed:20018852). O-
	GlcNAcylation of 'Ser-75' of EZH2 increases its stability, and facilitating the formation of
	H3K27me3 by the PRC2/EED-EZH2 complex (PubMed:24474760). Regulates circadian
	oscillation of the clock genes and glucose homeostasis in the liver. Stabilizes clock proteins
	ARNTL/BMAL1 and CLOCK through O-glycosylation, which prevents their ubiquitination and
	subsequent degradation. Promotes the CLOCK-ARNTL/BMAL1-mediated transcription of genes
	in the negative loop of the circadian clock such as PER1/2 and CRY1/2 (PubMed:12150998,
	PubMed:18288188, PubMed:19377461, PubMed:19451179, PubMed:20018868,
	PubMed:20200153, PubMed:21285374, PubMed:15361863).
	Aliases: UDP-N-acetylglucosaminepeptide N-acetylglucosaminyltransferase 110 kDa subunit
	(EC 2.4.1.255) (O-GlcNAc transferase subunit p110) (O-linked N-acetylglucosamine transferase
	110 kDa subunit) (OGT), OGT
UniProt:	015294
Pathways:	Regulation of Carbohydrate Metabolic Process
Application Details	

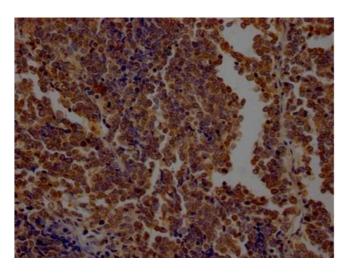
Format:	Liquid
Handling	
Restrictions:	For Research Use only
Application Notes:	Recommended dilution: WB:1:500-1:5000, IHC:1:50-1:200,

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Handling

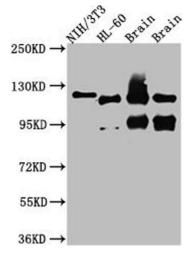
Buffer:	Rabbit IgG in phosphate buffered saline, pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

Images



Immunohistochemistry

Image 1. IHC image of ABIN7127651 diluted at 1:100 and staining in paraffin-embedded human lung cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10 % normal goat serum 30 min at RT. Then primary antibody (1 % BSA) was incubated at 4 °C overnight. The primary is detected by a Goat anti-rabbit IgG polymer labeled by HRP and visualized using 0.05 % DAB.



Western Blotting

Image 2. Western Blot Positive WB detected in: NIH/3T3 whole cell lysate, HL-60 whole cell lysate, Rat Brain whole cell lysate, Mouse Brain whole cell lysate All lanes: OGT antibody at 1:1000 Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 117, 104, 116, 75 kDa Observed band size: 117 kDa

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