

Datasheet for ABIN2784614 anti-ABAT antibody (Middle Region)

1 Image



Overview

O V CI V I C V V	
Quantity:	100 μL
Target:	ABAT
Binding Specificity:	Middle Region
Reactivity:	Human, Mouse, Rat, Cow, Dog, Goat, Guinea Pig, Horse, Rabbit
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This ABAT antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC)
Product Details	
Immunogen:	The immunogen is a synthetic peptide directed towards the middle region of human ABAT
Sequence:	YRSKERGQRG FSQEELETCM INQAPGCPDY SILSFMGAFH GRTMGCLATT
Predicted Reactivity:	Cow: 93%, Dog: 86%, Goat: 77%, Guinea Pig: 93%, Horse: 86%, Human: 100%, Mouse: 93%, Rabbit: 93%, Rat: 93%
Characteristics:	This is a rabbit polyclonal antibody against ABAT. It was validated on Western Blot using a cell lysate as a positive control.
Purification:	Affinity Purified
Target Details	
Target:	ABAT

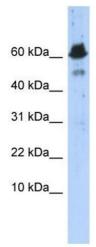
Target Details

Target Details		
Alternative Name:	ABAT (ABAT Products)	
Background:	4-aminobutyrate aminotransferase (ABAT) is responsible for catabolism of gamma-	
	aminobutyric acid (GABA), an important, mostly inhibitory neurotransmitter in the central	
	nervous system, into succinic semialdehyde. The active enzyme is a homodimer of 50-kD	
	subunits complexed to pyridoxal-5-phosphate. ABAT in liver and brain is controlled by 2	
	codominant alleles with a frequency in a Caucasian population of 0.56 and 0.44. The ABAT	
	deficiency phenotype includes psychomotor retardation, hypotonia, hyperreflexia, lethargy,	
	refractory seizures, and EEG abnormalities. 4-aminobutyrate aminotransferase (ABAT) is	
	responsible for catabolism of gamma-aminobutyric acid (GABA), an important, mostly	
	inhibitory neurotransmitter in the central nervous system, into succinic semialdehyde. The	
	active enzyme is a homodimer of 50-kD subunits complexed to pyridoxal-5-phosphate. The	
	protein sequence is over 95 % similar to the pig protein. GABA is estimated to be present in	
	nearly one-third of human synapses. ABAT in liver and brain is controlled by 2 codominant	
	alleles with a frequency in a Caucasian population of 0.56 and 0.44. The ABAT deficiency	
	phenotype includes psychomotor retardation, hypotonia, hyperreflexia, lethargy, refractory	
	seizures, and EEG abnormalities. Multiple alternatively spliced transcript variants encoding th	
	same protein isoform have been found for this gene.	
	Alias Symbols: GABA-AT, GABAT, NPD009	
	Protein Interaction Partner: LGR4, BAG3, ALDH5A1, ABAT,	
	Protein Size: 500	
Molecular Weight:	56 kDa	
Gene ID:	18	
NCBI Accession:	NM_020686, NP_065737	
JniProt:	P80404	
Pathways:	Monocarboxylic Acid Catabolic Process	
Application Details		
Application Notes:	Optimal working dilutions should be determined experimentally by the investigator.	
Comment:	Antigen size: 500 AA	
Restrictions:	For Research Use only	

Handling

Format:	Liquid
Concentration:	Lot specific
Buffer:	Liquid. Purified antibody supplied in 1x PBS buffer with 0.09 % (w/v) sodium azide and 2 % sucrose.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-20 °C
Storage Comment:	For short term use, store at 2-8°C up to 1 week. For long term storage, store at -20°C in small aliquots to prevent freeze-thaw cycles.

Images



Western Blotting

Image 1. WB Suggested Anti-ABAT Antibody Titration:

0.2-1 ug/ml

ELISA Titer: 1:1562500

Positive Control: Transfected 293T