



OAT Monoclonal Antibody

Catalog No	BYmab-00711
Isotype	lgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	OAT
Protein Name	Ornithine aminotransferase mitochondrial
Immunogen	Synthesized peptide derived from OAT . at AA range: 100-180
Specificity	OAT Monoclonal Antibody detects endogenous levels of OAT protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	OAT; Ornithine aminotransferase; mitochondrial; Ornithine delta-aminotransferase; Ornithineoxo-acid aminotransferase
Observed Band	48kD
Cell Pathway	Mitochondrion matrix .
Tissue Specificity	Alzheimer cortex,Brain,Cerebral cortex,Kidney,Liver,Placenta,Subthalamic nucleus,Ut
Function	catalytic activity:L-ornithine + a 2-oxo acid = L-glutamate 5-semialdehyde + an L-amino acid.,cofactor:Pyridoxal phosphate.,disease:Defects in OAT are the cause of hyperornithinemia with gyrate atrophy of choroid and retina (HOGA) [MIM:258870]. HOGA is a slowly progressive blinding autosomal recessive disorder.,pathway:Amino-acid biosynthesis; L-proline biosynthesis; L-glutamate 5-semialdehyde from L-ornithine: step 1/1.,similarity:Belongs to the class-III pyridoxal-phosphate-dependent aminotransferase family.,subunit:Homotetramer.,
Background	ornithine aminotransferase(OAT) Homo sapiens This gene encodes the mitochondrial enzyme ornithine aminotransferase, which is a key enzyme in the pathway that converts arginine and ornithine into the major excitatory and
	Nanjing BYabscience technology Co.,Ltd



国内优质抗体供应商

精准的 WB 检测服务

24H 在线服务,欢迎咨询



Products Images	
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.
matters needing attention	Avoid repeated freezing and thawing!
	inhibitory neurotransmitters glutamate and GABA. Mutations that result in a deficiency of this enzyme cause the autosomal recessive eye disease Gyrate Atrophy. Alternatively spliced transcript variants encoding different isoforms have been described. Related pseudogenes have been defined on the X chromosome. [provided by RefSeq, Jan 2010],

Nanjing BYabscience technology Co.,Ltd