

## HADH rabbit monoclonal antibody

Catalog # H00003033-K Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human HADH peptide using ARM Technology.
Immunogen	A synthetic peptide of human HADH is used for rabbit immunization.  Customer or Abnova will decide on the preferred peptide sequence.
Host	Rabbit
Library Construction	Non-fusion antibody library from rabbit spleen ( <u>ARM Technology</u> ).
Expression	Overexpression vector and transfection into 293H cell line.
Reactivity	Human
Purification	Protein A
Isotype	lgG
Quality Control Testing	Antibody reactive against human HADH peptide by ELISA and mammalian transfected lysate by We stern Blot.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.
Deliverable	Up to three rabbit lgG clones of 100 ug each will be delivered to customer.
Note	<ol> <li>Customer may provide cell or tissue lysate for antibody screening.</li> <li>Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering in cluding F(ab)<sub>2</sub>, lgG, scFv and different Fc and non-Fc conjugates per customer request.</li> </ol>

## **Applications**

Western Blot (Transfected lysate)

Protocol Download



ELISA

Gene Info — HADH	
Entrez GenelD	3033
GeneBank Accession#	<u>HADH</u>
Gene Name	HADH
Gene Alias	HAD, HADH1, HADHSC, HHF4, M/SCHAD, MGC8392, SCHAD
Gene Description	hydroxyacyl-Coenzyme A dehydrogenase
Omim ID	<u>231530</u> <u>601609</u> <u>609975</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene is a member of the 3-hydroxyacyl-CoA dehydrogenase gene family. The encoded prote in functions in the mitochondrial matrix to catalyze the oxidation of straight-chain 3-hydroxyacyl-Co As as part of the beta-oxidation pathway. Its enzymatic activity is highest with medium-chain-lengt h fatty acids. Mutations in this gene cause one form of familial hyperinsulinemic hypoglycemia. The human genome contains a related pseudogene. [provided by RefSeq
Other Designations	L-3-hydroxyacyl-Coenzyme A dehydrogenase L-3-hydroxyacyl-Coenzyme A dehydrogenase, shor t chain

## Pathway

- Butanoate metabolism
- Caprolactam degradation
- Fatty acid elongation in mitochondria
- Fatty acid metabolism
- Geraniol degradation
- Lysine degradation
- Metabolic pathways
- Tryptophan metabolism
- Valine



## Disease

- Alcoholism
- Diabetes Mellitus
- Genetic Predisposition to Disease
- Hyperinsulinism