# ASAH1 rabbit monoclonal antibody

Catalog # H00000427-K

Specification

Size 100 ug x up to 3

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Product Description	Rabbit monoclonal antibody raised against a human ASAH1 peptide using ARM Technology.
Immunogen	A synthetic peptide of human ASAH1 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 (ARM Technology).
Expression	Overexpression vector and transfection into 293H cell line.
Reactivity	Human
Purification	Protein A
lsotype	lgG
Quality Control Testing	Antibody reactive against human ASAH1 peptide by ELISA and mammalian transfected lysate by W estern 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 IgG 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 — ASAH1

Gene Bank Accession# Gene Name Gene Alias Gene Description Omim ID Gene Ontology Gene Summary	<u>427</u>
Gene Name Alias A Gene Alias A Gene Description N Omim ID 2 Gene Ontology H Gene Summary 7	
Gene Alias A Gene Description M Omim ID 2 Gene Ontology E Gene Summary 7 V	ASAH1
Gene Description Omim ID Gene Ontology Gene Summary	ASAH1
Omim ID 2 Gene Ontology E Gene Summary	AC, ASAH, FLJ21558, FLJ22079, PHP, PHP32
Gene Ontology	N-acylsphingosine amidohydrolase (acid ceramidase) 1
Gene Summary	228000
y t r e	Hyperlink
	This gene encodes a heterodimeric protein consisting of a nonglycosylated alpha subunit and a gl ycosylated beta subunit that is cleaved to the mature enzyme posttranslationally. The encoded pro tein catalyzes the synthesis and degradation of ceramide into sphingosine and fatty acid. Mutatio ns in this gene have been associated with a lysosomal storage disorder known as Farber diseas e. Multiple transcript variants encoding several distinct isoforms have been identified for this gene . [provided by RefSeq
Other Designations	N-acylsphingosine amidohydrolase 1 OTTHUMP00000122482 acylsphingosine deacylase putativ

### Pathway

- Lysosome
- <u>Metabolic pathways</u>
- Sphingolipid metabolism

#### Disease

- <u>Cardiovascular Diseases</u>
- Diabetes Mellitus
- Edema



**Product Information** 

• Tobacco Use Disorder