

DEGS2 rabbit monoclonal antibody

Catalog # H00123099-K Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human DEGS2 peptide using ARM Technology.
Immunogen	A synthetic peptide of human DEGS2 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 DEGS2 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 lgG clones of 100 ug each will be delivered to customer.
Note	 Customer may provide cell or tissue lysate for antibody screening. Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering in cluding F(ab)₂, lgG, scFv and different Fc and non-Fc conjugates per customer request.

Applications

Western Blot (Transfected lysate)

Protocol Download



ELISA

Gene Info — DEGS2	
Entrez GenelD	123099
GeneBank Accession#	DEGS2
Gene Name	DEGS2
Gene Alias	C14orf66, DES2, FADS8
Gene Description	degenerative spermatocyte homolog 2, lipid desaturase (Drosophila)
Omim ID	610862
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a bifunctional enzyme that is involved in the biosynthesis of phytosphingolipids in human skin and in other phytosphingolipid-containing tissues. This enzyme can act as a sphing olipid delta(4)-desaturase, and also as a sphingolipid C4-hydroxylase. [provided by RefSeq
Other Designations	degenerative spermatocyte homolog 2, lipid desaturase sphingolipid C4-hydroxylase/delta 4-des aturase sphingolipid delta 4 desaturase/C-4 hydroxylase sphingolipid delta(4)-desaturase/C4-hydroxylase

Pathway

- Metabolic pathways
- Sphingolipid metabolism

Disease

- Genetic Predisposition to Disease
- Narcolepsy