

SPR rabbit monoclonal antibody

Catalog # H00006697-K Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human SPR peptide using ARM Technology.
Immunogen	A synthetic peptide of human SPR 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
Isotype	lgG
Quality Control Testing	Antibody reactive against human SPR peptide by ELISA and mammalian transfected lysate by West ern 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 — SPR	
Entrez GenelD	<u>6697</u>
GeneBank Accession#	<u>SPR</u>
Gene Name	SPR
Gene Alias	SDR38C1
Gene Description	sepiapterin reductase (7,8-dihydrobiopterin:NADP+ oxidoreductase)
Omim ID	<u>182125</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes an aldo-keto reductase that catalyzes the NADPH-dependent reduction of pte ridine derivatives and is important in the biosynthesis of tetrahydrobiopterin (BH4). Mutations in th is gene result in DOPA-responsive dystonia due to sepiaterin reductase deficiency. A pseudogen e has been identified on chromosome 1. [provided by RefSeq
Other Designations	short chain dehydrogenase/reductase family 38C, member 1

Pathway

- Folate biosynthesis
- Metabolic pathways

Disease

- Autistic Disorder
- Bipolar Disorder
- Dystonic Disorders
- Genetic Predisposition to Disease
- Parkinson disease
- Parkinsonian Disorders