

APTX rabbit monoclonal antibody

Catalog # H00054840-K Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human APTX peptide using ARM Technology.
Immunogen	A synthetic peptide of human APTX 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 APTX 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	 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 — APTX	
Entrez GenelD	<u>54840</u>
GeneBank Accession#	APTX
Gene Name	APTX
Gene Alias	AOA, AOA1, AXA1, EAOH, EOAHA, FHA-HIT, FLJ20157, MGC1072
Gene Description	aprataxin
Omim ID	208920 606350 607426
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a member of the histidine triad (HIT) superfamily, some of which have nucleoti de-binding and diadenosine polyphosphate hydrolase activities. The encoded protein may play a role in single-stranded DNA repair. Mutations in this gene have been associated with ataxia-ocula r apraxia. Multiple transcript variants encoding distinct isoforms have been identified for this gene, however, the full length nature of some variants has not been determined. [provided by RefSeq
Other Designations	OTTHUMP00000021188 ataxia 1, early onset with hypoalbuminemia

Disease

- Genetic Predisposition to Disease
- Prostatic Neoplasms