

EYA4 rabbit monoclonal antibody

Catalog # H00002070-K Size 100 ug x up to 3

Specification

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| Product Description | Rabbit monoclonal antibody raised against a human EYA4 peptide using ARM Technology. |
| Immunogen | A synthetic peptide of human EYA4 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 | IgG |
| Quality Control Testing | Antibody reactive against human EYA4 peptide by ELISA and mammalian transfected lysate by Western 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 | 1. Customer may provide cell or tissue lysate for antibody screening. 2. Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering including F(ab) ₂ , IgG, scFv and different Fc and non-Fc conjugates per customer request. |

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- ELISA

Gene Info — EYA4

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| Entrez GeneID | 2070 |
| GeneBank Accession# | EYA4 |
| Gene Name | EYA4 |
| Gene Alias | CMD1J, DFNA10 |
| Gene Description | eyes absent homolog 4 (Drosophila) |
| Omim ID | 601316 603550 605362 |
| Gene Ontology | Hyperlink |
| Gene Summary | This gene encodes a member of the eyes absent (EYA) family of proteins. The encoded protein may act as a transcriptional activator through its protein phosphatase activity, and it may be important for eye development, and for continued function of the mature organ of Corti. Mutations in this gene are associated with postlingual, progressive, autosomal dominant hearing loss at the deafness, autosomal dominant nonsyndromic sensorineural 10 locus. Defects in this gene are also associated with dilated cardiomyopathy 1J. Three transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq] |
| Other Designations | OTTHUMP00000017235 OTTHUMP00000040267 dJ78N10.1 (eyes absent eyes absent 4 |

Disease

- [Tobacco Use Disorder](#)