

HSD17B10 rabbit monoclonal antibody

Catalog # H00003028-K Size 100 ug x up to 3

| Specification | |
|-------------------------|---|
| Product Description | Rabbit monoclonal antibody raised against a human HSD17B10 peptide using ARM Technology. |
| Immunogen | A synthetic peptide of human HSD17B10 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 HSD17B10 peptide by ELISA and mammalian transfected lysate b y 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 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 — HSD17B10 | |
|----------------------|--|
| Entrez GenelD | 3028 |
| GeneBank Accession# | HSD17B10 |
| Gene Name | HSD17B10 |
| Gene Alias | 17b-HSD10, ABAD, CAMR, DUPXp11.22, ERAB, HADH2, HCD2, MHBD, MRPP2, MRX17, MRX31, MRXS10, SCHAD, SDR5C1 |
| Gene Description | hydroxysteroid (17-beta) dehydrogenase 10 |
| Omim ID | 300256 300438 |
| Gene Ontology | <u>Hyperlink</u> |
| Gene Summary | This gene encodes 3-hydroxyacyl-CoA dehydrogenase type II, a member of the short-chain dehyd rogenase/reductase superfamily. The gene product is a mitochondrial protein that catalyzes the o xidation of a wide variety of fatty acids, alcohols, and steroids. The protein has been implicated in the development of Alzheimer's disease, and mutations in the gene are the cause of 2-methyl-3-h ydroxybutyryl-CoA dehydrogenase deficiency (MHBD). Several alternatively spliced transcript variants have been identified, but the full-length nature of only two transcript variants has been determined. [provided by RefSeq |
| Other Designations | 17-beta-hydroxysteroid dehydrogenase type 10 3-hydroxy-2-methylbutyryl-CoA dehydrogenase A B-binding alcohol dehydrogenase OTTHUMP00000023348 OTTHUMP00000023349 amyloid-be ta binding polypeptide amyloid-beta peptide binding alcohol dehydrogenase mental reta |

Pathway

- Metabolic pathways
- Valine