

ARG63484 anti-ACADM antibody

Package: 100 µg
Store at: -20°C

Summary

| | |
|---------------------|---|
| Product Description | Goat Polyclonal antibody recognizes ACADM |
| Tested Reactivity | Hu |
| Tested Application | IHC-P, WB |
| Specificity | This antibody is expected to recognise both reported isoforms. |
| Host | Goat |
| Clonality | Polyclonal |
| Isotype | IgG |
| Target Name | ACADM |
| Species | Human |
| Immunogen | C-RLIVAREHIDKYKN |
| Conjugation | Un-conjugated |
| Alternate Names | ACAD1; MCADH; EC 1.3.8.7; Medium-chain specific acyl-CoA dehydrogenase, mitochondrial; MCAD |

Application Instructions

| Application table | Application | Dilution |
|-------------------|--|------------------|
| | IHC-P | 3 - 5 µg/ml |
| | WB | 0.01 - 0.1 µg/ml |
| Application Note | WB: Recommend incubate at RT for 1h. IHC-P: Antigen Retrieval: Steam tissue section in Citrate buffer (pH 6.0). * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist. | |

Properties

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|---------------------|--|
| Form | Liquid |
| Purification | Purified from goat serum by antigen affinity chromatography. |
| Buffer | Tris saline (pH 7.3), 0.02% Sodium azide and 0.5% BSA. |
| Preservative | 0.02% Sodium azide |
| Stabilizer | 0.5% BSA |
| Concentration | 0.5 mg/ml |
| Storage instruction | For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed before use. |

Bioinformation

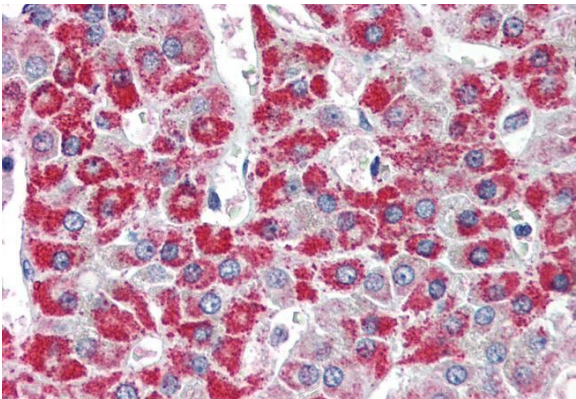
| | |
|----------------|--|
| Database links | GeneID: 34 Human Swiss-port # P11310 Human |
| Background | This gene encodes the medium-chain specific (C4 to C12 straight chain) acyl-Coenzyme A dehydrogenase. The homotetramer enzyme catalyzes the initial step of the mitochondrial fatty acid beta-oxidation pathway. Defects in this gene cause medium-chain acyl-CoA dehydrogenase deficiency, a disease characterized by hepatic dysfunction, fasting hypoglycemia, and encephalopathy, which can result in infantile death. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008] |
| Research Area | Cancer antibody; Cell Biology and Cellular Response antibody; Metabolism antibody; Signaling Transduction antibody |
| Calculated Mw | 47 kDa |
| PTM | Acetylation at Lys-307 and Lys-311 in proximity of the cofactor-binding sites reduces catalytic activity (By similarity). These sites are deacetylated by SIRT3. |

Images



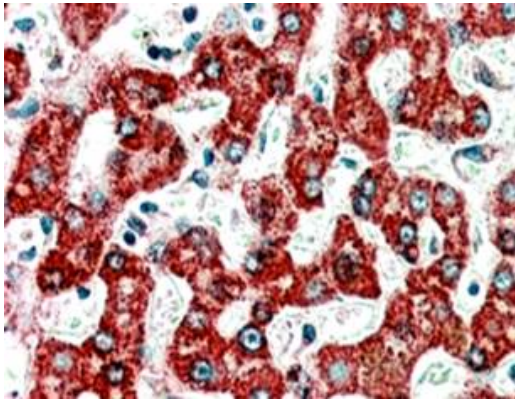
ARG63484 anti-ACADM antibody WB image

Western Blot: Human Heart lysate (RIPA buffer, 35 µg total protein per lane) stained with ARG63484 anti-ACADM antibody at 0.05 µg/ml dilution.



ARG63484 anti-ACADM antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human adrenal gland tissue. Antigen Retrieval: Steam tissue section in Citrate buffer (pH 6.0). The tissue section was stained with ARG63484 anti-ACADM antibody at 3.75 µg/ml dilution followed by AP-staining.



ARG63484 anti-ACADM antibody IHC-P image

Immunohistochemistry: paraffin embedded Human Liver. (Steamed antigen retrieval with citrate buffer pH 6) stained with ARG63484 anti-ACADM antibody at 3.8 µg/ml dilution followed by AP-staining.