

Summary

| Production Name | ACADM Rabbit Monoclonal Antibody | |
|-----------------|--|--|
| Description | Recombinant Rabbit Monoclonal antibody | |
| Host | Rabbit | |
| Application | WB,ICC/IF | |
| Reactivity | Human, Mouse, Rat | |

Performance

| Conjugation | Unconjugated |
|--------------|---|
| Modification | Unmodified |
| lsotype | IgG |
| Clonality | Monoclonal Antibody |
| Form | Liquid |
| Storage | Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw |
| | cycles. |
| Buffer | 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% Sodium azide and 0.05% |
| | BSA |
| Purification | Affinity Purified |

Immunogen

| Gene Name | ACADM |
|-------------------|--|
| Alternative Names | ACADM; Medium-chain specific acyl-CoA dehydrogenase; mitochondrial; MCAD |
| Gene ID | 34 |
| SwissProt ID | P11310 |

Application

| Dilution Ratio | WB: 1/500-1/1000 IF: 1/50-1/200 |
|------------------|--|
| Molecular Weight | Calculated MW: 47 kDa; Observed MW: 47 kDa |

Background

Product Name: ACADM Rabbit Monoclonal Antibody Catalog #: AMRe01599

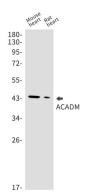


This gene encodes the medium-chain specific (C4 to C12 straight chain) acyl-Coenzyme A dehydrogenase. The homotetramer enzyme catalyzes the initial step of the mitochondrial fatty acid beta-oxidation pathway. Defects in this gene cause medium-chain acyl-CoA dehydrogenase deficiency, a disease characterized 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.

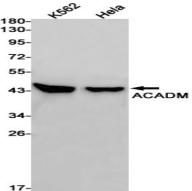
Research Area

Signal Transduction

Image Data

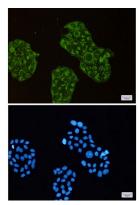


Western blot analysis of ACADM in mouse heart, rat heart lysates using ACADM antibody.



Western blot analysis of ACADM in K562, Hela lysates using ACADM antibody.





Immunocytochemistry analysis of ACADM(green) in Hela using ACADM antibody, and DAPI(blue)

Note For research use only.