

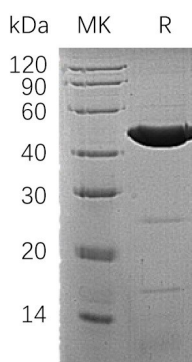
**Product Name: Recombinant Human ACADM (N-6His)**  
**Catalog #: PEH0007**



## Summary

|                                 |   |
|---------------------------------|---|
| <b>Name</b>                     | ACADM/MCAD  |
| <b>Purity</b>                   | Greater than 95% as determined by reducing SDS-PAGE   |
| <b>Endotoxin level</b>          | <1 EU/μg as determined by LAL test.   |
| <b>Construction</b>             | Recombinant Human Medium-Chain Specific Acyl-CoA Dehydrogenase, Mitochondrial is produced by our E.coli expression system and the target gene encoding Lys26-Asn421 is expressed with a 6His tag at the N-terminus. |
| <b>Accession #</b>              | P11310  |
| <b>Host</b>                     | E.coli  |
| <b>Species</b>                  | Human   |
| <b>Predicted Molecular Mass</b> | 45.9 KDa  |
| <b>Formulation</b>              | Supplied as a 0.2 μm filtered solution of 20mM Acetate, 10% Trehalose, 0.05% Tween 80, pH 5.0.  |
| <b>Shipping</b>                 | The product is shipped on dry ice/polar packs. Upon receipt, store it immediately at the temperature listed below.  |
| <b>Stability&amp;Storage</b>    | Store at ≤-70°C, stable for 6 months after receipt. Store at ≤-70°C, stable for 3 months under sterile conditions after opening. Please minimize freeze-thaw cycles.  |
| <b>Reconstitution</b>           |   |

## SDS-PAGE image



## Background

**Alternative Names** Medium-Chain Specific Acyl-CoA Dehydrogenase Mitochondrial; MCAD; ACADM

**Background** Medium-Chain Specific Acyl-CoA Dehydrogenase (ACADM) is a mitochondrial fatty

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acid beta-oxidation that belongs to the acyl-CoA dehydrogenase family. ACADM is a homotetramer enzyme that catalyzes the initial step of the mitochondrial fatty acid beta-oxidation pathway. ACADM is specific for acyl chain lengths of 4 to 16. It is essential for converting these particular fatty acids to energy, especially during fasting periods. Defects in ACADM cause medium-chain acyl-CoA dehydrogenase deficiency, a disease characterized by hepatic dysfunction, fasting hypoglycemia, and encephalopathy, which can result in infantile death.

**Note**

For Research Use Only , Not for Diagnostic Use.