bsm-60484R

[Primary Antibody]

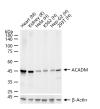
ACADM Recombinant Rabbit mAb



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- DATASHEET		
Host: Rabbit	lsotype: IgG/Kappa	Applications: WB (1:500-2000)
Clonality: Recombinant	CloneNo.: H5A10	IHC-P (1:50-200) IHC-F (1:50-200)
Target: ACADM		IF (1:50-200) Flow-Cyt (1:50-100)
Immunogen: A synthesized peptide derived from human ACADM: 150-200/421.		ICC/IF (1:50-200)
Purification: affinity purified by Protein A		Reactivity: Human, Mouse, Rat
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		
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]		Subcellular Location:

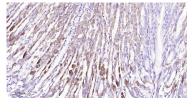
- VALIDATION IMAGES



25 ug total protein per lane of various lysates (see on figure) probed with ACADM monoclonal antibody, unconjugated (bsm-60484R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



Paraformaldehyde-fixed, paraffin embedded Rat Stomach; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with ACADM Monoclonal Antibody, Unconjugated (bsm-60484R) at 1:200 overnight at 4°C, followed by conjugation to the bs-0295G-HRP and DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded Mouse Stomach; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with ACADM Monoclonal Antibody, Unconjugated (bsm-60484R) at 1:200 overnight at 4°C, followed by conjugation to the bs-0295G-HRP and DAB (C-0010) staining.

The Hela (H) cells were fixed with 4% PFA (10 min at r.t.) and then permeabilized with 90% icecold methanol for 20 min at -20°C,the cells then were incubated in 5%BSA to block non-specific protein-protein interactions (30 min at r.t.).Primary Antibody (green):Rabbit Anti-ACADM antibody (bsm-60484R): 1:50-100/10^6 cells; Secondary Antibody (white blue): Goat anti-Rabbit IgG-BF488 (bs-60295G-BF488): 1 µg/test. Blank control (black): PBS. Acquisition of 20,000 events was performed.