

2,4-dienoyl-CoA reductase (DECR1) monoclonal antibody

Cat. no. A21982

Components: 100 µg monoclonal antibody

Lot no.: See product label

Clone/PAD: 8B1AD10

Isotype: Mouse IgG1, Kappa

Gene ID: 1666 **Gene Symbol:** DECR1

Alternative Names: 2,4-dienoyl-CoA reductase, mitochondrial; 2,4-dienoyl-CoA reductase [NADPH];

4-enoyl-CoA reductase [NADPH]; DECR; NADPH; SDR18C1

Concentration: 1 mg/mL in Hepes-Buffered Saline (HBS) with 0.02% sodium azide as a

preservative

mAb PURITY: Near homogeneity as judged by SDS-PAGE. The antibody was produced *in vitro*

using hybridomas grown in serum-free medium, and then purified by

biochemical fractionation.

Reactivity: Human

Immunogen: Human liver mitochondria

Validated Applications: Immunoprecipitation (only from isolated mitochondria), Immunocytochemistry,

Immunohistochemistry, In-Cell ELISA

Suggested Working 1 µg/mL for Immunocytochemistry

Concentration: (This is a starting working concentration. The optimal antibody concentration should be

determined empirically for each specific application.)

Storage: Store at 2–8°C. Do not freeze.

Expiration Date: See product label.

Target Background:

DECR1 encodes an accessory enzyme which participates in the beta-oxidation and metabolism of unsaturated fatty enoyl-CoA esters.

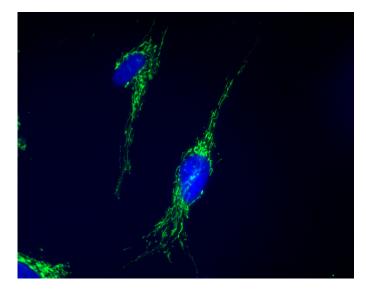
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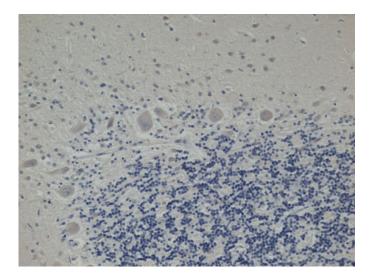
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Immunocytochemistry image of 2,4-dienoyl-CoA reductase (DECR1) monoclonal antibody. Human HDFn cells were fixed in 4% paraformaldehyde for 20 minutes and then permeabilized with 0.1% Triton X-100 for 15 minutes. The cells were incubated with 5 μ g/mL of the antibody overnight at 4°C. GAM-FITC was used as a secondary antibody at a 1/1,000 dilution for 1 hour (green). 10% Goat serum was used as the blocking agent for all blocking steps. The cell nuclei were counterstained with DAPI (blue).



Immunohistochemistry image of 2,4-dienoyl-CoA reductase (DECR1) monoclonal antibody. Immunohistochemical localization of mitochondrial and metabolic enzymes in sections of normal, aged human cerebellar tissue that was formalin-fixed and paraffin-embedded. Immunolabeling was carried out with primary antibodies diluted in TBST/4% BSA at room temperature for one hour. Immunodetection was carried out using a commercial HRP Detection Kit according the manufacturer's instructions.

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