

ACLY monoclonal antibody

Catalog: MB66502

Host: Mouse

Reactivity: Human, Mouse, Monkey

Background:

ATP-citrate lyase (ACL) is a homotetramer that catalyzes the formation of acetyl-CoA and oxaloacetate (OAA) in the cytosol, which is the key step for the biosynthesis of fatty acids, cholesterol and acetylcholine, as well as for gluconeogenesis. Nutrients and hormones regulate the expression level and phosphorylation of ATP-citrate lyase. It is phosphorylated by GSK-3 on Thr446 and Ser450. Ser455 of ATP-citrate lyase has been reported to be phosphorylated by PKA and Akt. Phosphorylation on Ser455 abolishes the homotropic allosteric regulation by citrate and enhances the catalytic activity of the enzyme.

Product:

Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3.

Molecular Weight:

~ 125 kDa

Swiss-Prot:

P53396

Purification&Purity:

The antibody was purified by immunogen affinity chromatography.

Applications:

WB (1/500 - 1/1000), IF/ICC (1/50 - 1/100), FC (1/50 - 1/100)

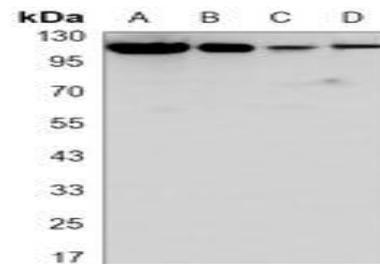
Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

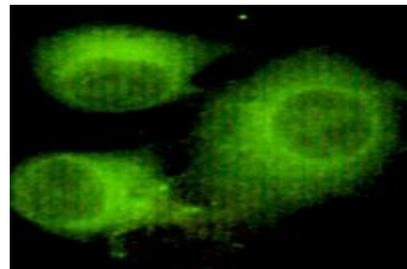
Specificity:

Recognizes endogenous levels of ACLY protein.

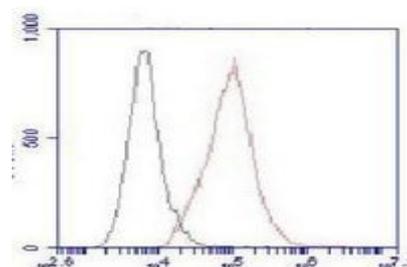
DATA:



Western blot analysis of ACLY expression in NIH3T3 (A), K562 (B), COS7 (C), HeLa (D) whole cell lysates.



Immunofluorescent analysis of ACLY staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.



Note:

For research use only, not for use in diagnostic procedure.

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