ACLY Rabbit mAb

Catalog No.: A3719 Recombinant 2 Publications



Basic Information

Observed MW

125kDa

Calculated MW

121kDa

Category

SMab Recombinant Monoclonal Antibody

Applications

WB,IF/ICC,IP,ELISA

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC0281

Background

ATP citrate lyase is the primary enzyme responsible for the synthesis of cytosolic acetyl-CoA in many tissues. The enzyme is a tetramer (relative molecular weight approximately 440,000) of apparently identical subunits. It catalyzes the formation of acetyl-CoA and oxaloacetate from citrate and CoA with a concomitant hydrolysis of ATP to ADP and phosphate. The product, acetyl-CoA, serves several important biosynthetic pathways, including lipogenesis and cholesterogenesis. In nervous tissue, ATP citrate-lyase may be involved in the biosynthesis of acetylcholine. Multiple transcript variants encoding distinct isoforms have been identified for this gene.

Recommended Dilutions

WB 1:1000 - 1:6000

IF/ICC 1:50 - 1:200

IP 0.5μg-4μg antibody for

200µg-400µg extracts of whole cells

ELISA Recommended starting

concentration is 1
µg/mL. Please optimize
the concentration
based on your specific

assay requirements.

Contact

www.abclonal.com

Immunogen Information

Gene IDSwiss Prot
47
P53396

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 1000-1101 of human ACLY (P53396).

Synonyms

ACL; ATPCL; CLATP; ACLY

Product Information

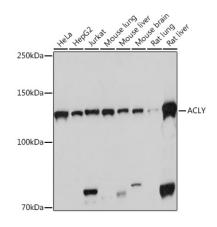
SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.

Validation Data



Western blot analysis of various lysates using ACLY Rabbit mAb (A3719) at 1:1000

dilution.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000

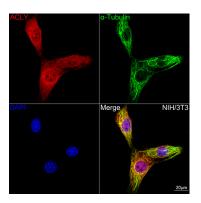
dilution.

Lysates/proteins: 25µg per lane.

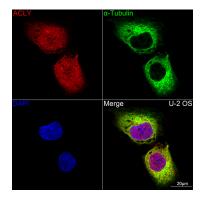
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

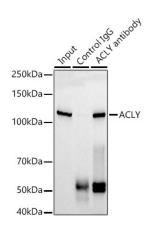
Exposure time: 3s.



Confocal imaging of NIH/3T3 cells using ACLY Rabbit mAb (A3719,dilution 1:100)(Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.



Confocal imaging of U-2 OS cells using ACLY Rabbit mAb (A3719,dilution 1:100)(Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.



Immunoprecipitation analysis of 300 μ g extracts from HepG2 cells using 3 μ g ACLY antibody (A3719). Western blot was performed from the immunoprecipitate using ACLY antibody (A3719) at a dilution of 1:1000.