# SANTA CRUZ BIOTECHNOLOGY, INC.

# p-ACCα (Ser 78/Ser 80)-R: sc-30447-R



# BACKGROUND

Acetyl-CoA carboxylase (ACC) is a complex multifunctional enzyme system which catalyzes the carboxylation of acetyl-CoA to malonyl-CoA, the ratelimiting step in fatty acid synthesis. Exercise diminishes the activity of acetyl-CoA carboxylase in human muscle. ACC $\alpha$  (ACC1) is the rate-limiting enzyme in the biogenesis of long-chain fatty acids, and ACC $\beta$  (ACC2) may control mitochondrial fatty acid oxidation. These two isoforms of ACC control the amount of fatty acids in the cells. The catalytic function of ACC $\alpha$  is regulated by phosphorylation (inactive) and dephosphorylation (active) of targeted serine residues and by allosteric transformation by citrate or palmitoyl-CoA, which serve as the enzyme's short-term regulatory mechanism. The gene encoding ACC $\alpha$  maps to human chromosome 17 and encodes a form of ACC, which is the major ACC in lipogenic tissues. The catalytic core of ACC $\beta$  is homologous to that of the ACC $\alpha$ , except for an additional peptide of about 150 amino acids at the N-terminus.

# REFERENCES

- 1. Kim, K.H. 1997. Regulation of mammalian acetyl-coenzyme A carboxylase. Annu. Rev. Nutr. 17: 77-99.
- 2. Dean, D., et al. 2000. Exercise diminishes the activity of acetyl-CoA carboxylase in human muscle. Diabetes 49: 1295-1300.

#### CHROMOSOMAL LOCATION

Genetic locus: ACACA (human) mapping to 17q12.

#### SOURCE

 $p\text{-}ACC\alpha$  (Ser 78/Ser 80)-R is a rabbit polyclonal antibody raised against a short amino acid sequence containing Ser 78 and Ser 80 phosphorylated ACC $\alpha$  of human origin.

### PRODUCT

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-30447 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

# **APPLICATIONS**

p-ACC $\alpha$  (Ser 78/Ser 80)-R is recommended for detection of Ser 78 and Ser 80 dually phosphorylated ACC $\alpha$  of human and rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for ACC  $\alpha$  siRNA (h): sc-40312, ACC  $\alpha$  shRNA Plasmid (h): sc-40312-SH and ACC  $\alpha$  shRNA (h) Lentiviral Particles: sc-40312-V.

Molecular Weight of p-ACCa: 265 kDa.

Positive Controls: rat heart extract: sc-2393, DU 145 cell lysate: sc-2268 or K-562 whole cell lysate: sc-2203.

#### **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto B Blocking Reagent: sc-2335 (use 50 mM NaF, sc-24988, as diluent), Western Blotting Luminol Reagent: sc-2048 and Lambda Phosphatase: sc-200312A. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

## DATA





Western blot analysis of ACC $\alpha$  phosphorylation in untreated (**A**, **C**), and lambda protein phosphatase (sc-200312A) treated (**B**, **D**) K-562 whole cell lysates. Antibodies tested include p-ACC $\alpha$  (Ser 78/Ser 80)-R: sc-30447-R (**A**, **C**) and ACC $\alpha$  (D-5): sc-137104 (**B**, **D**)

Western blot analysis of ACC $\alpha$  phosphorylation in untreated (**A**) and lambda protein phosphatase (sc-200312A) treated (**B**) rat heart tissue extracts. Antibody tested is p-ACC $\alpha$  (Ser 78/Ser 80): sc-30447 (**A**)

#### SELECT PRODUCT CITATIONS

Chen, L., et al. 2011. Cadmium induction of reactive oxygen species activates the mTOR pathway, leading to neuronal cell death. Free Radic. Biol. Med. 50: 624-632.

# STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

#### **RESEARCH USE**

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

#### PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

