## SANTA CRUZ BIOTECHNOLOGY, INC.

# ACOX1 (H-140): sc-98499



## BACKGROUND

ACOX1 (acyl-coenzyme A oxidase 1), also known as SCOX or PALMCOX, is a 660 amino acid protein that localizes to the peroxisome and belongs to the acyl-CoA oxidase family. Existing as two alternatively spliced isoforms, ACOX1 uses FAD as a cofactor to catalyze the desaturation of very long chain acyl-CoA proteins to 2-*trans*-enoyl-CoA proteins, a reaction that utilizes oxygen and produces hydrogen peroxide. Defects in the gene encoding ACOX1 are the cause of pseudoneonatal adrenoleukodystrophy (pseudo-NALD), which is a single-enzyme disorder that is characterized by seizures, mental retardation, leukody-strophy, mild hepatomegaly and hearing deficits.

#### REFERENCES

- 1. Pacot, C., et al. 1993. Biochemical properties of liver peroxisomes from rat, guinea pig and human species and the influence of hormonal status on rat liver acyl-CoA oxidase mRNA content. Biochimie 75: 235-242.
- Aoyama, T., et al. 1994. Molecular cloning and functional expression of a human peroxisomal acyl-coenzyme A oxidase. Biochem. Biophys. Res. Commun. 198: 1113-1118.

#### CHROMOSOMAL LOCATION

Genetic locus: ACOX1 (human) mapping to 17q25.1; Acox1 (mouse) mapping to 11 E2.

#### SOURCE

ACOX1 (H-140) is a rabbit polyclonal antibody raised against amino acids 436-571 mapping near the C-terminus of ACOX1 of human origin.

#### PRODUCT

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

#### APPLICATIONS

ACOX1 (H-140) is recommended for detection of ACOX1 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

ACOX1 (H-140) is also recommended for detection of ACOX1 in additional species, including equine, canine and bovine.

Suitable for use as control antibody for ACOX1 siRNA (h): sc-94104, ACOX1 siRNA (m): sc-140817, ACOX1 shRNA Plasmid (h): sc-94104-SH, ACOX1 shRNA Plasmid (m): sc-140817-SH, ACOX1 shRNA (h) Lentiviral Particles: sc-94104-V and ACOX1 shRNA (m) Lentiviral Particles: sc-140817-V.

Molecular Weight of ACOX1: 74 kDa.

#### **RESEARCH USE**

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

#### STORAGE

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

### DATA





ACOX1 (H-140): sc-98499. Immunoperoxidase staining of formalin fixed, paraffin-embedded human brain tissue showing cytoplasmic staining of neuronal and nilal cells.

AC0X1 (H-140): sc-98499. Immunoperoxidase staining of formalin fixed, paraffin-embedded human liver tissue showing cytoplasmic staining of hepatocytes and bile duct cells.

#### SELECT PRODUCT CITATIONS

- Liu, I.M., et al. 2011. Regulation of obesity and lipid disorders by extracts from *Angelica acutiloba* root in high-fat diet-induced obese rats. Phytother. Res. 26: 223-230.
- Ghosh, S., et al. 2011. Altered glutathione homeostasis in heart augments cardiac lipotoxicity associated with diet-induced obesity in mice. J. Biol. Chem. 286: 42483-42493.
- 3. Chang, C.J., et al. 2011. Kaempferol regulates the lipid-profile in high-fat diet-fed rats through an increase in hepatic PPAR $\alpha$  levels. Planta Med. 77: 1876-1882.
- 4. Chang, C.J., et al. 2012. Myricetin increases hepatic peroxisome proliferator-activated receptor  $\alpha$  protein expression and decreasesp lasma lipids and adiposity in rats. Evid. Based Complement. Alternat. Med. 2012: 787152.
- Tzeng, T.F., et al. 2012. Vinegar-baked radix bupleuri regulates lipid disorders via a pathway dependent on peroxisome-proliferator-activated receptor-α in high-fat-diet-induced obese rats. Evid. Based Complement. Alternat. Med. 2012: 827278.
- McIntosh, A.L., et al. 2013. Liver fatty acid binding protein gene-ablation exacerbates weight gain in high-fat fed female mice. Lipids 48: 435-448.

#### **PROTOCOLS**

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