

ACOX1 (153CT43.1.1): sc-517306

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, leukodystrophy, mild hepatomegaly and hearing deficits.

REFERENCES

- Pacot, C. and Latruffe, N. 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.
- Varanasi, U., et al. 1994. Isolation of the human peroxisomal acyl-CoA oxidase gene: organization, promoter analysis, and chromosomal localization. *Proc. Natl. Acad. Sci. USA* 91: 3107-3111.
- Fan, C.Y., et al. 1996. Hepatocellular and hepatic peroxisomal alterations in mice with a disrupted peroxisomal fatty acyl-coenzyme A oxidase gene. *J. Biol. Chem.* 271: 24698-24710.
- Fujiwara, C., et al. 2000. Catalase-less peroxisomes. Implication in the milder forms of peroxisome biogenesis disorder. *J. Biol. Chem.* 275: 37271-37277.
- Suzuki, Y., et al. 2002. Peroxisomal acyl CoA oxidase deficiency. *J. Pediatr.* 140: 128-130.
- Online Mendelian Inheritance in Man, OMIM™. 2002. Johns Hopkins University, Baltimore, MD. MIM Number: 609751. World Wide Web URL: <http://www.ncbi.nlm.nih.gov/omim/>
- Ferdinandusse, S., et al. 2007. Clinical, biochemical, and mutational spectrum of peroxisomal acyl-coenzyme A oxidase deficiency. *Hum. Mutat.* 28: 904-912.

CHROMOSOMAL LOCATION

Genetic locus: ACOX1 (human) mapping to 17q25.1.

SOURCE

ACOX1 (153CT43.1.1) is a mouse monoclonal antibody raised against a recombinant protein corresponding to ACOX1 of human origin.

PRODUCT

Each vial contains 100 µg IgG₁ in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

RESEARCH USE

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

APPLICATIONS

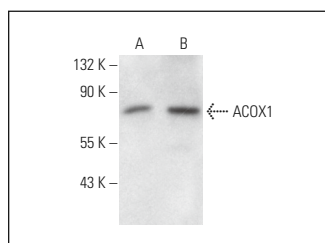
ACOX1 (153CT43.1.1) is recommended for detection of ACOX1 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500).

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

Molecular Weight of ACOX1: 74 kDa.

Positive Controls: K-562 whole cell lysate: sc-2203 or human liver extract: sc-363766.

DATA



ACOX1 (153CT43.1.1): sc-517306. Western blot analysis of ACOX1 expression in K-562 whole cell lysate (A) and human liver tissue extract (B).

SELECT PRODUCT CITATIONS

- Li, G., et al. 2019. MicroRNA-15a regulates the differentiation of intramuscular preadipocytes by targeting ACAA1, ACOX1 and SCP2 in chickens. *Int. J. Mol. Sci.* 20: 4063.
- Tao, S., et al. 2021. Bixin attenuates high-fat diet-caused liver steatosis and inflammatory injury through Nrf2/PPAR α signals. *Oxid. Med. Cell. Longev.* 2021: 6610124.
- Huang, T.Y., et al. 2021. Combined effects of a ketogenic diet and exercise training alter mitochondrial and peroxisomal substrate oxidative capacity in skeletal muscle. *Am. J. Physiol. Endocrinol. Metab.* 320: E1053-E1067.
- Li, L., et al. 2021. Dapagliflozin alleviates hepatic steatosis by restoring autophagy via the AMPK-mTOR pathway. *Front. Pharmacol.* 12: 589273.
- Engel, C., et al. 2022. Altered peroxisome proliferator-activated receptor α signaling in variably diseased peripheral arterial segments. *Front. Cardiovasc. Med.* 9: 834199.
- Xu, Z., et al. 2022. Canagliflozin ameliorates nonalcoholic fatty liver disease by regulating lipid metabolism and inhibiting inflammation through induction of autophagy. *Yonsei Med. J.* 63: 619-631.

STORAGE

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