SANTA CRUZ BIOTECHNOLOGY, INC.

L-FABP (H-120): sc-50380



BACKGROUND

Fatty acid-binding proteins, designated FABPs, are a family of homologous cytoplasmic proteins that are expressed in a highly tissue-specific manner and play an integral role in the balance between lipid and carbohydrate metabolism. FABPs mediate fatty acid (FA) and/or hydrophobic ligand uptake, transport and targeting within their respective tissues. The mechanisms underlying these actions can give rise to both passive diffusional uptake and proteinmediated transmembrane transport of FAs. FABPs are expressed in adipocytes (A-FABP), brain (B-FABP), epithelium (E-FABP, psoriasis-associated FABP,

PA-FABP), striated muscle and heart (H-FABP, mammary-derived growth inhibitor or MDGI), intestine (I-FABP), liver (L-FABP), myelin (M-FABP) and testis (T-FABP). Liver-specific FABP (L-FABP) expression is modulated by developmental, hormonal, dietary and pharmacological factors and is required for cholesterol synthesis and metabolism.

CHROMOSOMAL LOCATION

Genetic locus: FABP1 (human) mapping to 2p11.2; Fabp1 (mouse) mapping to 6 C1.

SOURCE

L-FABP (H-120) is a rabbit polyclonal antibody raised against amino acids 7-126 mapping within an internal region of L-FABP of human origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

L-FABP (H-120) is recommended for detection of L-FABP of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ 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).

L-FABP (H-120) is also recommended for detection of L-FABP in additional species, including canine and porcine.

Suitable for use as control antibody for L-FABP siRNA (h): sc-41243, L-FABP siRNA (m): sc-41244, L-FABP shRNA Plasmid (h): sc-41243-SH, L-FABP shRNA Plasmid (m): sc-41244-SH, L-FABP shRNA (h) Lentiviral Particles: sc-41243-V and L-FABP shRNA (m) Lentiviral Particles: sc-41244-V.

Molecular Weight of L-FABP: 14 kDa.

Positive Controls: L-FABP (m): 293T Lysate: sc-121261, Hep G2 cell lysate: sc-2227 or mouse liver extract: sc-2256.

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.

DATA



L-FABP (H-120): sc-50380. Western blot analysis of L-FABP expression in non-transfected 293T: sc-117752 (**A**), mouse L-FABP transfected 293T: sc-121261 (**B**) and Hep G2 (**C**) whole cell lysates.



L-FABP (H-120): sc-50380. Immunoperoxidase staining of formalin fixed, paraffin-embedded human kidney tissue showing cytoplasmic and nuclear staining of cells in tubules (**A**). Immunoperoxidase staining of formalin fixed, paraffin-embedded human rectum tissue showing cytoplasmic and nuclear staining of glandular cells (**B**).

SELECT PRODUCT CITATIONS

- Labonté, E.D., et al. 2008. Reduced absorption of saturated fatty acids and resistance to diet-induced obesity and diabetes by ezetimibe-treated and Npc111-/- mice. Am. J. Physiol. Gastrointest. Liver Physiol. 295: G776-G783.
- Chaerkady, R., et al. 2008. A quantitative proteomic approach for identification of potential biomarkers in hepatocellular carcinoma. J. Proteome Res. 7: 4289-4298.
- Liao, C.C., et al. 2014. The inhibition of oleic acid induced hepatic lipogenesis and the promotion of lipolysis by caffeic acid via up-regulation of AMP-activated kinase. J. Sci. Food Agric. 94: 1154-1162.

PROTOCOLS

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Try L-FABP (F-9): sc-271591 or L-FABP (C-4):

sc-374537, our highly recommended monoclonal alternatives to L-FABP (H-120).