

### **Product Data Sheet**

# **Anti-ACSVL1 Antibody**

Catalog # Source Reactivity Applications

CQA2547 Rabbit H, M, R WB, IH

**Description** Rabbit polyclonal antibody to ACSVL1

Immunogen Recombinant full length protein of human ACSVL1

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of ACSVL1 protein.

**Clonality** Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

**Dilution** WB (1/500 - 1/2000), IH (1/50 - 1/200)

Gene Symbol SLC27A2

ACSVL1; FACVL1; FATP2; VLACS; Very long-chain acyl-CoA synthetase; VLACS; VLCS;

Fatty acid transport protein 2; FATP-2; Fatty-acid-coenzyme A ligase, very long-chain

1; Long-chain-fatty-acid--CoA ligase; Solute carrier family 27 member 2; THCA-CoA

ligase; Very long-chain-fatty-acid-CoA ligase

Entrez Gene 11001 (Human); 26458 (Mouse)

**SwissProt** O14975 (Human); O35488 (Mouse)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

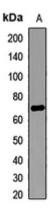
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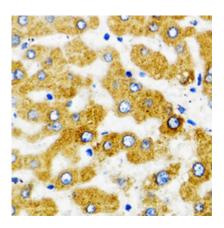




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Western blot analysis of ACSVL1 expression in HepG2 (A) whole cell lysates. (Predicted band size: 64; 70 kD; Observed band size: 70 kD)



Immunohistochemical analysis of ACSVL1 staining in human liver formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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