

A Reliable Research Partner in Life Science and Medicine

## **ACSL4 Polyclonal Antibody**

Catalog No. E-AB-14661

*Note:* Centrifuge before opening to ensure complete recovery of vial contents.

### **Description**

**Reactivity** Human, Mouse, Rat

**Immunogen** Recombinant protein of human ACSL4

Host Rabbit Isotype IgG

**Purification** Affinity purification

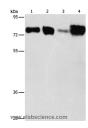
**Conjugation** Unconjugated

**Buffer** PBS with 0.05% sodium azide and 50% glycerol, PH7.4

## **Applications** Recommended Dilution

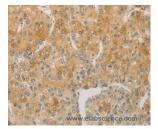
WB 1:1000-1:5000 IHC 1:50-1:200

### Data

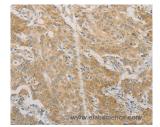


Western Blot analysis of Hepg2 and hela cell, Human fetal kidney and liver tissue using ACSL4 Polyclonal Antibody at dilution of 1:650

Calculated Mw:79kDa



Immunohistochemistry of paraffin-embedded Human liver cancer using ACSL4 Polyclonal Antibody at dilution of 1:60



Immunohistochemistry of paraffin-embedded Human gastric cancer using ACSL4 Polyclonal Antibody at dilution of 1:60

# **Preparation & Storage**

Storage Store at -20°C. Avoid freeze / thaw cycles.

#### For Research Use Only

Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017

Web: <u>www.elabscience.com</u> Email: <u>techsupport@elabscience.com</u>





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## **Background**

The protein encoded by this gene is an isozyme of the long-chain fatty-acid-coenzyme A ligase family. Although differing in substrate specificity, subcellular localization, and tissue distribution, all isozymes of this family convert free long-chain fatty acids into fatty acyl-CoA esters, and thereby play a key role in lipid biosynthesis and fatty acid degradation. This isozyme preferentially utilizes arachidonate as substrate. The absence of this enzyme may contribute to the mental retardation or Alport syndrome. Alternative splicing of this gene generates 2 transcript variants.

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