SCCPDH Polyclonal Antibody

Catalog Number: E-AB-19034



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

Description

Reactivity Human, Mouse, Rat

Immunogen Fusion protein of human SCCPDH

Host Rabbit Isotype IgG

Purification Antigen affinity purification

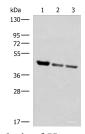
Conjugation Unconjugated

Formulation PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

Applications Recommended Dilution

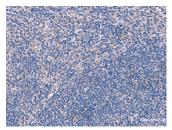
WB 1:1000-1:5000 IHC 1:50-1:300 ELISA 1:5000-1:10000

Data

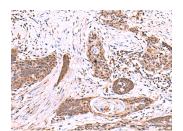


Western blot analysis of Human fetal liver tissue PC-3 and A172 cell lysates using SCCPDH Polyclonal Antibody at dilution of 1:1000

Observed Mw:Refer to figures Calculated Mw:47 kDa



Immunohistochemistry of paraffin-embedded Human tonsil tissue using SCCPDH Polyclonal Antibody at dilution of 1:85(×200)



Immunohistochemistry of paraffin-embedded Human esophagus cancer tissue using SCCPDH Polyclonal Antibody at dilution of 1:85(×200)

Preparation & Storage

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

Background

For Research Use Only

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SCCPDH (Probable saccharopine dehydrogenase) is a 429 amino acid protein that belongs to the saccharopine dehydrogenase family. The SCCPDH gene is conserved in chimpanzee, dog, cow, mouse, rat, chicken, fruit fly, mosquito and C.elegans, and maps to human chromosome 1q44. Chromosome 1 is the largest human chromosome spanning about 260 million base pairs and making up 8% of the human genome. There are about 3,000 genes on chromosome 1, and considering the great number of genes there are also a large number of diseases associated with chromosome 1. Stickler syndrome, Parkinsons, Gaucher disease and Usher syndrome are also associated with chromosome 1. A breakpoint has been identified in 1q which disrupts the DISC1 gene and is linked to schizophrenia. Aberrations in chromosome 1 are found in a variety of cancers including head and neck cancer, malignant melanoma and multiple myeloma.

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