产品名称: TSSC3 Rabbit Polyclonal Antibody

产品货号: APRab19374



产品概述 (Summary)

产品名称 (Production Name) TSSC3 Rabbit Polyclonal Antibody

描述 (Description) Rabbit polyclonal Antibody

宿主 (Host) Rabbit

应用 (Application) IHC,ICC/IF,ELISA 种属反应性 (Reactivity) Human,Rat,Mouse

产品性能 (Performance)

偶联物 (Conjugation)Unconjugated修饰 (Modification)Unmodified

同种型 (Isotype) IgG

克隆 (Clonality) Polyclonal 形式 (Form) Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid 存放说明 (Storage)

freeze/thaw cycles.

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% 储存溶液 (Buffer)

New type preservative N.

纯化方式 (Purification) Affinity purification

免疫原信息 (Immunogen)

基因名 (Gene Name) PHLDA2

PHLDA2; BWR1C; HLDA2; IPL; TSSC3; Pleckstrin homology-like domain family

A member 2; Beckwith-Wiedemann syndrome chromosomal region 1 別名 (Alternative Names)

candidate gene C protein; Imprinted in placenta and liver protein; Tumor-

suppressing STF cDNA 3 protein; T

基因 ID (Gene ID) 7262.0

Q53GA4.The antiserum was produced against synthesized peptide derived 蛋白ID (SwissProt ID)

from human PHLDA2. AA range:81-130

产品应用 (Application)

稀释比 (Dilution Ratio) IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:10000-1:20000

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蛋白分子量 (Molecular Weight)

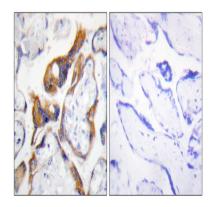
研究背景 (Background)

This gene is located in a cluster of imprinted genes on chromosome 11p15.5, which is considered to be an important tumor suppressor gene region. Alterations in this region may be associated with the Beckwith-Wiedemann syndrome, Wilms tumor, rhabdomyosarcoma, adrenocortical carcinoma, and lung, ovarian, and breast cancer. This gene has been shown to be imprinted, with preferential expression from the maternal allele in placenta and liver. [provided by RefSeq, Oct 2010], domain: The PH domain binds phosphoinositides with a broad specificity. It may compete with the PH domain of some other proteins, thereby interfering with their binding to phosphatidylinositol 4,5-bisphosphate (PIP2) and phosphatidylinositol 3,4,5-triphosphate (PIP3), function: Plays a role in regulating placenta growth. May act via its PH domain that competes with other PH domain-containing proteins, thereby preventing their binding to membrane lipids.,induction:Maternal PHLDA2 allele is activated, while paternal Phlda2 is repressed due to genomic imprinting. Downregulated by hypoxia. Although highly similar to PHLDA3 protein, it is not regulated by p53/TP53, miscellaneous: The PHLDA2 locus is imprinted. Loss of imprinting results in overexpression. Imprinting is dependent on RNAi machinery, similarity: Belongs to the PHLDA2 family, similarity: Contains 1 PH domain, tissue specificity: Expressed in placenta and adult prostate gland. In placenta, it is present in all cells of the villous cytotrophoblast. The protein is absent in cells from hydatidiform moles. Hydatidiform mole is a gestation characterized by abnormal development of both fetus and trophoblast. The majority of hydatidiform moles are associated with an excess of paternal to maternal genomes and are likely to result from the abnormal expression of imprinted genes (at protein level). Expressed at low levels in adult liver and lung, and fetal liver. Expressed in adult brain and neuroblastoma, medullablastoma and glioblastoma cell lines.,

研究领域 (Research Area)

Cell Biology; Cell Cycle; Cell Cycle Inhibitors; Epigenetics and Nuclear Signaling; Transcription; Cancer susceptibility; Tumor Suppressors

图片 (Image Data)



Immunohistochemistry analysis of paraffin-embedded human placenta, using PHLA2 Antibody. The picture on the right is blocked with the synthesized peptide.

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注意事项 (Note)

For research use only.

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