abcam

Product datasheet

Anti-SUR1 antibody ab217633

1 References 2 图像

概述

产品名称 Anti-SUR1抗体

描述 兔多克隆抗体to SUR1

宿主 Rabbit

适用于: IHC-P, WB 经测试应用 种属反应性 **与反应:** Mouse, Rat

预测可用于: Human 📤

免疫原 Synthetic peptide within Human SUR1 aa 332-355 conjugated to keyhole limpet haemocyanin.

The exact sequence is proprietary.

Database link: Q09428

阳性对照 IHC-P: Mouse intestine tissue.

常规说明 The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

> Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

存储溶液 pH: 7.40

Preservative: 0.02% Proclin 300

Constituents: 50% Glycerol (glycerin, glycerine), 1% BSA, 48.98% TBS, 1X

纯度 Protein A purified

克隆 多克隆 同种型 lgG

The Abpromise guarantee

Abpromise™承诺保证使用ab217633于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
IHC-P		1/100 - 1/500.
WB		1/1000.

靶标

功能

疾病相关

Putative subunit of the beta-cell ATP-sensitive potassium channel (KATP). Regulator of ATP-sensitive K(+) channels and insulin release.

Defects in ABCC8 are a cause of leucine-induced hypoglycemia (LIH) [MIM:240800]; also known as leucine-sensitive hypoglycemia of infancy. LIH is a rare cause of hypoglycemia and is described as a condition in which symptomatic hypoglycemia is provoked by high protein feedings. Hypoglycemia is also elicited by administration of oral or intravenous infusions of a single amino acid, leucine.

Defects in ABCC8 are the cause of familial hyperinsulinemic hypoglycemia type 1 (HHF1) [MIM:256450]; also known as persistent hyperinsulinemic hypoglycemia of infancy (PHHI) or congenital hyperinsulinism. HHF is the most common cause of persistent hypoglycemia in infancy and is due to defective negative feedback regulation of insulin secretion by low glucose levels. It causes nesidioblastosis, a diffuse abnormality of the pancreas in which there is extensive, often disorganized formation of new islets. Unless early and aggressive intervention is undertaken, brain damage from recurrent episodes of hypoglycemia may occur.

Defects in ABCC8 are a cause of diabetes mellitus permanent neonatal (PNDM) [MIM:606176]. PNDM is a rare form of diabetes distinct from childhood-onset autoimmune diabetes mellitus type 1. It is characterized by insulin-requiring hyperglycemia that is diagnosed within the first months of life. Permanent neonatal diabetes requires lifelong therapy.

Defects in ABCC8 are the cause of transient neonatal diabetes mellitus type 2 (TNDM2) [MIM:610374]. Neonatal diabetes is a form of diabetes mellitus defined by the onset of mild-to-severe hyperglycemia within the first months of life. Transient neonatal diabetes remits early, with a possible relapse during adolescence.

序列相似性

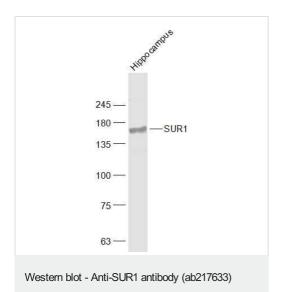
Belongs to the ABC transporter superfamily. ABCC family. Conjugate transporter (TC 3.A.1.208) subfamily.

Contains 2 ABC transmembrane type-1 domains.

Contains 2 ABC transporter domains.

细胞定位 Membrane.

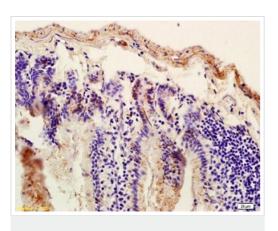
图片



Anti-SUR1 antibody (ab217633) at 1/1000 dilution + Rat hippocampus lysates

Secondary

Conjugated secondary antibody at 1/20000 dilution



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SUR1 antibody (ab217633)

Immunohistochemical analysis of formalin-fixed and paraffinembedded mouse intestine tissue labeling SUR1 with ab217633 at 1/200 dilution, followed by conjugation to the secondary antibody and DAB staining.

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