abcam

Product datasheet

Anti-EDN3 antibody ab197374

2 图像

概述

产品名称 Anti-EDN3抗体

描述 兔多克隆抗体to EDN3

宿主 Rabbit

经测试应用 适用于: IHC-P

种属反应性 与反应: Human

免疫原 Fusion protein within Human EDN3 aa 1 to the C-terminus (internal sequence). The exact

> immunogen sequence used to generate this antibody is proprietary information. If additional detail on the immunogen is needed to determine the suitability of the antibody for your needs, please contact our Scientific Support team to discuss your requirements. The identity of the protein

fusion partner is GST. Database link: P14138

阳性对照 Human esophagus and thyroid cancer 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

Preservative: 0.05% Sodium azide

Constituents: 50% Glycerol (glycerin, glycerine), 49% PBS

纯度 Immunogen affinity purified

ΙgG

克隆 多克隆 同种型

The Abpromise guarantee

Abpromise™承诺保证使用ab197374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		1/25 - 1/100.

靶标

功能

Endothelins are endothelium-derived vasoconstrictor peptides.

组织特异性

Expressed in trophoblasts and placental stem villi vessels, but not in cultured placental smooth

muscle cells.

疾病相关

Defects in EDN3 are the cause of Hirschsprung disease type 4 (HSCR4) [MIM:613712]; also known as aganglionic megacolon (MGC). A genetic disorder of neural crest development characterized by the absence of intramural ganglion cells in the hindgut; often resulting in intestinal

obstruction.

Defects in EDN3 are a cause of congenital central hypoventilation syndrome (CCHS)

[MIM:209880]; also known as congenital failure of autonomic control or Ondine curse. CCHS is a rare disorder characterized by abnormal control of respiration in the absence of neuromuscular or lung disease, or an identifiable brain stem lesion. A deficiency in autonomic control of respiration results in inadequate or negligible ventilatory and arousal responses to hypercapnia and

hypoxemia.

Defects in EDN3 are a cause of Waardenburg syndrome type 4 (WS4B) [MIM:613265]; also known as Waardenburg-Shah syndrome. WS4B is characterized by the association of

Waardenburg features (depigmentation and deafness) and the absence of enteric ganglia in the

distal part of the intestine (Hirschsprung disease).

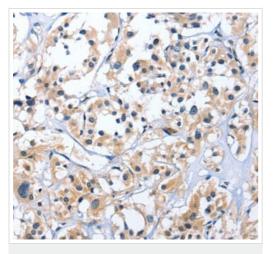
序列相似性

Belongs to the endothelin/sarafotoxin family.

细胞定位

Secreted.

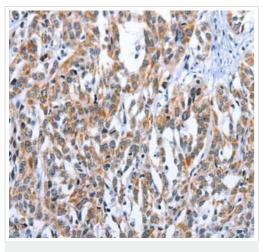
图片



cancer tissue, labeling EDN3 with ab197374 at 1/30 dilution.

Immunohistochemical analysis of paraffin-embedded Human thyroid

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-EDN3 antibody (ab197374)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-EDN3 antibody (ab197374)

Immunohistochemical analysis of paraffin-embedded Human esophagus cancer tissue, labeling EDN3 with ab197374 at 1/30 dilution.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- · We investigate all quality concerns to ensure our products perform to the highest standards

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