abcam

Product datasheet

Anti-Wnt10a antibody ab62051

4 References 画像数 4

製品の概要

製品名	Anti-Wnt10a antibody
製品の詳細	Rabbit polyclonal to Wnt10a
由来種	Rabbit
アプリケーション	**適用あり:** IHC-P
種交差性	**交差種:** Human
免疫原	Synthetic peptide corresponding to Wnt10a (internal sequence) conjugated to Keyhole Limpet Haemocyanin (KLH).
特記事項	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

製品の特性

製品の状態	Liquid
保存方法	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.
バッファー	pH: 7.70 Preservative: 0.1% Sodium azide Constituent: PBS
精製度	Immunogen affinity purified
ポリ/モノ	ポリクローナル
アイソタイプ	IgG

アプリケーション

The Abpromise guarantee	**Abpromise保証は、** 次のテスト済みアプリケーションにおけるab62051の使用に適用されます

アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IHC-P		Use a concentration of 10 µg/ml.

ターゲット情報

機能

Ligand for members of the frizzled family of seven transmembrane receptors. Probable developmental protein. May be a signaling molecule important in CNS development. Is likely to signal over only few cell diameters.

関連疾患

Defects in WNT10A are a cause of ectodermal dysplasia anhidrotic (EDA) [MIM:224900]; also known ectodermal dysplasia hypohidrotic autosomal recessive (HED). Ectodermal dysplasia defines a heterogeneous group of disorders due to abnormal development of two or more ectodermal structures. EDA is characterized by sparse hair (atrichosis or hypotrichosis), abnormal or missing teeth and the inability to sweat due to the absence of sweat glands.
Note=Most patients carrying WNT10A mutations present with sweating anomalies. However, comparison with EDA cases harboring mutations in the ectodysplasin pathway identifies some phenotypic differences. Dermatological features (anomalies of hair and sweat glands) are less severe in patients carrying WNT10A mutations and facial dysmorphism can be absent. The dental phenotype consists in microdontia, whereas teeth agenesis is more frequent in patients carrying mutations in the ectodysplasin pathway.
Defects in WNT10A are a cause of odonto-onycho-dermal dysplasia (OODD) [MIM:257980]. OODD is a rare autosomal recessive ectodermal dysplasia in which the presenting phenotype is dry hair, severe hypodontia, smooth tongue with marked reduction of fungiform and filiform papillae, onychodysplasia, keratoderma and hyperhidrosis of palms and soles, and hyperkeratosis of the skin.
Defects in WNT10A are a cause of Schopf-Schulz-Passarge syndrome (SSPS) [MIM:224750]. SSPS is rare ectodermal dysplasia, characterized chiefly by cysts of the eyelid margins, palmoplantar keratoderma, hypodontia, hypotrichosis and nail dystrophy. Multiple eyelid apocrine hidrocystomas are the hallmark of this condition, although they usually appear in adulthood. The concomitant presence of eccrine syringofibroadenoma in most patients and of other adnexal skin tumors in some affected subjects indicates that Schopf-Schulz-Passarge syndrome is a genodermatosis with skin appendage neoplasms.

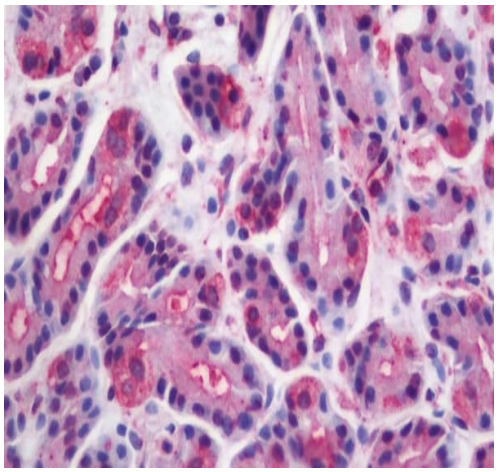
配列類似性

Belongs to the Wnt family.

細胞内局在

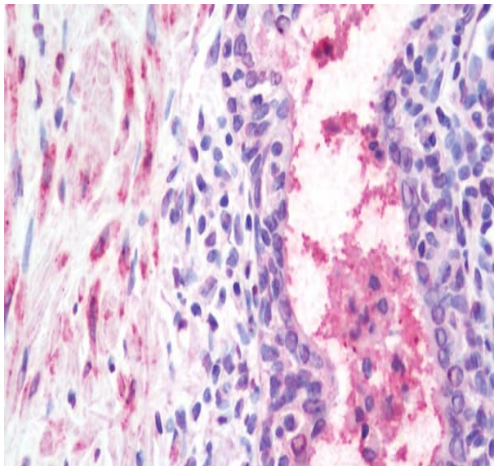
Secreted > extracellular space > extracellular matrix.

画像



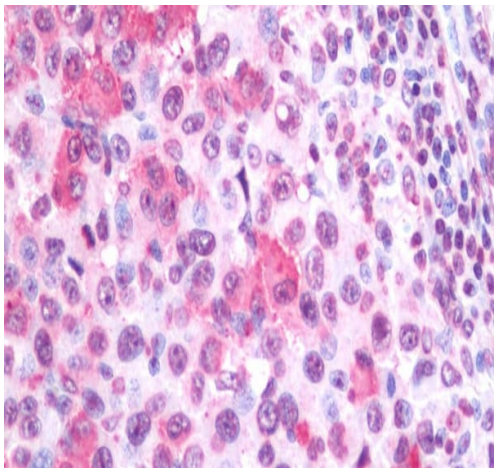
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Wnt10a antibody (ab62051)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human stomach tissue sections labelling Wnt10a with ab62051.



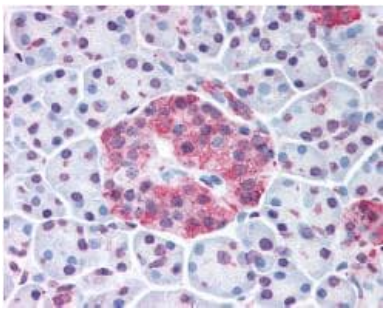
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Wnt10a antibody (ab62051)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human prostate tissue sections labelling Wnt10a with ab62051.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Wnt10a antibody (ab62051)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human melanoma (lymph node) tissue sections labelling Wnt10a with ab62051.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Wnt10a antibody (ab62051)

ab62051, at 10µg/ml, staining Wnt10a in Formalin Fixed Paraffin Embedded Human Pancreatic Islets tissue by Immunohistochemistry.

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