abcam

Product datasheet

Anti-GDF1 antibody [EPR5815] ab124706

ועלשעבע RabMAb

1 References 画像数3

製品の概要

製品名 Anti-GDF1 antibody [EPR5815]

製品の詳細 Rabbit monoclonal [EPR5815] to GDF1

由来種 Rabbit

アプリケーション **適用あり:** WB

適用なし: Flow Cyt,ICC/IF or IHC-P

種交差性 交差種: Human

交差が予測される動物種: Mouse, Rat 🔷

Synthetic peptide. This information is proprietary to Abcam and/or its suppliers. 免疫原

ポジティブ・コントロール U87-MG and Human fetal brain lysates.

特記事項 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

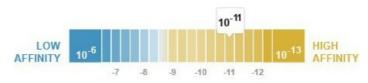
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

 $K_D = 8.40 \times 10^{-11} M$ 解離定数(KD値)



Learn more about K_D

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

精製度 Protein A purified

ポリモノ モノクローナル **クローン名** EPR5815

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab124706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 |
|----------|-----------|-------------------------------------------------------|
| WB | | 1/1000 - 1/10000. Predicted molecular weight: 39 kDa. |

追加情報 Is unsuitable for Flow Cyt,ICC/IF or IHC-P.

ターゲット情報

機能

May mediate cell differentiation events during embryonic development.

組織特異性

Expressed in the brain.

関連疾患

Defects in GDF1 are a cause of conotruncal heart malformations (CTHM) [MIM:217095]. A group of congenital heart defects involving the outflow tracts. Examples include truncus arteriosus communis, double-outlet right ventricle and transposition of great arteries. Truncus arteriosus communis is characterized by a single outflow tract instead of a separate aorta and pulmonary artery. In transposition of the great arteries, the aorta arises from the right ventricle and the pulmonary artery from the left ventricle. In double outlet of the right ventricle, both the pulmonary artery and aorta arise from the right ventricle.

Defects in GDF1 are the cause of transposition of the great arteries dextro-looped type 3 (DTGA3) [MIM:613854]. A congenital heart defect consisting of complete inversion of the great vessels, so that the aorta incorrectly arises from the right ventricle and the pulmonary artery incorrectly arises from the left ventricle. This creates completely separate pulmonary and systemic circulatory systems, an arrangement that is incompatible with life. The presence or absence of associated cardiac anomalies defines the clinical presentation and surgical management of patients with transposition of the great arteries.

Defects in GDF1 are a cause of tetralogy of Fallot (TOF) [MIM:187500]. A congenital heart anomaly which consists of pulmonary stenosis, ventricular septal defect, dextroposition of the aorta (aorta is on the right side instead of the left) and hypertrophy of the right ventricle. In this condition, blood from both ventricles (oxygen-rich and oxygen-poor) is pumped into the body often causing cyanosis.

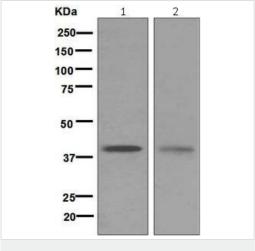
配列類似性

Belongs to the TGF-beta family.

細胞内局在

Secreted.

画像



Western blot - Anti-GDF1 antibody [EPR5815] (ab124706)

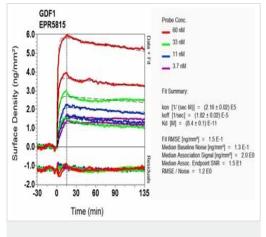
All lanes: Anti-GDF1 antibody [EPR5815] (ab124706) at 1/1000 dilution

Lane 1: U87-MG cell lysate

Lane 2: Human fetal brain lysates cell lysate

Lysates/proteins at 10 µg per lane.

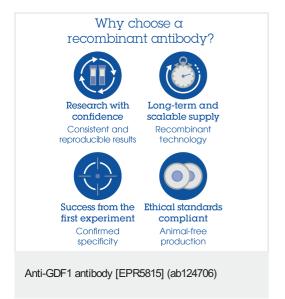
Predicted band size: 39 kDa



Ol-RD Scanning - Anti-GDF1 antibody [EPR5815] (ab124706)

Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about K_D



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