# abcam

### Product datasheet

## Anti-FGF21 antibody [EPR8314(2)] ab171941



★★★★★★ 1 Abreviews 64 References 画像数7

製品の概要

製品名 Anti-FGF21 antibody [EPR8314(2)]

製品の詳細 Rabbit monoclonal [EPR8314(2)] to FGF21

由来種 Rabbit

特異性 The immunogen used for this product shares 6 continuous identical amino acids with SIKE1.

Cross-reactivity with this protein has not been confirmed experimentally.

Expression levels of the target protein vary with sample type and some optimisation may be required (PMID: 27285327). For western blot using cell lines, it may be necessary to collect cell culture supernatant for endogenous FGF21 detection as the target protein is readily secreted

(PMID: 24041694; PMID: 26691139).

適用あり: WB, IHC-P アプリケーション

適用なし: ICC/IF

種交差性 交差種: Mouse, Rat, Human

免疫原 Synthetic peptide within Human FGF21 aa 1-100. The exact sequence is proprietary.

Database link: **Q9NSA1** 

ポジティブ・コントロール Recombinant human FGF21 + lgG1 fusion protein (Fc Chimera Active) (ab108556) can be used

as a positive control in WB. WB: Human fetal liver lysate, mouse spleen, rat spleen. IHC-P:

Human stomach

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 +4°C short term (1-2 weeks). Allow to warm to room temp and agitate

gently before aliquotting. Store at -20°C long term. Avoid freeze / thaw cycle.

パッファー Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 0.05% BSA, 40% Glycerol

精製度 Protein A purified

**ポリモ**ノクローナル **ウローン名** EPR8314(2)

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ab171941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	<b>★★★</b> ☆☆ <u>(1)</u>	1/1000. Predicted molecular weight: 22 kDa.
IHC-P		1/250. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. See IHC antigen retrieval protocols.

追加情報 Is unsuitable for ICC/IF.

#### ターゲット情報

機能 Stimulates glucose uptake in differentiated adipocytes via the induction of glucose transporter

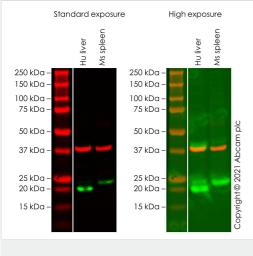
SLC2A1/GLUT1 expression (but not SLC2A4/GLUT4 expression). Activity requires the presence

of KLB

**配列類似性** Belongs to the heparin-binding growth factors family.

細胞内局在 Secreted.

#### 画像



Western blot - Anti-FGF21 antibody [EPR8314(2)] (ab171941)

**All lanes**: Anti-FGF21 antibody [EPR8314(2)] (ab171941) at 1/1000 dilution

Lane 1: Human Liver cell lysate

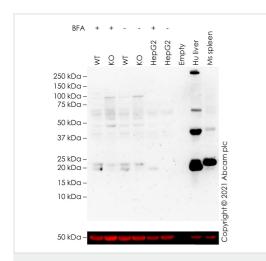
Lane 2: Mouse Spleen cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

**Predicted band size:** 22 kDa **Observed band size:** 21 kDa

False colour image of Western blot: Anti-FGF21 antibody [EPR8314(2)] staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab171941 was shown to bind specifically to FGF21. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween<sup>®</sup> 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit lgG H&L (IRDye<sup>®</sup> 800CW) preabsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye<sup>®</sup> 680RD) preabsorbed (ab216776) at 1/20000 dilution.



Western blot - Anti-FGF21 antibody [EPR8314(2)] (ab171941)

**All lanes :** Anti-FGF21 antibody [EPR8314(2)] (ab171941) at 1/1000 dilution

**Lane 1 :** Wild-type HeLa Treated BFA (5 ug/mL, 6 h) cell lysate at 20 μg

**Lane 2 :** FGF21 knockout HeLa Treated BFA (5  $\mu$ mL, 6 h) cell lysate at 20  $\mu$ g

**Lane 3 :** Wild-type HeLa Vehicle Control BFA (0 ug/mL, 6 h) cell lysate at 20  $\mu$ g

**Lane 4 :** FGF21 knockout HeLa Vehicle Control BFA (0  $\mu$ 0 ug/mL, 6 h) cell lysate at 20  $\mu$ g

Lane 5 : HepG2 Treated BFA (5  $\mu$ mL, 6 h) cell lysate at 20  $\mu$ g

Lane 6: HepG2 cell lysate at 20 µg

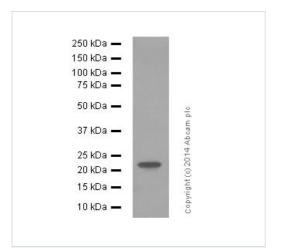
Lane 7: Empty

Lane 8: Human Liver cell lysate at 10 µg
Lane 9: Mouse Spleen cell lysate at 10 µg

Performed under reducing conditions.

**Predicted band size:** 22 kDa **Observed band size:** 21 kDa

False colour image of Western blot: Anti-FGF21 antibody [EPR8314(2)] staining at 1/1000 dilution, shown in black; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab171941 was shown to bind specifically to FGF21. A band was observed at 21 kDa in treated wild-type HeLa cell lysates with no signal observed at this size in FGF21 knockout cell line ab265974 (knockout cell lysate ab256915). To generate this image, wild-type and FGF21 knockout HeLa cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 5 % milk in TBS-0.1 % Tween<sup>®</sup> 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times before development and with highsensitivity chemiluminescence substrate and imaged with 16 minutes exposure time. Secondary antibodies used were HRP conjugated Goat anti-Rabbit (H+L) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) at 1/20000 dilution.



Western blot - Anti-FGF21 antibody [EPR8314(2)] (ab171941)

Anti-FGF21 antibody [EPR8314(2)] (ab171941) at 1/1000 dilution (purified) + Mouse spleen tissue lysate at 10 µg

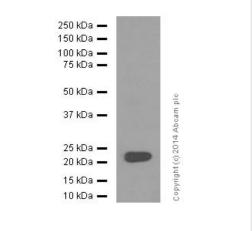
#### Secondary

HRP goat anti-rabbit (H+L) at 1/1000 dilution

Predicted band size: 22 kDa Observed band size: 22 kDa

Blocking buffer: 5% NFDM/TBST

Dilution buffer: 5% NFDM/TBST



Western blot - Anti-FGF21 antibody [EPR8314(2)] (ab171941)



Anti-FGF21 antibody [EPR8314(2)] (ab171941) at 1/5000 dilution (purified) + Rat spleen tissue lysate at 20 μg

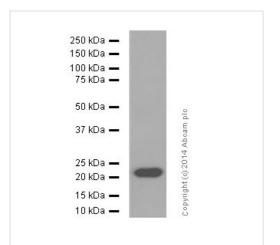
#### Secondary

HRP goat anti-rabbit (H+L) at 1/1000 dilution

Predicted band size: 22 kDa Observed band size: 22 kDa

Blocking buffer: 5% NFDM/TBST

Dilution buffer: 5% NFDM/TBST



Western blot - Anti-FGF21 antibody [EPR8314(2)] (ab171941)

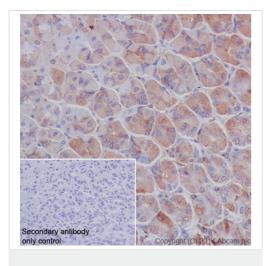
Anti-FGF21 antibody [EPR8314(2)] (ab171941) at 1/5000 dilution (purified) + Human fetal liver tissue lysate at 20 µg

#### Secondary

HRP goat anti-rabbit (H+L) at 1/1000 dilution

Predicted band size: 22 kDa Observed band size: 22 kDa

Blocking buffer: 5% NFDM/TBST Dilution buffer: 5% NFDM/TBST



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-FGF21 antibody [EPR8314(2)] (ab171941)

Immunohistochemical staining of paraffin embedded human stomach with purified ab171941 at a working dilution of 1/250. The secondary antibody used is ab97051, a HRP-conjugated goat antirabbit IgG (H+L), at a dilution of 1/500. The sample is counterstained with hematoxylin. Antigen retrieval was perfomed using Tris-EDTA buffer, pH 9.0. PBS was used instead of the primary antibody as the negative control, and is shown in the inset.



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