abcam

Product datasheet

HRP Anti-Insulin degrading enzyme / IDE antibody [EPR6099] ab201836



リコンピナント

RabMAb

画像数3

製品の概要

製品名 HRP Anti-Insulin degrading enzyme / IDE antibody [EPR6099]

製品の詳細 HRP Rabbit monoclonal [EPR6099] to Insulin degrading enzyme / IDE

由来種 Rabbit HRP 標識

アプリケーション **適用あり:** WB 種交差性 交差種: Human

交差が予測される動物種: Mouse, Rat 🔷

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

ポジティブ・コントロール WB: HeLa, HepG2, A375, and K562 whole cell 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 +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Stable for 12 months at -20°C. Store In the Dark.

バッファー pH: 7.40

Preservative: 0.1% Proclin 300 Solution

Constituents: 30% Glycerol (glycerin, glycerine), PBS, 1% BSA

精製度 Protein A purified

ポリモノ モノクローナル

アイソタイプ

ΙgG

アプリケーション

Abpromise保証は、次のテスト済みアプリケーションにおけるab201836の使用に適用されます The Abpromise guarantee 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		1/5000. Detects a band of approximately 118 kDa (predicted molecular weight: 118 kDa).

ターゲット情報

機能 Plays a role in the cellular breakdown of insulin, IAPP, glucagon, bradykinin, kallidin and other

> peptides, and thereby plays a role in intercellular peptide signaling. Degrades amyloid formed by APP and IAPP. May play a role in the degradation and clearance of naturally secreted amyloid

beta-protein by neurons and microglia.

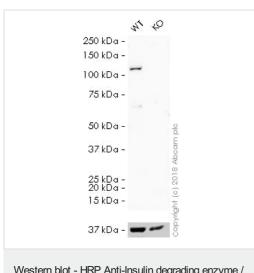
配列類似性 Belongs to the peptidase M16 family.

翻訳後修飾 The N-terminus is blocked.

細胞内局在 Cytoplasm. Cell surface. Present at the cell surface of neuron cells. The membrane-associated

isoform is approximately 5 kDa larger than the known cytosolic isoform.

画像



Western blot - HRP Anti-Insulin degrading enzyme /

IDE antibody [EPR6099] (ab201836)

All lanes: HRP Anti-Insulin degrading enzyme / IDE antibody

[EPR6099] (ab201836) at 1/5000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: IDE (Insulin degrading enzyme / IDE) knockout HAP1

whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 118 kDa Observed band size: 118 kDa

Exposure time: 20 minutes

ab201836 was shown to specifically react with Insulin degrading

enzyme / IDE in wild-type HAP1 cells as signal was lost in IDE (Insulin degrading enzyme / IDE) knockout cells. Wild-type and IDE (Insulin degrading enzyme / IDE) knockout samples were subjected to SDS-PAGE. Ab201836 and ab184095 (Mouse monoclonal [mAbcam 9484] to GAPDH - Loading Control (Alexa Fluor 68 680) loading control) were incubated overnight at 4°C at 1/5000 dilution and 1/20000 dilution respectively. The loading control was imaged using the Licor Odyssey CLx prior to blots being developed with ECL technique.

1 2 3 4

250 kDa —

150 kDa —

100 kDa —

75 kDa —

37 kDa —

25 kDa —

20 kDa —

15 kDa —

15 kDa —

15 kDa —

15 kDa —

16 kDa —

17 kDa —

18 kDa —

19 kDa —

19 kDa —

10 kDa —

10 kDa —

10 kDa —

Western blot - HRP Anti-Insulin degrading enzyme / IDE antibody [EPR6099] (ab201836)

All lanes : HRP Anti-Insulin degrading enzyme / IDE antibody [EPR6099] (ab201836) at 1/5000 dilution

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2 : HepG2 (Human hepatocellular liver carcinoma cell line) Whole Cell Lysate

Lane 3 : A375 (Human melanoma cell line) Whole Cell LysateLane 4 : K562 (Human erythromyeloblastoid leukemia cell line)Nuclear Lysate

Lysates/proteins at 10 µg per lane.

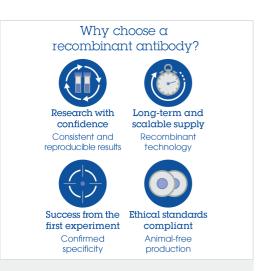
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 118 kDa Observed band size: 118 kDa

Exposure time: 30 seconds

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 2% Bovine Serum Albumin before being incubated with ab201836 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406**.



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