# abcam

# Product datasheet

# Anti-Insulin degrading enzyme / IDE antibody [EPR6099] - BSA and Azide free ab247897



ייבע RabMAb

#### 画像数3

#### 製品の概要

製品名 Anti-Insulin degrading enzyme / IDE antibody [EPR6099] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR6099] to Insulin degrading enzyme / IDE - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: Flow Cyt (Intra), WB

適用なし: IHC-P or IP

種交差性 交差種: Mouse, Human

交差が予測される動物種: Rat 4

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, HAP1, K562 and HepG2 cell lysates.

特記事項 ab247897 is the carrier-free version of ab109538.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.

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

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

**バッファー** pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

**ポリ/モノ** モノクローナル

**クローン名** EPR6099

アイソタイプ lgG

#### 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ab247897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
Flow Cyt (Intra)		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 118 kDa.

追加情報 Is unsuitable for IHC-P or IP.

#### 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.

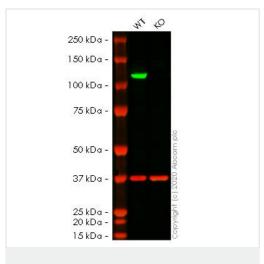
**配列類似性** Belongs to the peptidase M16 family.

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 - Anti-Insulin degrading enzyme / IDE antibody [EPR6099] - BSA and Azide free (ab247897)

**All lanes :** Anti-Insulin degrading enzyme / IDE antibody [EPR6099] (ab109538) at 1/1000 dilution

Lane 1 : Wild-type HeLa cell lysate

Lane 2 : IDE knockout HeLa cell lysate

Lysates/proteins at 20 µg per lane.

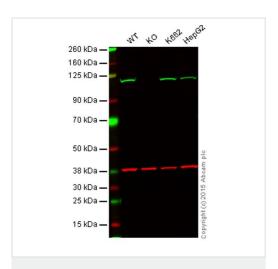
Performed under reducing conditions.

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

This data was developed using the same antibody clone in a different buffer formulation (<u>ab109538</u>).

Lanes 1-2: Merged signal (red and green). Green - <u>ab109538</u> observed at 118 kDa. Red - Anti-GAPDH antibody [6C5] - Loading Control (<u>ab8245</u>) observed at 37 kDa.

ab109538 was shown to react with IDE in wild-type HeLa cells in western blot. Loss of signal was observed when knockout cell line ab261755 (knockout cell lysate ab257197) was used. Wild-type HeLa and IDE knockout HeLa cell lysates were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab109538 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) overnight at 4°C at a 1 in 1000 dilution and a 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye®800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye®680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-Insulin degrading enzyme / IDE antibody [EPR6099] - BSA and Azide free (ab247897)

**All lanes :** Anti-Insulin degrading enzyme / IDE antibody [EPR6099] (ab109538) at 1/2000 dilution

Lane 1: Wild-type HAP1 cell lysate

Lane 2: IDE knockout HAP1 cell lysate,

Lane 3: K562 cell lysate

Lane 4: HepG2 cell lysate

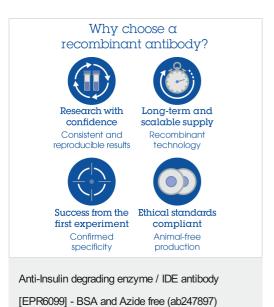
Lysates/proteins at 20 µg per lane.

Predicted band size: 118 kDa

This data was developed using the same antibody clone in a different buffer formulation (<u>ab109538</u>).

**Lanes 1 - 4**: Merged signal (red and green). Green - <u>ab109538</u> observed at 118 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

<u>ab109538</u> was shown to specifically react with IDE in wild-type HAP1 cells. No band was observed when IDE knockout samples were examined. Wild-type and IDE knockout samples were subjected to SDS-PAGE. <u>ab109538</u> and <u>ab8245</u> (loading control to GAPDH) were both diluted 1/2000 and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed <u>ab216773</u> and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed <u>ab216776</u> secondary antibodies at 1/10,000 dilution for 1 hour at room temperature before imaging.



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