# abcam

# Product datasheet

# Anti-CD13 antibody [EPR4058] - Low endotoxin, Azide free ab227111



リコンピナント

RabMAb

3 References 画像数 17

# 製品の概要

製品名 Anti-CD13 antibody [EPR4058] - Low endotoxin, Azide free

製品の詳細 Rabbit monoclonal [EPR4058] to CD13 - Low endotoxin, Azide free

由来種 Rabbit

アプリケーション 適用あり: IHC-P, WB, ICC/IF

適用なし: Flow Cyt or IP

種交差性 交差種: Mouse, Rat, Human

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: THP-1, PANC-1, Rat kidney, HAP1, HeLa and Mouse kidney lysates; IHC-P: human kidney,

liver, hepatocellular carcinoma, prostatic carcinoma, astrocytoma and breast tissues, mouse and

rat kidney tissues; ICC/IF: THP-1 cells

特記事項 ab227111 is the carrier-free version of ab108310.

Our <u>carrier-free</u> 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 cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our <u>conjugation kits</u> 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<sup>®</sup> 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

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monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Our <u>Low endotoxin, azide-free formats</u> have low endotoxin level (≤ 1 EU/ml, determined by the LAL assay) and are free from azide, to achieve consistent experimental results in functional assays.

#### 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** EPR4058

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ab227111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 at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Predicted molecular weight: 110 kDa.
ICC/IF		Use at an assay dependent concentration.

追加情報

Is unsuitable for Flow Cyt or IP.

# ターゲット情報

#### 機能

Broad specificity aminopeptidase. Plays a role in the final digestion of peptides generated from hydrolysis of proteins by gastric and pancreatic proteases. May play a critical role in the pathogenesis of cholesterol gallstone disease. May be involved in the metabolism of regulatory peptides of diverse cell types including small intestinal and tubular epithelial cells, macrophages, granulocytes and synaptic membranes from the CNS. Found to cleave antigen peptides bound to major histocompatibility complex class II molecules of presenting cells and to degrade neurotransmitters at synaptic junctions. Is also implicated as a regulator of IL-8 bioavailability in the endometrium, and therefore may contribute to the regulation of angiogenesis. Is used as a marker for acute myeloid leukemia and plays a role in tumor invasion. In case of human

coronavirus 229E (HCoV-229E) infection, serves as receptor for HCoV-229E spike glycoprotein.

Mediates as well human cytomegalovirus (HCMV) infection.

Expressed in epithelial cells of the kidney, intestine, and respiratory tract; granulocytes, monocytes, fibroblasts, endothelial cells, cerebral pericytes at the blood-brain barrier, synaptic membranes of cells in the CNS. Also expressed in endometrial stromal cells, but not in the endometrial glandular cells. Found in the vasculature of tissues that undergo angiogenesis and in malignant gliomas and lymph node metastases from multiple tumor types but not in blood vessels of normal tissues. A soluble form has been found in plasma. It is found to be elevated in plasma and effusions of cancer patients.

**配列類似性** Belongs to the peptidase M1 family.

ドメイン Amino acids 260-353 are essential to mediate susceptibility to infection with HCoV-229E (in

porcine/human chimeric studies) and more specifically amino acids 288-295 (mutagenesis

studies).

翻訳後修飾 Sulfated.

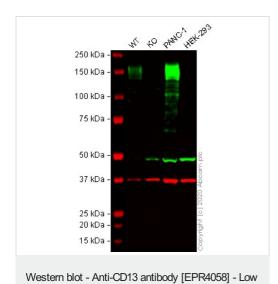
N- and O-glycosylated.

May undergo proteolysis and give rise to a soluble form.

**細胞内局在** Cell membrane. Cytoplasm > cytosol. A soluble form has also been detected.

## 画像

組織特異性



endotoxin, Azide free (ab227111)

**All lanes :** Anti-CD13 antibody [EPR4058] (<u>ab108310</u>) at 1/1000 dilution

Lane 1: Wild-type THP-1 cell lysate

Lane 2: ANPEP knockout THP-1 cell lysate

Lane 3: PANC-1 cell lysate
Lane 4: HEK-293 cell lysate

Lysates/proteins at 30 µg per lane.

Performed under reducing conditions.

Predicted band size: 110 kDa

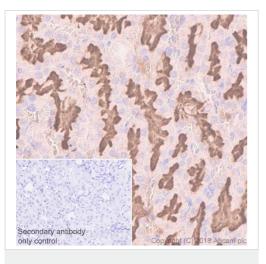
Observed band size: 160 kDa

This data was developed using the same antibody clone in a different buffer formulation (ab108310).

**Lanes 1 - 4:** Merged signal (red and green). Green - <u>ab108310</u> observed at 160 kDa. Red - loading control <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) observed at 37kDa.

<u>ab108310</u> was shown to react with CD13 in wild-type THP-1 cells in western blot with loss of signal observed in ANPEP knockout cell

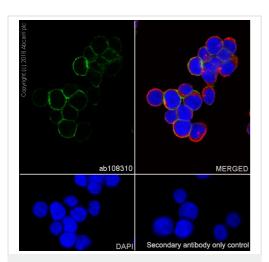
line <u>ab273759</u> (knockout cell lysate <u>ab275505</u>). Wild-type and ANPEP knockout THP-1 cell lysates were subjected to SDS-PAGE. Membranes were blocked in fluorescent western blot (TBS-based) blocking solution before incubation with <u>ab108310</u> and <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) overnight at 4°C at a 1 in 1000 dilution and a 1 in 20000 dilution respectively. Blots were incubated with Goat anti-Rabbit IgG H&L (IRDye<sup>®</sup> 800CW) preabsorbed (<u>ab216773</u>) and Goat anti-Mouse IgG H&L (IRDye<sup>®</sup> 680RD) preabsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat kidney tissue sections labeling CD13 with purified <u>ab108310</u> at 1/1600 dilution (0.43 µg/ml). Heat mediated antigen retrieval was performed using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain. This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab108310</u>)

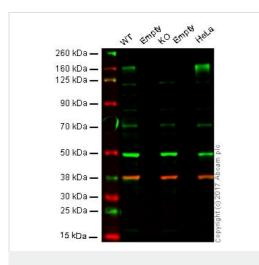


Immunocytochemistry/ Immunofluorescence - Anti-CD13 antibody [EPR4058] - Low endotoxin, Azide free (ab227111)

Confocal image showing membranous staining in THP-1 cells

ab108310 (purified) at 1/100 staining CD13 in the THP-1 (human monocytic leukemia monocyte) cell line by ICC/IF
(Immunocytochemistry/immunofluorescence). Cells were fixed with 100% methanol. Samples were incubated with primary antibody 1/500. ab150077 An Alexa Fluor® 488-conjugated Goat anti-Rabbit IgG at 1/1000 was used as the secondary antibody. Ab195889
Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) 1/200 was used as a counter stain and DAPI was used as a nuclear counter stain.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).



Western blot - Anti-CD13 antibody [EPR4058] - Low endotoxin, Azide free (ab227111)

This WB data was generated using the same anti-CD13 antibody clone, EPR4058, in a different buffer formulation (cat# <u>ab108310</u>).

Lane 1: Wild type HAP1 whole cell lysate (20 μg)

Lane 2: Empty

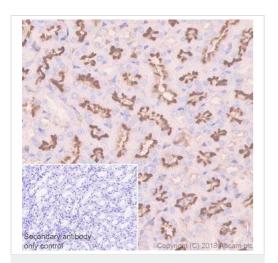
Lane 3: CD13 (KO) whole cell lysate (20 µg)

Lane 4: Empty

Lane 5: HeLa whole cell lysate (20 µg)

**Lanes 1 - 5:** Merged signal (red and green). Green - <u>ab108310</u> observed at 160 kDa. Red - loading control, <u>ab18058</u>, observed at 130 kDa.

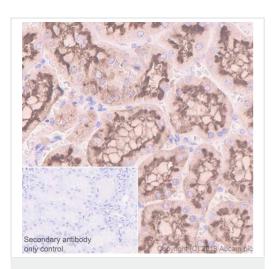
ab108310 was shown to recognize CD13 when CD13 knockout samples were used, along with additional cross-reactive bands. Wild-type and CD13 knockout samples were subjected to SDS-PAGE. Ab108310 and ab18058 (Mouse anti Vinculin loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/10000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

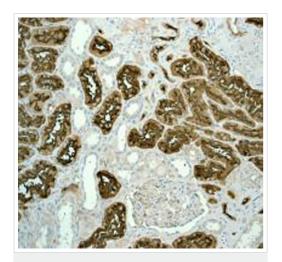
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse kidney tissue sections labeling CD13 with purified <a href="mailto:ab108310">ab108310</a> at 1/1600 dilution (0.43 µg/ml). Heat mediated antigen retrieval was performed using <a href="mailto:ab93684">ab93684</a> (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain. This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue sections labeling CD13 with purified <a href="mailto:ab108310">ab108310</a> at 1/1600 dilution (0.43 µg/ml). Heat mediated antigen retrieval was performed using <a href="mailto:ab93684">ab93684</a> (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain. This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<a href="mailto:ab108310">ab108310</a>)

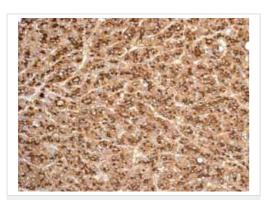


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

<u>ab108310</u> (unpurified), at 1/250, staining CD13 in human kidney tissue by immunohistochemistry.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).



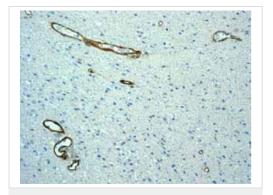
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

<u>ab108310</u> (unpurified) showing positive staining in Hepatocellular carcinoma tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab108310</u>).

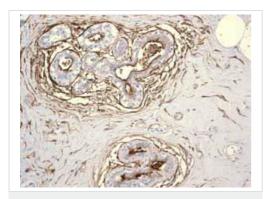
Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody
[EPR4058] - Low endotoxin, Azide free (ab227111)

<u>ab108310</u> showing positive staining in Astrocytoma tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).



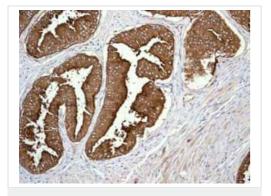
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

<u>ab108310</u> showing positive staining in Normal breast tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



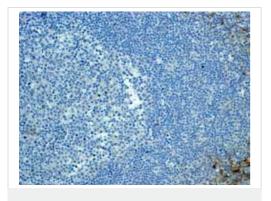
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

ab108310 showing positive staining in Prostatic carcinoma tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.

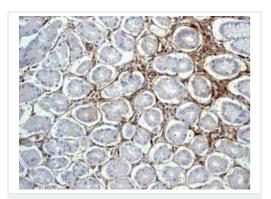


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

ab108310 showing positive staining in Normal tonsil tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab108310</u>).



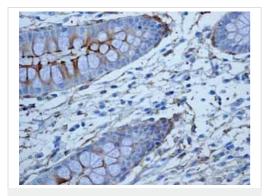
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

ab108310 showing positive staining in Normal stomach tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



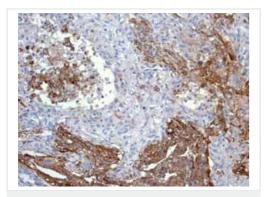
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

ab108310 showing positive staining in Normal colon tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab108310**).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

<u>ab108310</u> showing positive staining in Lung adenocarcinoma tissue.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab108310).



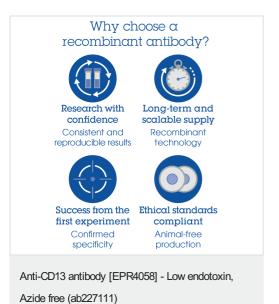
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD13 antibody

[EPR4058] - Low endotoxin, Azide free (ab227111)

This IHC data was generated using the same anti-CD13 antibody clone, EPR4058, in a different buffer formulation (cat# ab108310).

<u>ab108310</u> showing positive staining in Normal liver tissue.

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



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