abcam

Product datasheet

Anti-IL-18 antibody [EPR19954-188] - BSA and Azide free ab243295





RabMAb



概述

产品名称 Anti-IL-18抗体[EPR19954-188] - BSA and Azide free

塩 免单克隆抗体[EPR19954-188] to IL-18 - BSA and Azide free

宿主 Rabbit

经测试应用 适用于: Flow Cyt (Intra), WB, IHC-P

不适用于: ICC/IF

种属反应性 与反应: Human

免疫原 Recombinant full length protein. This information is proprietary to Abcam and/or its suppliers.

阳性对照 IHC-P: Human tonsil, liver and kidney tissues. Flow Cyt (intra): PC-3 cells.

常规说明 ab243295 is the carrier-free version of ab243091.

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[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] 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 monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

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性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C. Do Not Freeze.

存储溶液 pH: 7.2

Constituent: PBS

无载体 是

纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR19954-188

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab243295于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Flow Cyt (Intra)		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 22, 14 kDa (predicted molecular weight: 22 kDa).
IHC-P	★★★☆☆ (1)	Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

应用说明 Is unsuitable for ICC/IF.

靶标

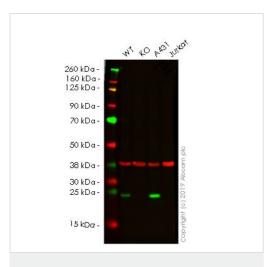
功能 Augments natural killer cell activity in spleen cells and stimulates interferon gamma production in

T-helper type I cells.

序列相似性 Belongs to the IL-1 family.

细**胞定位** Secreted.

图片



Western blot - Anti-IL-18 antibody [EPR19954-188] - BSA and Azide free (ab243295)

All lanes : Anti-IL-18 antibody [EPR19954-188] (**ab243091**) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: IL18 knockout HeLa cell lysate

Lane 3: A431 cell lysate

Lane 4: Jurkat cell lysate

Lysates/proteins at 20 µg per lane.

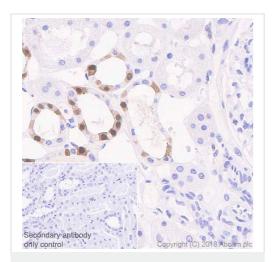
Performed under reducing conditions.

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

This data was developed using the same antibody clone in a different buffer formulation (<u>ab243091</u>).

Lanes 1-4: Merged signal (red and green). Green - <u>ab243091</u> observed at 22 kDa. Red - loading control <u>ab8245</u> observed at 37 kDa.

<u>ab243091</u> Anti-IL-18 antibody [EPR19954-188] was shown to specifically react with IL-18 in wild-type HeLa cells. Loss of signal was observed when knockout cell line <u>ab265274</u> (knockout cell lysate <u>ab256952</u>) was used. Wild-type and IL-18 knockout samples were subjected to SDS-PAGE. <u>ab243091</u> and Anti-alpha Tubulin antibody [EP1332Y] - Microtubule Marker (<u>ab52866</u>) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. 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 in 20000 dilution for 1 hour at room temperature before imaging.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IL-18 antibody

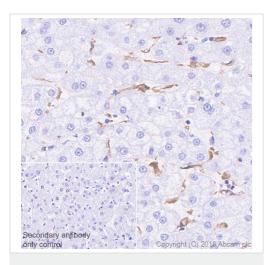
[EPR19954-188] - BSA and Azide free (ab243295)

Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling IL-18 with <u>ab243091</u> at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic and nuclear staining on distal convoluted tubules of human kidney (PMID: 17687255). Counter stained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab243091).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IL-18 antibody

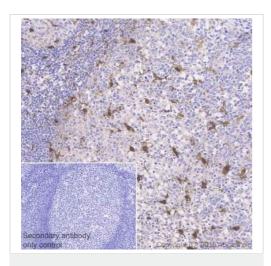
[EPR19954-188] - BSA and Azide free (ab243295)

Immunohistochemical analysis of paraffin-embedded human liver tissue labeling IL-18 with <u>ab243091</u> at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic staining on Kupffer cells in human liver (PMID: 19084941). Counter stained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab243091).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IL-18 antibody

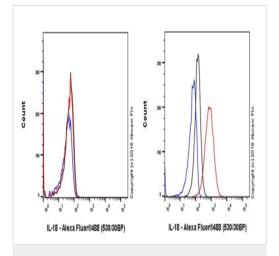
[EPR19954-188] - BSA and Azide free (ab243295)

Immunohistochemical analysis of paraffin-embedded human tonsil tissue labeling IL-18 with <u>ab243091</u> at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Positive staining on subsets of macrophages in tonsil (PMID: 28842466). Counter stained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab243091).

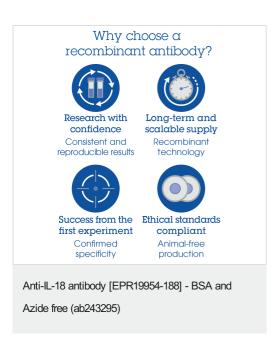


Flow Cytometry (Intracellular) - Anti-IL-18 antibody [EPR19954-188] - BSA and Azide free (ab243295)

Intracellular flow cytometric analysis of 4% paraformal dehyde-fixed, 90% methanol-permeabilized Jurkat (human T cell leukemia cell line from peripheral blood) (left panel) and PC-3 (human prostate adenocarcinoma cell line) (right panel) cells labeling IL-18 with ab243091 at 1/50 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730) (black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150077) at 1/2000 dilution was used as the secondary antibody.

Negative control: Jurkat (PMID 15086390).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab243091).



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