abcam

Product datasheet

Anti-Ikaros antibody [EPR13790] - BSA and Azide free ab206645



ועלטעבעו RabMAb

画像数 5

製品の概要

製品名 Anti-Ikaros antibody [EPR13790] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR13790] to Ikaros - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: Flow Cyt (Intra), IHC-P, WB

交差種: Mouse, Human 種交差性

交差が予測される動物種: 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 Jurkat, Daudi, Raji, Ramos and MOLT4 cell lysates; Human thymus and mouse spleen tissues;

MOLT4 cells.

特記事項 ab206645 is the carrier-free version of ab191394.

> 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[®] 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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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** EPR13790

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ab206645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. ab199376 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Detects a band of approximately 50-70 kDa (predicted molecular weight: 58 kDa).

ターゲット情報

機能 Transcription regulator of hematopoietic cell differentiation (PubMed:17934067). Binds gamma-

satellite DNA (PubMed:17135265, PubMed:19141594). Plays a role in the development of lymphocytes, B- and T-cells. Binds and activates the enhancer (delta-A element) of the CD3-delta gene. Repressor of the TDT (fikzfterminal deoxynucleotidyltransferase) gene during thymocyte differentiation. Regulates transcription through association with both HDAC-dependent and HDAC-independent complexes. Targets the 2 chromatin-remodeling complexes, NuRD and BAF (SWI/SNF), in a single complex (PYR complex), to the beta-globin locus in adult erythrocytes. Increases normal apoptosis in adult erythroid cells. Confers early temporal competence to retinal progenitor cells (RPCs) (By similarity). Function is isoform-specific and is modulated by

dominant-negative inactive isoforms (PubMed:17135265, PubMed:17934067).

組織特異性 Abundantly expressed in thymus, spleen and peripheral blood Leukocytes and lymph nodes.

Lower expression in bone marrow and small intestine.

関連疾患 Defects in IKZF1 are frequent occurrences (28.6%) in acute lymphoblasic leukemia (ALL). Such

alterations or deletions lead to poor prognosis for ALL.

Chromosomal aberrations involving IKZF1 are a cause of B-cell non-Hodgkin lymphomas (B-cell

NHL). Translocation t(3;7)(q27;p12), with BCL6.

Belongs to the Ikaros C2H2-type zinc-finger protein family.

Contains 6 C2H2-type zinc fingers.

ドメイン The N-terminal zinc-fingers 2 and 3 are required for DNA binding as well as for targeting IKFZ1 to

pericentromeric heterochromatin.

The C-terminal zinc-finger domain is required for dimerization.

Phosphorylation controls cell-cycle progression from late G(1) stage to S stage.

Hyperphosphorylated during G2/M phase. Dephosphorylated state during late G(1) phase. Phosphorylation on Thr-140 is required for DNA and pericentromeric location during mitosis. CK2

is the main kinase, in vitro. GSK3 and CDK may also contribute to phosphorylation of the C-terminal serine and threonine residues. Phosphorylation on these C-terminal residues reduces the DNA-binding ability. Phosphorylation/dephosphorylation events on Ser-13 and Ser-295 regulate TDT expression during thymocyte differentiation. Dephosphorylation by protein phosphatase 1 regulates stability and pericentromeric heterochromatin location. Phosphorylated in both lymphoid and non-lymphoid tissues (By similarity). Phosphorylation at Ser-361 and Ser-364 downstream of

SYK induces nuclear translocation.

Sumoylated. Simulataneous sumoylation on the 2 sites results in a loss of both HDAC-dependent and HDAC-independent repression. Has no effect on pericentromeric heterochromatin location.

Desumoylated by SENP1.

Polyubiquitinated.

細胞内局在 Cytoplasm; Nucleus. In resting lymphocytes, distributed diffusely throughout the nucleus. Localizes

to pericentromeric heterochromatin in proliferating cells. This localization requires DNA binding which is regulated by phosphorylation / dephosphorylation events and Nucleus. In resting lymphocytes, distributed diffusely throughout the nucleus. Localizes to pericentromeric heterochromatin in proliferating cells. This localization requires DNA binding which is regulated by

phosphorylation / dephosphorylation events (By similarity).

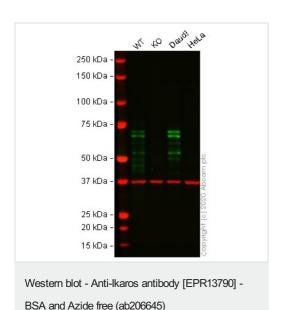
製品の状態

配列類似性

翻訳後修飾

There are 7 isoforms produced by alternative splicing.

画像



All lanes : Anti-lkaros antibody [EPR13790] (ab191394) at 1/10000 dilution

Lane 1: Wild-type Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lane 2: IKZF1 knockout Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lane 3 : Daudi (Human Burkitt's lymphoma cell line) whole cell lysate

Lane 4: HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

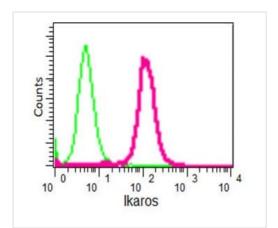
Predicted band size: 58 kDa

Observed band size: 50-70 kDa

This data was developed using the same antibody clone in a different buffer formulation (<u>ab191394</u>).

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab191394</u> observed at 50-70 kDa. Red - loading control <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) observed at 37kDa.

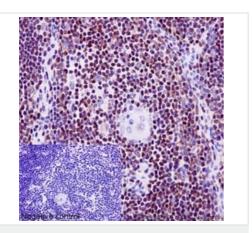
<u>ab191394</u> was shown to react with Ikaros in wild-type Jurkat cells in western blot with loss of signal observed in IKZF1 knockout sample. Wild-type and IKZF1 knockout Jurkat cell lysates were subjected to SDS-PAGE. Membranes were blocked in 3% milk in TBS-T (0.1% Tween[®]) before incubation with <u>ab191394</u> and <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) overnight at 4°C at a 1 in 10000 dilution and a 1 in 20000 dilution respectively. Blots were incubated with Goat anti-Rabbit IgG H&L (IRDye[®] 800CW) preabsorbed (<u>ab216773</u>) and Goat anti-Mouse IgG H&L (IRDye[®] 680RD) preabsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Flow Cytometry (Intracellular) - Anti-Ikaros antibody [EPR13790] - BSA and Azide free (ab206645)

Intracellular flow cytometric analysis of MOLT4 cells (paraformaldehyde-fixed, 2%) labeling lkaros with **ab191394** at 1/130 dilution (red) or a rabbit lgG (negative) (green), followed by Goat anti rabbit lgG (FITC) secondary at 1/150 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab191394**).



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

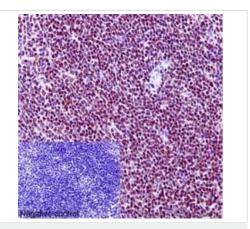
[EPR13790] - BSA and Azide free (ab206645)

This IHC data was generated using the same anti-lkaros antibody clone, EPR13790, in a different buffer formulation (cat# <u>ab191394</u>).

Immunohistochemical analysis of paraffin-embedded Human thumus tissue labeling lkaros with ab191394 at 1/1400

thymus tissue labeling lkaros with **ab191394** at 1/1400 dilution followed by pre-diluted HRP Polymer for Rabbit lgG secondary antibody and counter-stained with Hematoxylin. Inset: Negative control: using PBS instead of primary antibody.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



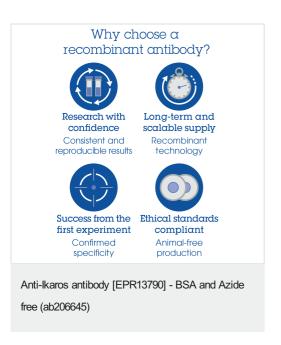
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ikaros antibody
[EPR13790] - BSA and Azide free (ab206645)

This IHC data was generated using the same anti-lkaros antibody clone, EPR13790, in a different buffer formulation (cat# <u>ab191394</u>).

Immunohistochemical analysis of paraffin-embedded Mouse spleen tissue labeling Ikaros with <u>ab191394</u> at 1/1400 dilution followed by pre-diluted HRP Polymer for Rabbit IgG secondary antibody and counter-stained with Hematoxylin.

Inset: Negative control: using PBS instead of primary antibody.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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