# abcam

# Product datasheet

# Anti-FOXO3A antibody [EPR1950] - BSA and Azide free ab232393



リコンピナント

RabMAb

#### 画像数3

# 製品の概要

特記事項

製品名 Anti-FOXO3A antibody [EPR1950] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR1950] to FOXO3A - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: Flow Cyt (Intra), WB

種交差性 交差種: Human

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 Flow Cyt (intra): HeLa cells. WB: HEK-293 and HEK-293T cell lysates.

ab232393 is the carrier-free version of ab109629.

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<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **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** EPR1950

アイソタイプ IgG

#### 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ab232393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: 71 kDa.

# ターゲット情報

機能 Transcriptional activator which triggers apoptosis in the absence of survival factors, including

neuronal cell death upon oxidative stress. Recognizes and binds to the DNA sequence 5'-

[AG]TAAA[TC]A-3'.

組織特異性 Ubiquitous.

関連疾患 Note=A chromosomal aberration involving FOXO3 is found in secondary acute leukemias.

Translocation t(6;11)(q21;q23) with MLL/HRX.

**配列類似性** Contains 1 fork-head DNA-binding domain.

翻訳後修飾 In the presence of survival factors such as IGF-1, phosphorylated on Thr-32 and Ser-253 by

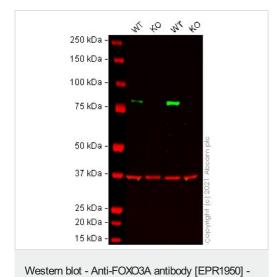
AKT1/PKB. This phosphorylated form then interacts with 14-3-3 proteins and is retained in the cytoplasm. Survival factor withdrawal induces dephosphorylation and promotes translocation to the nucleus where the dephosphorylated protein induces transcription of target genes and triggers apoptosis. Although AKT1/PKB doesn't appear to phosphorylate Ser-315 directly, it may activate other kinases that trigger phosphorylation at this residue. Phosphorylated by STK4 on Ser-209

upon oxidative stress, which leads to dissociation from YWHAB/14-3-3-beta and nuclear

translocation. Phosphorylated by PIM1.

**細胞内局在** Cytoplasm > cytosol. Nucleus. Translocates to the nucleus upon oxidative stress and in the

absence of survival factors.



BSA and Azide free (ab232393)

**All lanes :** Anti-FOXO3A antibody [EPR1950] (<u>ab109629</u>) at 1/1000 dilution

Lane 1: Wild-type HEK-293T cell lysate

Lane 2: Human FOXO3 (FOXO3A) knockout HEK-293T cell lysate (ab256922)

Lane 3: Wild-type HEK-293 cell lysate

Lane 4: Human FOXO3 (FOXO3A) knockout HEK-293 cell lysate (ab261649)

Lysates/proteins at 20 µg per lane.

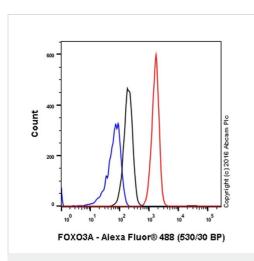
Performed under reducing conditions.

Predicted band size: 71 kDa
Observed band size: 82 kDa

This data was developed using the same antibody clone in a different buffer formulation (ab109629).

**Lanes 1 - 4:** Merged signal (red and green). Green - <u>ab109629</u> observed at 82 kDa. Red - loading control <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) observed at 37 kDa.

ab109629 was shown to react with FOXO3A in wild-type cells in Western blot with loss of signal observed in FOXO3 knockout cells. Wild-type and FOXO3 knockout cell lysates were subjected to SDS-PAGE. Membranes were blocked in 3 % milk in TBS-T (0.1 % Tween®) before incubation with ab109629 and ab8245 (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® 800CW) preabsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 h at room temperature before imaging.



Flow Cytometry (Intracellular) - Anti-FOXO3A antibody [EPR1950] - BSA and Azide free (ab232393)

Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma) cells labeling FOXO3A with unpurified  $\underline{ab109629}$  at 1/150 dilution (10µg/ml) (red). Cells were fixed with 4% paraformaldehyde and permeabilized with 90% methanol. A Goat anti-rabbit lgG (Alexa Fluor® 488) (1/2000 dilution) was used as the secondary antibody. Rabbit monoclonal lgG (Black) was used as the isotype control, cells without incubation with primary antibody and secondary antibody (Blue) were used as the unlabeled control.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab109629).



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