# abcam

## Product datasheet

## Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free ab248343

יובעבלי RabMAb

## 画像数7

#### 製品の概要

製品名 Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR1891(2)] to Bad (phospho S112) - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: Dot blot, IP, WB

適用なし: ICC/IF or IHC-P

種交差性 交差種: Rat. Human

交差が予測される動物種: Mouse 🔷

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

特記事項 ab248343 is the carrier-free version of ab129192.

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

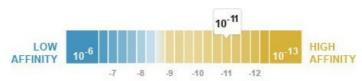
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#### 製品の特性

製品の状態 Liquid

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

**解離定数(K<sub>D</sub>値)** K<sub>D</sub> = 4.70 x 10 <sup>-11</sup> M



Learn more about K<sub>D</sub>

**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** EPR1891(2)

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ab248343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
Dot blot		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 18 kDa.

追加情報 Is unsuitable for ICC/IF or IHC-P.

#### ターゲット情報

機能 Promotes cell death. Successfully competes for the binding to Bcl-X(L), Bcl-2 and Bcl-W, thereby

affecting the level of heterodimerization of these proteins with BAX. Can reverse the death

 $repressor\ activity\ of\ Bcl-X(L),\ but\ not\ that\ of\ Bcl-2\ (By\ similarity).\ Appears\ to\ act\ as\ a\ link\ between$ 

growth factor receptor signaling and the apoptotic pathways.

組織特異性 Expressed in a wide variety of tissues.

配列類似性 Belongs to the Bcl-2 family.

ドメイン Intact BH3 motif is required by BIK, BID, BAK, BAD and BAX for their pro-apoptotic activity and

for their interaction with anti-apoptotic members of the Bcl-2 family.

翻訳後修飾 Phosphorylated on one or more of Ser-75, Ser-99, Ser-118 and Ser-134 in response to survival

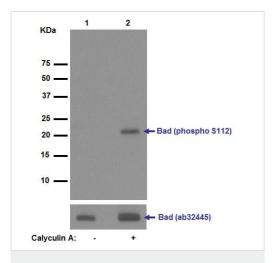
stimuli, which blocks its pro-apoptotic activity. Phosphorylation on Ser-99 or Ser-75 promotes

heterodimerization with 14-3-3 proteins. This interaction then facilitates the phosphorylation at Ser-118, a site within the BH3 motif, leading to the release of Bcl-X(L) and the promotion of cell survival. Ser-99 is the major site of AKT/PKB phosphorylation, Ser-118 the major site of protein kinase A (CAPK) phosphorylation. Ser-75 is phosphorylated by AKT/PKB, protein kinase A and PIM2.

#### 細胞内局在

Mitochondrion outer membrane. Cytoplasm. Upon phosphorylation, locates to the cytoplasm.

#### 画像



Western blot - Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free (ab248343) **All lanes :** Anti-Bad (phospho S112) antibody [EPR1891(2)] (ab129192) at 1/5000 dilution (purified)

Lane 1: Untreated C6 cells

Lane 2: C6 cells treated with Calcyculin A

Lysates/proteins at 10 µg per lane.

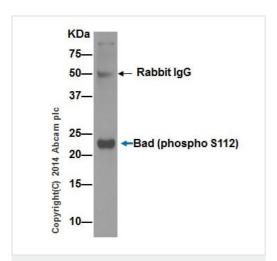
## **Secondary**

All lanes: HRP goat anti-rabbit (H+L) at 1/1000 dilution

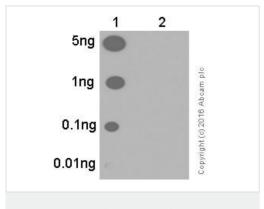
**Predicted band size:** 18 kDa **Observed band size:** 23 kDa

This data was developed using <u>ab129192</u>, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.



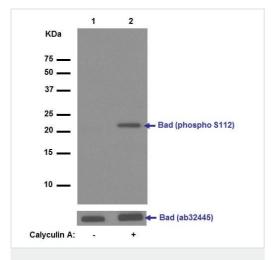
Immunoprecipitation - Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free (ab248343) This data was developed using **ab129192**, the same antibody clone in a different buffer formulation.**ab129192** (purified) at 1/30 immunoprecipitating Bad in HeLa cell lysate treated with Calcyculin A (Lane 1). For western blotting a HRP-conjugated anti-rabbit lgG was used as the secondary antibody (1/1000).Blocking buffer and concentration: 5% NFDM/TBST.Diluting buffer and concentration: 5% NFDM/TBST.



Dot Blot - Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free (ab248343)

This data was developed using <u>ab129192</u>, the same antibody clone in a different buffer formulation.Dot blot analysis of Bad (phospho S112) using <u>ab129192</u> primary antibody at a dilution of 1/1000. <u>ab97051</u> was used as a secondary antibody at a dilution of 1/100,000. **Lane 1:** Bad (pS112) phospho peptide

**Lane 2:** Bad non-phospho peptide **Blocking and diluting buffer:** 5% NFDM/TBST



Western blot - Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free (ab248343)

**All lanes :** Anti-Bad (phospho S112) antibody [EPR1891(2)] (ab129192) at 1/20000 dilution (purified)

Lane 1: Untreated HeLa cell lysate

Lane 2: HeLa cell lysate treated with Calcyculin A

Lysates/proteins at 10 µg per lane.

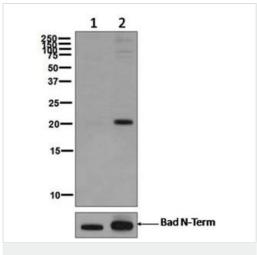
## Secondary

All lanes: HRP goat anti-rabbit (H+L) at 1/1000 dilution

**Predicted band size:** 18 kDa **Observed band size:** 23 kDa

This data was developed using <u>ab129192</u>, the same antibody clone in a different buffer formulation.

**Blocking and dilution buffer:** 5% NFDM/TBST.



Western blot - Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free (ab248343)

**All lanes :** Anti-Bad (phospho S112) antibody [EPR1891(2)] (ab129192) at 1/1000 dilution

Lane 1: HeLa cell lysates, untreated

Lane 2: HeLa cell lysates, treated with Calyculin A

Lysates/proteins at 10 µg per lane.

#### Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

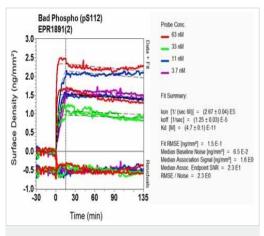
Predicted band size: 18 kDa

This data was developed using <u>ab129192</u>, the same antibody clone in a different buffer formulation.

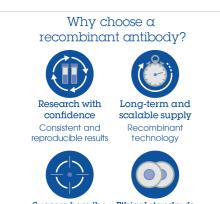
This data was developed using  $\underline{ab129192}$ , the same antibody clone in a different buffer formulation. Equilibrium disassociation constant ( $K_D$ )

Learn more about K<sub>D</sub>

Click here to learn more about KD



OI-RD Scanning - Anti-Bad (phospho S112) antibody [EPR1891(2)] - BSA and Azide free (ab248343)



first experiment Confirmed specificity

Success from the Ethical standards compliant Animal-free production

Anti-Bad (phospho S112) antibody [EPR1891(2)] -BSA and Azide free (ab248343)

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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