# abcam

# Product datasheet

# Anti-p23 antibody [EPR3846] ab92503





重组 RabMAb

★★★★★ 1 Abreviews 5 References 4 图像

概述

产品名称 Anti-p23抗体[EPR3846]

描述 兔单克隆抗体[EPR3846] to p23

宿主 Rabbit

经测试应用 适用于: WB, ICC/IF

不适用于: Flow Cyt or IHC-P

种属反应性 与反应: Human

预测可用于: Mouse, Rat 📤

免疫原 Synthetic peptide within Human p23 aa 50-150. The exact sequence is proprietary.

阳性对照 WB: HEK-293T, HeLa, Human spleen, Jurkat and HepG2 cell lysates ICC/IF: HeLa cells

This product is a recombinant monoclonal antibody, which offers several advantages including: 常规说明

- 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**.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

存储溶液 pH: 7.20

Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, 40% Glycerol (glycerin, glycerine), 9.85% Tris glycine, 50% Tissue

culture supernatant

纯度 Protein A purified

克降 单克隆 克隆编号 **EPR3846** 

### 应用

# The Abpromise guarantee Abpromise™承诺保证使用ab92503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
WB	*** <u>*</u>	1/10000 - 1/50000. Predicted molecular weight: 19 kDa.
ICC/IF		1/100 - 1/250.

应用说明

Is unsuitable for Flow Cyt or IHC-P.

## 靶标

功能 Molecular chaperone that localizes to genomic response elements in a hormone-dependent

manner and disrupts receptor-mediated transcriptional activation, by promoting disassembly of

transcriptional regulatory complexes.

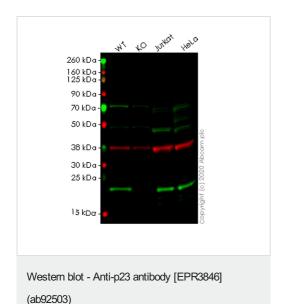
通路 Lipid metabolism; prostaglandin biosynthesis.

序列相似性 Belongs to the p23/wos2 family.

Contains 1 CS domain.

细胞定位 Cytoplasm.

# 图片



**All lanes :** Anti-p23 antibody [EPR3846] (ab92503) at 1/1000 dilution

Lane 1: Wild-type HEK-293T (Human epithelial cell line from embryonic kidney transformed with large T antigen) whole cell lysate

Lane 2: PTGES3 knockout HEK-293T (Human epithelial cell line from embryonic kidney transformed with large T antigen) whole cell lysate

**Lane 3 :** Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

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

Lysates/proteins at 20 µg per lane.

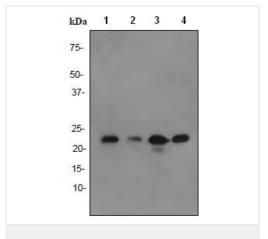
#### Secondary

All lanes: Goat anti-Rabbit lgG H&L (IRDye® 800CW)

**Predicted band size:** 19 kDa **Observed band size:** 23 kDa

**Lanes 1-4:** Merged signal (red and green). Green - ab92503 observed at 23 kDa. Red - loading control **ab8245** observed at 36 kDa.

ab92503 Anti-p23 antibody [EPR3846] was shown to specifically react with p23 in wild-type HEK-293T cells. Loss of signal was observed when knockout cell line <a href="mailto:ab266791">ab266791</a> (knockout cell lysate <a href="mailto:ab258151">ab258151</a>) was used. Wild-type and p23 knockout samples were subjected to SDS-PAGE. ab92503 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-p23 antibody [EPR3846] (ab92503)

**All lanes :** Anti-p23 antibody [EPR3846] (ab92503) at 1/10000 dilution

Lane 1: HeLa cell lysate

Lane 2: Human spleen lysate

Lane 3 : Jurkat cell lysate

Lane 4: HepG2 cell lysate

Lysates/proteins at 10 µg per lane.

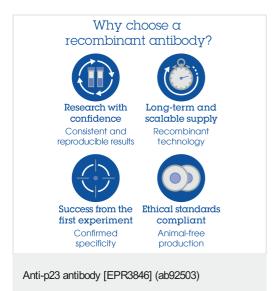
#### Secondary

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

Predicted band size: 19 kDa Observed band size: 23 kDa



Immunocytochemistry/ Immunofluorescence - Antip23 antibody [EPR3846] (ab92503) Immunofluorescent staining of HeLa cells, using ab92503 at a dilution of 1/100.



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

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