# abcam

# Product datasheet

# Anti-Tissue Factor antibody [EPR22548-232] - BSA and Azide free ab254010





RabMAb

1 References 7 图像

### 概述

产品名称 Anti-Tissue Factor抗体[EPR22548-232] - BSA and Azide free

描述 兔单克隆抗体[EPR22548-232] to组织Factor - BSA and Azide free

**宿主** Rabbit

经测试应用 适用于: WB, IHC-P, ICC/IF, Flow Cyt, IP

种属反应性 与反应: Human

免疫原 Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

阳性对照 IHC-P: Human cervix carcinoma and kidney tissue. ICC/IF: A431 cells. Flow Cyt: A431 cells. IP:

Tissue Factor IP in A431 whole cell lysate.

常规说明 ab254010 is the carrier-free version of ab228968.

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 **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<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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#### 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** EPR22548-232

**同种型** IgG

#### 应用

## The Abpromise guarantee Abpromise™承诺保证使用ab254010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		1/1000. Detects a band of approximately 47 kDa (predicted molecular weight: 33 kDa).
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.
ICC/IF		Use at an assay dependent concentration.
Flow Cyt		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.

# 靶标

功能 Initiates blood coagulation by forming a complex with circulating factor VII or VIIa. The [TF:VIIa]

complex activates factors IX or X by specific limited protolysis. TF plays a role in normal

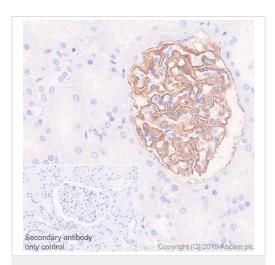
hemostasis by initiating the cell-surface assembly and propagation of the coagulation protease

cascade.

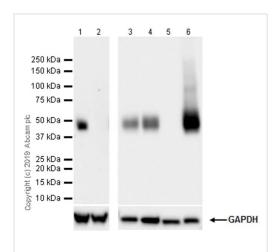
序列相似性 Belongs to the tissue factor family.

细胞定位 Membrane.

# 图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Tissue Factor antibody [EPR22548-232] - BSA and Azide free (ab254010)



Western blot - Anti-Tissue Factor antibody
[EPR22548-232] - BSA and Azide free (ab254010)

Immunohistochemical analysis of paraffinembedded human kidney tissue labeling Tissue Factor with <a href="mailto:ab228968">ab228968</a> at 1/500 dilution, followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (<a href="mailto:ab209101">ab209101</a>). Positive staining in human renal glomerulus (PMID: 7684196). The section was incubated with <a href="mailto:ab228968">ab228968</a> for 30 mins at RT. The immunostaining was performed on a Leica Biosystems BOND<sup>®</sup> RX instrument. Counterstained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Perform heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20mins.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab228968).

#### All lanes:

Lane 1: Wild-type HAP1 whole cell lysate at 60 µg

Lane 2: Tissue Factor knockout HAP1 whole cell lysate at 60 µg

Lane 3: A431 (human epidermoid carcinoma epithelial cell), whole cell lysate at 20 µg

**Lane 4 :** MDA-MB-231 (human breast adenocarcinoma epithelial cell), whole cell lysate at 20  $\mu g$ 

**Lane 5 :** MCF7 (human breast adenocarcinoma epithelial cell), whole cell lysate at 20  $\mu g$ 

Lane 6: BxPC-3 (human pancreas adenocarcinoma epithelial cell), whole cell lysate at 20 µg

#### Secondary

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/100000 dilution

**Predicted band size:** 33 kDa **Observed band size:** 47 kDa

<u>ab228968</u> was shown to specifically react with Tissue Factor in wild-type HAP1 cells as signal was lost in Tissue Factor knockout cells. Wild-type and Tissue Factor knockout samples were subjected to SDS-PAGE. <u>ab228968</u> and <u>ab181602</u> (Rabbit anti-

GAPDH loading control) were incubated 1 hour at room temperature at 1/1000 dilution and 1/200,000 dilution respectively. Blots were developed with Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated (ab97051) secondary antibody at 1/100,000 dilution for 1 hour at room temperature before imaging. The blot was developed on a BIO-RAD<sup>®</sup> ChemiDoc<sup>™</sup> MP instrument using the ECL technique.

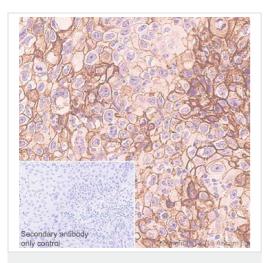
The molecular weight observed is consistent with what has been described in the literature (PMID: 28938620).

Low expression: MCF-7 (PMID: 24137414, 28938620).

Exposure times: Lanes 1-2: 70 secs; Lanes 3-6: 3.25 secs.

Blocking/Dilution buffer: 5% NFDM/TBST.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab228968).

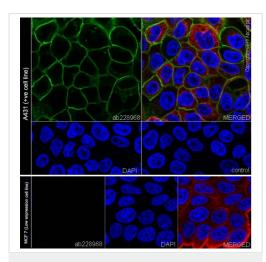


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Tissue Factor antibody
[EPR22548-232] - BSA and Azide free (ab254010)

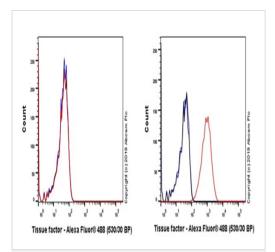
Immunohistochemical analysis of paraffin-embedded human cervix carcinoma tissue labeling Tissue Factor with <u>ab228968</u> at 1/500 dilution, followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>). Membranous and weak cytoplasmic staining on tumor cells of human cervix carcinoma (PMID: 26383146). The section was incubated with <u>ab228968</u> for 30 mins at RT. The immunostaining was performed on a Leica Biosystems BOND<sup>®</sup> RX instrument. Counterstained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

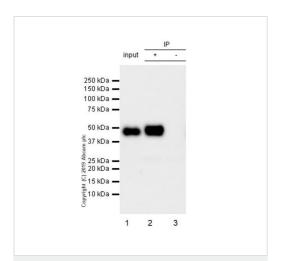
Perform heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20mins.



Immunocytochemistry/ Immunofluorescence - Anti-Tissue Factor antibody [EPR22548-232] - BSA and Azide free (ab254010)



Flow Cytometry - Anti-Tissue Factor antibody [EPR22548-232] - BSA and Azide free (ab254010)



Immunoprecipitation - Anti-Tissue Factor antibody [EPR22548-232] - BSA and Azide free (ab254010)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilizedA431 (human epidermoid carcinoma epithelial cell) and MCF7 () cells labeling Tissue Factor with <a href="mailto:ab228968">ab228968</a> at 1/100 dilution, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (<a href="mailto:ab150077">ab150077</a>) secondary antibody at 1/1000 dilution (green). Confocal image showing membranous staining in A431 cell line. Low expression: MCF7 PMID: 24137414, 28938620. The nuclear stain is DAPI (blue).

Counterstained with <u>ab195889</u> Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor<sup>®</sup> 594) at a 1/200 dilution (red). The negative control is the secondary antibody only.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab228968).

Flow cytometric analysis of MCF7 (human breast adenocarcinoma epithelial cell, Left) / A431 (human epidermoid carcinoma epithelial cell, Right) labeling Tissue Factor with <u>ab228968</u> at 1/500 (red) compared with a Rabbit monoclonal IgG (<u>ab172730</u>) (black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat anti rabbit IgG (Alexa Fluor<sup>®</sup> 488, <u>ab150077</u>), at 1/2000 dilution was used as the secondary antibody.

Low expression: MCF7 (PMID: 24137414, PMID: 28938620).

Gated on viable cells.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab228968).

Tissue Factor was immunoprecipitated from 0.35 mg A431 (human epidermoid carcinoma epithelial cell) whole cell lysate with <a href="mailto:ab228968">ab228968</a> at 1/30 dilution. Western blot was performed from the immunoprecipitate using <a href="mailto:ab228968">ab228968</a> at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (<a href="mailto:ab131366">ab131366</a>), was used for detection at 1/5000 dilution.

Lane 1: A431 whole cell lysate 10 µg (Input).

Lane 2: ab228968 IP in A431 whole cell lysate.

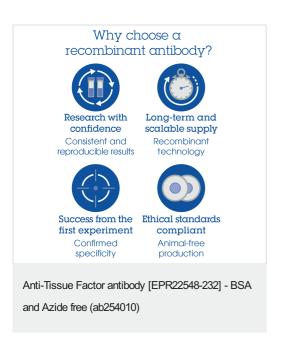
**Lane 3:** Rabbit monoclonal lgG (<u>ab172730</u>) instead of <u>ab228968</u> in A431 whole cell lysate.

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: 30 seconds.

This data was developed using the same antibody clone in a

different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab228968).



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