abcam

Product datasheet

Anti-TNFAIP3 antibody [EPR2663] ab92324



ייבער RabMAb

★★★★★ 1 Abreviews 31 References 画像数 10

製品の概要

製品名 Anti-TNFAIP3 antibody [EPR2663]

製品の詳細 Rabbit monoclonal [EPR2663] to TNFAIP3

由来種 Rabbit

特異性 Mouse species is recommended based on WB results, we do not guarantee IHC-P for

mouse.

アプリケーション 適用あり: WB, IHC-P

適用なし: Flow Cyt (Intra) or ICC/IF

種交差性 交差種: Mouse, Human

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 WB: WEHI-3 treated with TNF (ab9642), Jurkat treated with TNF (ab9642) + TPA, Jurkat treated

with 5ng/ml PMA for 48 hours and then treated with 2µg/ml PHA for 48 hours, HeLa, A549 and

Daudi cell lysates. IHC-P: Human kidney tissue.

特記事項 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**.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Stable for 12 months at -20°C.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol, 0.05% BSA

精製度 Protein A purified

ポリÆノ モノクローナル **ウローン名** EPR2663 **アイソタイプ** IgG

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab92324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。

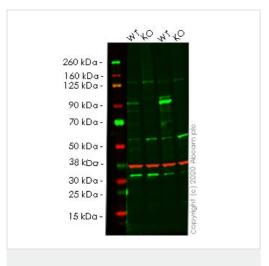
アプリケーション	Abreviews	特記事項
WB		1/1000 - 1/5000. Predicted molecular weight: 90 kDa.
IHC-P	**** (1)	1/50 - 1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. Mouse species is recommended based on WB results, we do not guarantee IHC-P for mouse.

追加情報 Is unsuitable for Flow Cyt (Intra) or ICC/IF.

ターゲット情報

機能	Ubiquitin-editing enzyme that contains both ubiquitin ligase and deubiquitinase activities. Essential component of a ubiquitin-editing protein complex, comprising also RNF11, ITCH and TAX1BP1, that ensures the transient nature of inflammatory signaling pathways. Upon TNF stimulation, deubiquitinates 'Lys-63'-polyubiquitin chains on RIPK1 and catalyzes the formation of 'Lys-48'-polyubiquitin chains. This leads to RIPK1 proteasomal degradation and consequently termination of the TNF- or LPS-mediated activation of NF-kappa-B. In vitro able to deubiquitinate both 'Lys-48'- and 'Lys-63' polyubiquitin chains. Inhibitor of programmed cell death. Has a role in the function of the lymphoid system.
配列類似性	Belongs to the peptidase C64 family. Contains 7 A20-type zinc fingers. Contains 1 OTU domain.
ドメイン	The A20-type zinc fingers mediate the ubiquitin ligase activity. The OTU domain mediates the deubiquitinase activity.
細胞内局在	Cytoplasm. Nucleus.

画像



Western blot - Anti-TNFAIP3 antibody [EPR2663] (ab92324)

All lanes : Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/1000 dilution

Lane 1: Wild-type A549 cell lysate

Lane 2: TNFAIP3 knockout A549 cell lysate

Lane 3: Wild-type HeLa cell lysate

Lane 4: TNFAIP3 knockout HeLa cell lysate

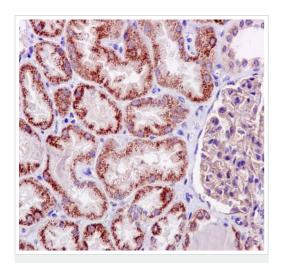
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 90 kDa **Observed band size:** 90 kDa

Lanes 1-4: Merged signal (red and green). Green - ab92324 observed at 90 kDa. Red - Anti-GAPDH antibody [6C5] - Loading Control (ab8245) observed at 37 kDa.

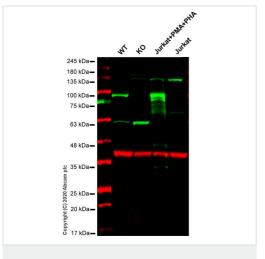
ab92324 was shown to react with TNFAIP3 in wild-type A549 cells in western blot. Loss of signal was observed when knockout cell line ab266946 (knockout cell lysate ab257114) was used. Wild-type A549 and TNFAIP3 knockout A549 cell lysates were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab92324 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) overnight at 4°C at a 1 in 1000 dilution and a 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.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TNFAIP3 antibody

[EPR2663] (ab92324)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue labelling TNFAIP3 with unpurified ab92324 at 1/50. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody. Counterstained with Hematoxylin.



Western blot - Anti-TNFAIP3 antibody [EPR2663] (ab92324)

All lanes : Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: TNFAIP3 knockout HeLa cell lysate

Lane 3: Jurkat cell treated with 5ng/ml PMA for 48 hours and then

treated with 2µg/ml PHA for 48 hours, whole cell lysate

Lane 4: Untreated Jurkat cell lysate

Lysates/proteins at 20 µg per lane.

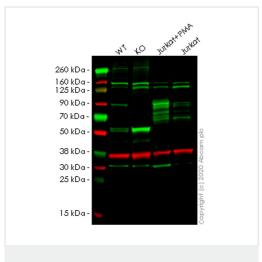
Performed under reducing conditions.

Predicted band size: 90 kDa **Observed band size:** 80 kDa

Lanes 1-4: Merged signal (red and green). Green - ab92324 observed at 80 kDa. Red - loading control, **ab8245** observed at 37 kDa.

ab92324 Anti-TNFAIP3 antibody [EPR2663] was shown to specifically react with TNFAIP3 in wild-type HeLa cells. Loss of signal was observed when knockout cell line **ab265983** (knockout

cell lysate <u>ab257112</u>) was used. Wild-type and TNFAIP3 knockout samples were subjected to SDS-PAGE. ab92324 and Anti-GAPDH antibody [6C5] - Loading Control (<u>ab8245</u>) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye[®] 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse IgG H&L (IRDye[®] 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 10000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-TNFAIP3 antibody [EPR2663] (ab92324)

All lanes : Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/1000 dilution

Lane 1 : Wild-type A549 (Human lung carcinoma cell line) whole cell lysate

Lane 2 : TNFAIP3 knockout A549 (Human lung carcinoma cell line) whole cell lysate

Lane 3: Jurkat (Human T cell leukemia cell line from peripheral blood) cell treated with 5ng/ml PMA for 48 hours and then treated with 2µg/ml PHA for 48 hours, whole cell lysate

Lane 4: Untreated Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (<u>ab216773</u>) at 1/10000 dilution

Predicted band size: 90 kDa
Observed band size: 80 kDa

Lanes 1-4: Merged signal (red and green). Green - ab92324 observed at 80 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

ab92324 Anti-TNFAIP3 antibody [EPR2663] was shown to specifically react with TNFAIP3 in wild-type A549 cells. Loss of signal was observed when knockout cell line <u>ab266945</u> (knockout cell lysate <u>ab266945</u> (knockout cell lysate <u>ab266945</u> (knockout samples were subjected to SDS-PAGE. ab92324 and Anti-GAPDH antibody [6C5] - Loading Control (<u>ab8245</u>) 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-TNFAIP3 antibody [EPR2663] (ab92324)

All lanes : Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/5000 dilution

Lane 1: WEHI-3 (Mouse leukemia lymphoblast) whole cell lysate

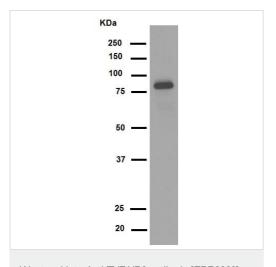
Lane 2: WEHI-3 treated with 20 ng/ml TNF alpha (ab9642) for 6 h

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 90 kDa Observed band size: 80 kDa



Western blot - Anti-TNFAIP3 antibody [EPR2663] (ab92324)

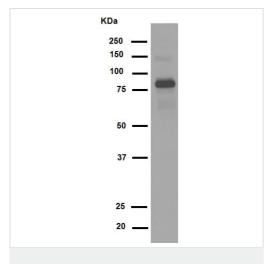
Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/2000 dilution (unpurified) + Jurkat cell lysate - treated with TNF (ab9642) and TPA at 10 μg

Secondary

Peroxidase-conjugated goat anti-rabbit IgG (H+L) at 1/1000 dilution

Predicted band size: 90 kDa
Observed band size: 80 kDa

Blocking and dilution buffer: 5% NFDM/TBST.



Western blot - Anti-TNFAIP3 antibody [EPR2663] (ab92324)

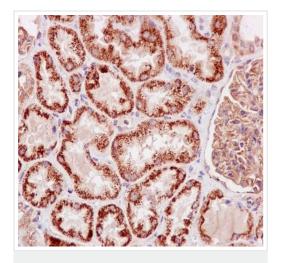
Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/3000 dilution (purified) + Jurkat cell lysate - treated with TNF (ab9642) and TPA at 10 μg

Secondary

Peroxidase-conjugated goat anti-rabbit IgG (H+L) at 1/1000 dilution

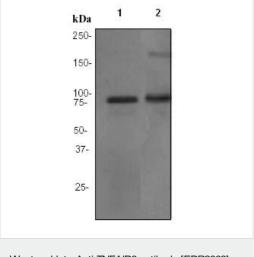
Predicted band size: 90 kDa **Observed band size:** 80 kDa

Blocking and dilution buffer: 5% NFDM/TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TNFAIP3 antibody
[EPR2663] (ab92324)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue labelling TNFAIP3 with purified ab92324 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Counterstained with Hematoxylin.



Western blot - Anti-TNFAIP3 antibody [EPR2663] (ab92324)

All lanes : Anti-TNFAIP3 antibody [EPR2663] (ab92324) at 1/1000 dilution (unpurified)

Lane 1: Jurkat cells treated with TNF (ab9642) and TPA

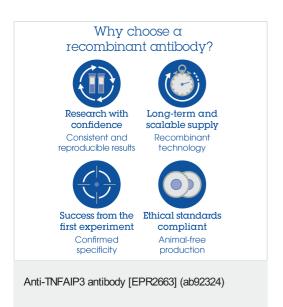
Lane 2: Daudi cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

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

Predicted band size: 90 kDa



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