abcam

Product datasheet

Anti-TNPO3 antibody [3152C2a] ab54353

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Overview

Product name Anti-TNPO3 antibody [3152C2a]

Description Mouse monoclonal [3152C2a] to TNPO3

Host species Mouse

Tested applications Suitable for: Flow Cyt, WB, IHC-P

Species reactivity Reacts with: Mouse, Human

Immunogen Recombinant fragment (Human) from near the N terminus.

Positive control HeLa whole cell lysate (<u>ab150035</u>); NIH3T3 whole cell lysate

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at 4°C (stable for up to 12 months). Store at -20°C or -80°C.

Storage buffer pH: 7.40

Preservative: 0.05% Sodium azide Constituents: 1% BSA, PBS

Purity Protein G purified

Purification notes This antibody was purified using protein G column chromatography from culture supernatant of

hybridoma cultured in a medium containing bovine IgG-depleted (approximately 95%) fetal bovine

serum.

ClonalityMonoclonalClone number3152C2aIsotypeIgG2b

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab54353 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Flow Cyt		Use 1µg for 10 ⁶ cells. ab170192 - Mouse monoclonal lgG2b, is suitable for use as an isotype control with this antibody.
WB	★★★★ (3)	1/50 - 1/100. Detects a band of approximately 90 kDa (predicted molecular weight: 110 kDa).
IHC-P		Use a concentration of 5 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Target

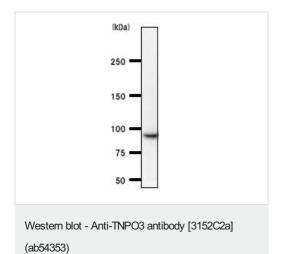
Function

Seems to function in nuclear protein import as nuclear transport receptor. In vitro, mediates the nuclear import of splicing factor SR proteins SFRS1 and SFRS2, by recognizing phosphorylated RS domains.

Cellular localization

Cytoplasm. Nucleus.

Images

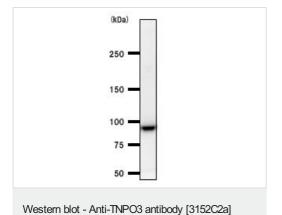


Anti-TNPO3 antibody [3152C2a] (ab54353) at 1/100 dilution + HeLa whole cell lysate at 25 μg

Secondary

anti Mouse IgG at 1/2500 dilution

Predicted band size: 110 kDa Observed band size: ~90 kDa



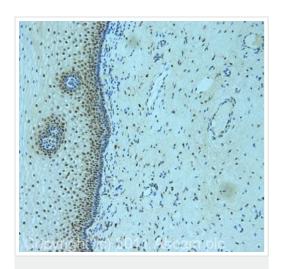
(ab54353)

Anti-TNPO3 antibody [3152C2a] (ab54353) at 1/50 dilution + NIH3T3 whole cell lysate at 25 μg

Secondary

anti Mouse IgG at 1/2500 dilution

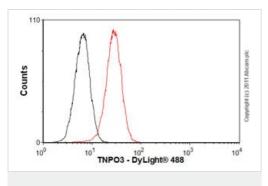
Predicted band size: 110 kDa Observed band size: ~90 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TNPO3 antibody
[3152C2a] (ab54353)

IHC image of ab54353 staining in human normal cervix formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pretreated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab54353, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



Flow Cytometry - Anti-TNPO3 antibody [3152C2a] (ab54353)

Overlay histogram showing HeLa cells stained with ab54353 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab54353, 1µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse lgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG2b [PLPV219] (ab91366, 2µg/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive signal in HeLa cells fixed with 80% methanol/permeabilized in 0.1% PBS-Tween used under the same conditions.

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