abcam

Product datasheet

Alexa Fluor® 647 Anti-Topoisomerase I antibody [EPR5375] ab196930

Recombinant

RabMAb

3 Images

Overview

Product name Alexa Fluor® 647 Anti-Topoisomerase I antibody [EPR5375]

Description Alexa Fluor® 647 Rabbit monoclonal [EPR5375] to Topoisomerase I

Host species Rabbit

Conjugation Alexa Fluor® 647. Ex: 652nm, Em: 668nm

Tested applications Suitable for: ICC/IF, Flow Cyt (Intra)

Species reactivity Reacts with: Human

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

Positive control ICC/IF: T47D cells. Flow Cyt (intra): T47D cells.

General notes 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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Properties

Form Liquid

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

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), PBS, 1% BSA

Purity Protein A purified

ClonalityMonoclonalClone numberEPR5375

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab196930 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
ICC/IF		1/100. This product gave a positive signal in T47D cells fixed with 4% formaldehyde (10 min) and 100% methanol (5 min)
Flow Cyt (Intra)		1/500. ab199093 - Rabbit monoclonal lgG (Alexa Fluor® 647), is suitable for use as an isotype control with this antibody.

Target

Function The reaction catalyzed by topoisomerases leads to the conversion of one topological isomer of

DNA to another.

Involvement in diseaseNote=A chromosomal aberration involving TOP1 is found in a form of therapy-related

myelodysplastic syndrome. Translocation t(11;20)(p15;q11) with NUP98.

Sequence similaritiesBelongs to the eukaryotic type I topoisomerase family.

Post-translational Sumoylated. Lys-117 is the main site of sumoylation. Sumoylation plays a role in partitioning

modifications

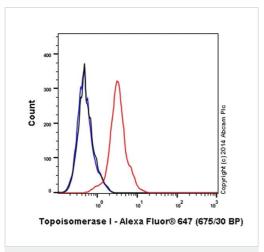
TOP1 between nucleoli and nucleoplasm. Levels are dramatically increased on camptothecin

(CPT) treatment.

Cellular localization Nucleus > nucleolus. Nucleus > nucleoplasm. Diffuse nuclear localization with some enrichment in

nucleoli. On CPT treatment, cleared from nucleoli into nucleoplasm. Sumolyated forms found in

both nucleoplasm and nucleoli.



Flow Cytometry (Intracellular) - Alexa Fluor® 647 Anti-Topoisomerase I antibody [EPR5375] (ab196930)

ab196930 ab195887

Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 647 Anti-Topoisomerase I antibody [EPR5375] (ab196930)

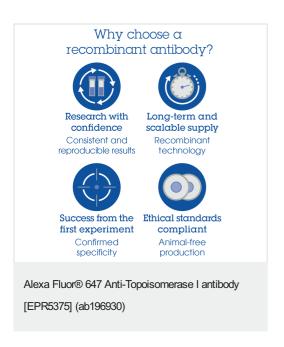
Overlay histogram showing T47D cells stained with ab196930 (red line). The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab196930, 1/500 dilution) for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) Alexa Fluor 647 used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a solid-state 25mW red diode laser (635 nm) and 675/30 bandpass filter.

This antibody gave a positive signal in T47D fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.

ab196930 staining Topoisomerase I in T47D cells. The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab196930 at a 1/100 dilution (shown in red) and ab195887, Mouse monoclonal to alpha Tubulin (Alexa Fluor[®] 488), at a 1/250 dilution (shown in green). Nuclear DNA was labelled with DAPI (shown in blue). Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

This product also gave a positive signal under the same testing conditions in HeLa cells fixed with 100% methanol (5 min).



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