abcam

Product datasheet

Anti-Survivin antibody [EPR2675] ab134170

אילשעבע RabMAb

★★★★★ <u>5 Abreviews</u> <u>13 References</u> 画像数 8

製品の概要

製品名	Anti-Survivin antibody [EPR2675]
製品の詳細	Rabbit monoclonal [EPR2675] to Survivin
由来種	Rabbit
アプリケーション	適用あり: Flow Cyt, WB, IHC-P, ICC/IF 適用なし: IP
種交差性	交差種: Mouse, Rat, Human
免疫原	Synthetic peptide within Human Survivin aa 50-150. The exact sequence is proprietary.
ポジティブ・コントロール	ICC/IF: NIH/3T3 cells; IHC-P: Human tonsil, Mouse and Rat spleen tissue; Flow Cyt: MCF7 cells. WB: HeLa, NIH/3T3, C6, Ramos and F9 whole cell lysates.
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 <u>see here</u>. Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb[®] patents</u>.

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 (glycerin, glycerine), 0.05% BSA
精製度	Protein A purified
ポリ/モノ	モノクローナル
クローン名	EPR2675

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ab134170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。

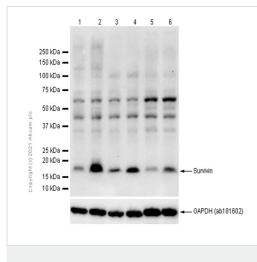
アプリケーション	Abreviews	特記事項
Flow Cyt		1/200.
WB	★ ★ ★ ★ ★ <u>(2)</u>	1/300 - 1/1000. Predicted molecular weight: 16 kDa.
IHC-P	★ ★ ★ ★ ☆ ☆ (1)	1/800. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
ICC/IF	* * * * * (1)	1/200.

追加情報

Is unsuitable for IP.

ターゲット情報

機能	Component of the chromosomal passenger complex (CPC), a complex that acts as a key regulator of mitosis. The CPC complex has essential functions at the centromere in ensuring correct chromosome alignment and segregation and is required for chromatin-induced microtubule stabilization and spindle assembly. The complex with RAN plays a role in mitotic spindle formation by serving as a physical scaffold to help deliver the RAN effector molecule TPX2 to microtubules. May play a role in neoplasia. May counteract a default induction of apoptosis in G2/M phase. Inhibitor of caspase-3 and caspase-7. Isoform 2 and isoform 3 do not appear to play vital roles in mitosis. Isoform 3 shows a marked reduction in its anti-apoptotic effects when compared with the displayed wild-type isoform.
組織特異性	Expressed only in fetal kidney and liver, and to lesser extent, lung and brain. Abundantly expressed in adenocarcinoma (lung, pancreas, colon, breast, and prostate) and in high-grade lymphomas. Also expressed in various renal cell carcinoma cell lines.
配列類似性	Belongs to the IAP family. Contains 1 BIR repeat.
発生段階	Expression is cell cycle-dependent and peaks at mitosis.
ドメイン	The BIR repeat is necessary and sufficient for HBXIP binding.
翻訳後修飾	Ubiquitination is required for centrosomal targeting. In vitro phosphorylation at Thr-117 by AURKB/STK12 prevents interaction with INCENP and localization to mitotic chromosomes.
細胞内局在	Cytoplasm. Nucleus. Chromosome. Chromosome > centromere. Cytoplasm > cytoskeleton > spindle. Localizes on chromosome arms and inner centromeres from prophase through metaphase and then transferring to the spindle midzone and midbody from anaphase through cytokinesis. Colocalizes with AURKB at mitotic chromosomes.



Western blot - Anti-Survivin antibody [EPR2675] (ab134170)

All lanes : Anti-Survivin antibody [EPR2675] (ab134170) at 1/1000 dilution (Purified)

Lane 1 : Untreated HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

Lane 2 : HeLa (Human cervix adenocarcinoma epithelial cell) treated with 100ng/ml nocodazole for 24 hours whole cell lysate Lane 3 : Untreated NIH/3T3 (Mouse embryonic fibroblast) whole cell lysate

Lane 4 : NIH/3T3 (Mouse embryonic fibroblast) treated with 100ng/ml nocodazole for 24 hours whole cell lysate
Lane 5 : Untreated C6 (Rat glial tumor glial cell) whole cell lysate
Lane 6 : C6 (Rat glial tumor glial cell) treated with 100ng/ml

Secondary

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

Predicted band size: 16 kDa

nocodazole for 24 hours whole cell lysate

All lanes : Anti-Survivin antibody [EPR2675] (ab134170) at 1/1000 dilution (Purified)

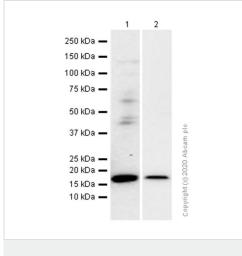
Lane 1 : Ramos (Human Burkitt's lymphoma B lymphocyte) whole cell lysate

Lane 2 : F9 (Mouse embryonal carcinoma epithelial cell) whole cell lysate

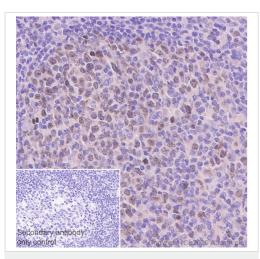
Secondary

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

Predicted band size: 16 kDa



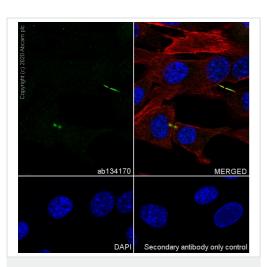
Western blot - Anti-Survivin antibody [EPR2675] (ab134170)



sections) analysis of human tonsil tissue sections labeling Survivin with purified ab134170 at 1/800 dilution (2.67 µg/mL). Heat mediated antigen retrieval was performed using Perform heat mediated antigen retrieval using **ab93684** (Tris/EDTA buffer, pH 9.0). Tissue was counterstained with Hematoxylin. Rabbit specific IHC polymer detection kit HRP/DAB (**ab209101**) secondary antibody was used at 1/0 dilution. PBS instead of the primary antibody was used as the negative control. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument.

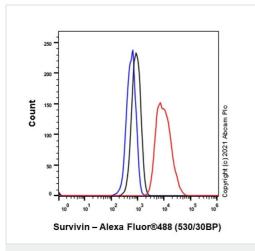
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Survivin antibody [EPR2675] (ab134170)



Immunocytochemistry/ Immunofluorescence - Anti-Survivin antibody [EPR2675] (ab134170)

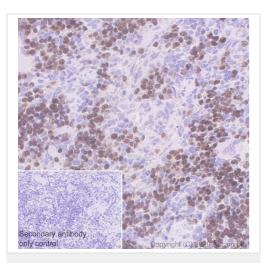
Immunocytochemistry analysis of NIH/3T3 (Mouse embryonic fibroblast) cells labeling Survivin with purified ab134170 at 1/200 dilution (10 μ g/mL). Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. Cells were counterstained with Ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594) 1/200 (2.5 μ g/mL). Goat anti rabbit IgG (Alexa Fluor[®] 488, **ab150077**) was used as the secondary antibody at 1/1000 (2 μ g/mL) dilution. DAPI (blue) was used as nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



1/200 dilution (10 μg/mL) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit lgG (Alexa Fluor® 488, **ab150077**) secondary antibody was used at 1/2000. lsotype control - Rabbit monoclonal lgG (Black). Unlabelled control - Cell without incubation with primary antibody and secondary antibody (Blue).

Flow Cytometry analysis of MCF7 (Human breast adenocarcinoma epithelial cell) cells labelling Survivin with purified ab134170 at

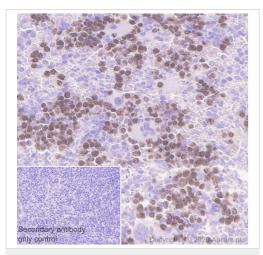
Flow Cytometry - Anti-Survivin antibody [EPR2675] (ab134170)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Survivin antibody [EPR2675] (ab134170)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat spleen tissue sections labeling Survivin with purified ab134170 at 1/800 dilution (2.67 µg/mL). Heat mediated antigen retrieval was performed using Perform heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0). Tissue was counterstained with Hematoxylin. Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>) secondary antibody was used at 1/0 dilution. PBS instead of the primary antibody was used as the negative control.

The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Survivin antibody [EPR2675] (ab134170)



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse spleen tissue sections labeling Survivin with purified ab134170 at 1/800 dilution (2.67 µg/mL). Heat mediated antigen retrieval was performed using Perform heat mediated antigen retrieval using **ab93684** (Tris/EDTA buffer, pH 9.0). Tissue was counterstained with Hematoxylin. Rabbit specific IHC polymer detection kit HRP/DAB (**ab209101**) secondary antibody was used at 1/0 dilution. PBS instead of the primary antibody was used as the negative control. The immunostaining was performed on a Leica Biosystems

BOND[®] RX instrument.

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