abcam

Product datasheet

Anti-SNAP23 antibody ab3340

★★★★★ 7 Abreviews 19 References 画像数 7

製品の概要

製品名 Anti-SNAP23 antibody

製品の詳細 Rabbit polyclonal to SNAP23

由来種 Rabbit

アプリケーション 適用あり: WB, ICC/IF, IP, IHC-P

種交差性 交差種: Mouse, Rat, Human, Recombinant fragment

免疫原 Synthetic peptide corresponding to Mouse SNAP23 aa 193-210.

Sequence:

NKNRIDIANTRAKKLIDS

(Peptide available as ab4956)

Run BLAST with
Run BLAST with

特記事項

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

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 or -

80°C. Avoid freeze / thaw cycle.

パッファー Constituents: 0.1% BSA, 99% PBS

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

1

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ab3340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	★ ★ ★ ★ 🚖 (6)	Use a concentration of 1 - 2 µg/ml. Detects a band of approximately 23 kDa. 23kDa band represents SNAP 23 from rat brain protein extract.
ICC/IF		Use a concentration of 2 µg/ml.
IP	★★★★☆ (1)	Use at an assay dependent concentration. Used at a concentration of 2 ug/ml for 1 hr (see Abreview).
IHC-P		Use a concentration of 1 µg/ml.

ターゲット情報

機能	Essential component of	the high affinity rec	ceptor for the general m	embrane fusion machinery and

an important regulator of transport vesicle docking and fusion.

組織特異性 Ubiquitous. Highest levels where found in placenta.

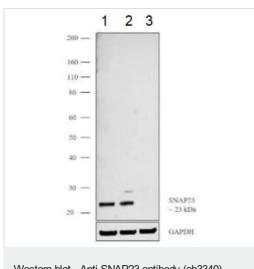
配列類似性 Belongs to the SNAP-25 family.

Contains 2 t-SNARE coiled-coil homology domains.

細胞内局在 Cell membrane. Cell membrane. Cell junction, synapse, synaptosome. Mainly localized to the

plasma membrane.

画像



Western blot - Anti-SNAP23 antibody (ab3340)

All lanes: Anti-SNAP23 antibody (ab3340) at 1 µg/ml

Lane 1: Untransfected U-87 MG (Human glioblastomaastrocytoma epithelial cell line) whole cell lysates

Lane 2: Non-specific scrambled siRNA transfected U-87 MG

whole cell lysates

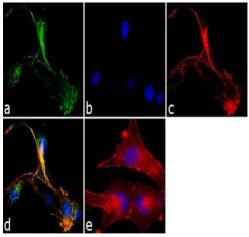
Lane 3: SNAP23 KO U-87 MG (Human glioblastoma-astrocytoma

epithelial cell line) whole cell lysates

Secondary

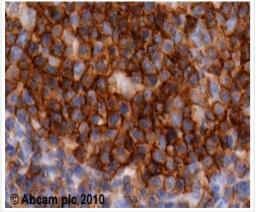
All lanes : Goat anti-Rabbit lgG (H+L) Superclonal™ Secondary Antibody, HRP conjugate at 0.25 µg/ml

Decrease in signal upon siRNA mediated knock down confirms that antibody is specific to SNAP23.



Immunocytochemistry/ Immunofluorescence - Anti-

SNAP23 antibody (ab3340) magnification.



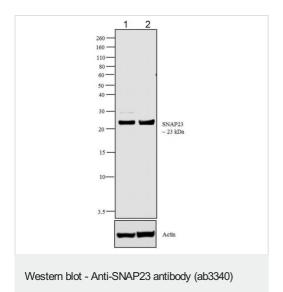
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SNAP23 antibody (ab3340)

90% confluent log phase U-87 MG (Human glioblastomaastrocytoma epithelial cell line) cells with ab3340. The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 10 minutes, and blocked with 1% BSA for 1 hour at room temperature. The cells were labeled with ab3340 at 2µg/mL in 0.1% BSA and incubated for 3 hours at room temperature and then labeled with Goat anti-Rabbit IgG (H+L) Superclonal™ Secondary Antibody, Alexa Fluor® 488 conjugate at a dilution of 1:2000 for 45 minutes at room temperature (Panel a: green). Nuclei (Panel b: blue) were stained with DAPI. F-actin (Panel c: red) was stained with Alexa Fluor® 555 Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing membrane localization. Panel e shows the no primary antibody control. The images were captured at 60X

Immunofluorescence analysis of SNAP-23 was performed using

ab3340 (1ug/ml) staining SNAP23 in human tonsil using an automated system (DAKO Autostainer Plus). Using this protocol there is strong staining of cellular membrane compartments of the lymphatic nodules.

Sections were rehydrated and antigen retrieved with the Dako 3 in 1 AR buffer EDTA pH 9.0 in a DAKO PT link. Slides were peroxidase blocked in 3% H2O2 in methanol for 10 mins. They were then blocked with Dako Protein block for 10 minutes (containing casein 0.25% in PBS) then incubated with primary antibody for 20 min and detected with Dako envision flex amplification kit for 30 minutes. Colorimetric detection was completed with Diaminobenzidine for 5 minutes. Slides were counterstained with Haematoxylin and coverslipped under DePeX. Please note that, for manual staining, optimization of primary antibody concentration and incubation time is recommended. Signal amplification may be required.



All lanes: Anti-SNAP23 antibody (ab3340) at 2 µg/ml

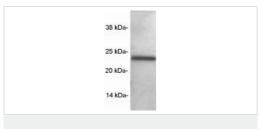
Lane 1 : U-87 MG (Human glioblastoma-astrocytoma epithelial cell line) whole cell lysate

Lane 2: HepG2 (Human liver hepatocellular carcinoma cell line) whole cell lysate

Lysates/proteins at 30 µg/ml per lane.

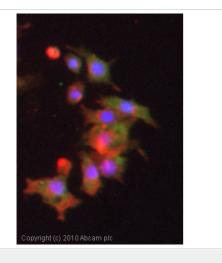
Secondary

All lanes : Goat anti-Rabbit lgG (H+L) Superclonal $^{\text{TM}}$ Secondary Antibody, HRP conjugate at 0.4 μ g/ml



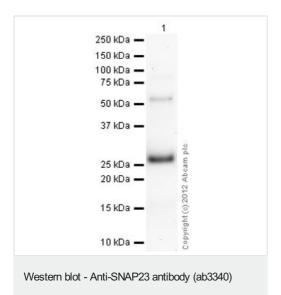
Western blot - Anti-SNAP23 antibody (ab3340)

Anti-SNAP23 antibody (ab3340) + Rat brain tissue



Immunocytochemistry/ Immunofluorescence - Anti-SNAP23 antibody (ab3340)

ICC/IF image of ab3340 stained PC-12 (Rat adrenal gland pheochromocytoma cell line) cells. The cells were 4% PFA fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab3340, 5 μ g/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat antirabbit lgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 μ M.



Anti-SNAP23 antibody (ab3340) at 1 μ g/ml + Recombinant Human SNAP23 protein (ab79180) at 0.001 μ g

Secondary

Goat Anti-Rabbit IgG H&L (HRP) preadsorbed (ab97080) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 2 minutes

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