# abcam

## Product datasheet

## Anti-SNX27 antibody [1C6] ab77799

★★★★★ 10 Abreviews 38 References 画像数 4

#### 製品の概要

製品名 Anti-SNX27 antibody [1C6]

製品の詳細 Mouse monoclonal [1C6] to SNX27

由来種 Mouse

アプリケーション 適用あり: Flow Cyt (Intra), WB, IP, ICC/IF

種交差性 交差種: Human

交差が予測される動物種: Monkey 4 非交差種: Mouse

免疫原 Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

エピトープ N-terminal amino acids 1-267.

ポジティブ・コントロール This antibody gave a positive signal in the following whole cell lysates: HeLa; A431; HEK293;

A549; MDA-MB-231. ICC/IF: Jurkat cells

特記事項 This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

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. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

**バッファー** pH: 7.40

Preservative: 0.02% Sodium azide Constituents: PBS, 6.97% L-Arginine

精製度 Protein G purified

ポリ/モノ モノクローナル

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クローン名	1C6
<b>₹</b> ID- <b>₹</b>	Sp2/0
アイソタイプ	lgG1

## アプリケーション

**The Abpromise guarantee** <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab77799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 (Intra)		Use $0.1\text{-}1\mu\text{g}$ for $10^6$ cells. $\underline{ab170190}$ - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.
WB	<b>★★★★</b> ★ (6)	Use a concentration of 10 µg/ml. Detects a band of approximately 61 kDa (predicted molecular weight: 61 kDa).  Abcam recommends using 3% milk as the blocking agent.
IP		Use at an assay dependent concentration.
ICC/IF	<b>★★★★☆</b> (4)	Use a concentration of 5 µg/ml.

## ターゲット情報

機能	Involved in endocytic trafficking (By similarity). In T lymphocytes, participates in endocytic recycling
	pathway. Recruits PSCDBP and HT4R to early endosomes.

組織特異性 Expressed in cells of hematopoietic origin (at protein level).

配列類似性
Belongs to the sorting nexin family.
Contains 1 PDZ (DHR) domain.
Contains 1 PX (phox homology) domain.
Contains 1 Ras-associating domain.

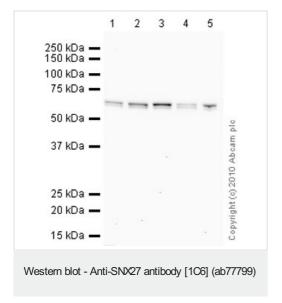
ドメイン The PDZ domain mediates the interaction with DGKZ, PSCDBP and HT4R and is responsible for

vesicular localization.

細胞内局在 Cytoplasm > cytosol. Early endosome. In T-lymphocytes, recruited from the cytosol to sorting

endosomes by phosphoinositide-3-kinase products.

### 画像



All lanes: Anti-SNX27 antibody [1C6] (ab77799) at 1 µg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2 : A431 (Human epithelial carcinoma cell line) Whole Cell Lysate

**Lane 3**: HEK293 (Human embryonic kidney cell line) Whole Cell Lysate

**Lane 4**: A549 (Human lung adenocarcinoma epithelial cell line) Whole Cell Lysate

**Lane 5**: MDA-MB-231 (Human breast adenocarcinoma cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

#### Secondary

**All lanes :** Goat polyclonal to Mouse IgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

Developed using the ECL technique.

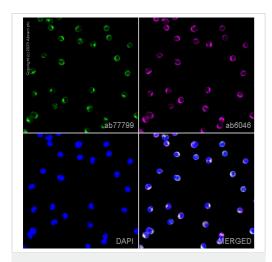
Performed under reducing conditions.

**Predicted band size:** 61 kDa **Observed band size:** 61 kDa

Exposure time: 15 minutes

Blocking: 5% Milk in TBS-T.

Two bands are observed very close together, at around 60-kDa. There are four reported isoforms of the SNX27 protein (Swissprot). We hypothesise that this antibody detects both isoform 1 and isoform 2, which are 61-kDa and 60-kDa respectively. As the immunogen is derived from the N-terminal region of SNX27, the antibody is not expected to react with isoforms 3 or 4.

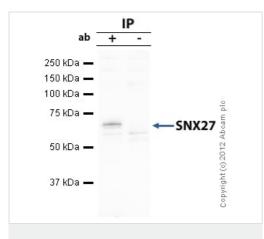


Immunocytochemistry/ Immunofluorescence - Anti-SNX27 antibody [1C6] (ab77799)

Flow Cytometry (Intracellular) - Anti-SNX27 antibody [1C6] (ab77799)

ab77799 staining SNX27 in Jurkat cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Tween 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 ab77799 at 5µg/ml and ab6046, Rabbit polyclonal to beta Tubulin - Loading Control. Cells were then incubated with ab150117, Goat polyclonal Secondary Antibody to Mouse IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (shown in green) and ab150080, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 594) at 1/1000 dilution (shown in pseudocolour magenta). Nuclear DNA was labelled with DAPI (shown in blue). Image was acquired with a confocal microscope (Leica-Microsystems TCS SP8) and a single confocal section is shown.

Overlay histogram showing Jurkat cells stained with ab77799 (red line). The cells were fixed with 80% methanol (5 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 (ab77799, 1µg/1x10<sup>6</sup> cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse IgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG1 [ICIGG1] (ab91353, 2µg/1x10<sup>6</sup> cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive signal in Jurkat cells fixed with 4% paraformaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



Immunoprecipitation - Anti-SNX27 antibody [1C6] (ab77799)

SNX27 was immunoprecipitated using 0.5mg Hek293 whole cell extract, 5µg of Mouse monoclonal to SNX27 and 50µl of protein G magnetic beads (+). No antibody was added to the control (-). The antibody was incubated under agitation with Protein G beads for 10min, Hek293 whole cell extract lysate diluted in RIPA buffer was added to each sample and incubated for a further 10min under agitation.

Proteins were eluted by addition of  $40\mu l$  SDS loading buffer and incubated for 10min at  $70^{\circ}C$ ;  $10\mu l$  of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with ab77799.

Secondary: Goat polyclonal to mouse IgG light chain specific (HRP) at 1/5000 dilution.

Band: 61kDa: SNX27.

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