# SNAP47 (D-11): sc-514428



The Power to Question

### **BACKGROUND**

In eukaryotic cells, the Golgi apparatus receives newly synthesized proteins from the endoplasmic reticulum and delivers them after covalent modification to their destination in the cell. For membrane-directed proteins, this process is believed to be carried out via vesicular transport. Correct vesicular transport is determined by specific pairing of vesicle-associated SNAREs (v-SNAREs) with those on the target membrane (t-SNAREs). This complex then recruits soluble NSF attachment proteins (SNAPs) and N-ethylmaleimide-sensitive factor (NSF) to form the highly stable SNAP receptor (SNARE) complex. The formation of a SNARE complex pulls the vesicle and target membrane together and may provide the energy to drive fusion of the lipid bilayers. SNAP47 (synaptosomal-associated protein 47), also known as epididymis luminal protein 170, is a 464 amino acid protein that is ubiquitously expressed with highest levels found in nervous tissue. There are four isoforms of SNAP47 that are produced as a result of alternative splicing events.

### **REFERENCES**

- 1. Holt, M., et al. 2006. Identification of SNAP-47, a novel Qbc-SNARE with ubiquitous expression. J. Biol. Chem. 281: 17076-17083.
- Leabu, M. 2006. Membrane fusion in cells: molecular machinery and mechanisms. J. Cell. Mol. Med. 10: 423-427.
- 3. Lang, T. and Jahn, R. 2008. Core proteins of the secretory machinery. Handb. Exp. Pharmacol. 184: 107-127.
- 4. Jena, B.P. 2008. Assembly and disassembly of SNAREs in membrane fusion. Methods Cell Biol. 90: 157-182.
- Stein, A., et al. 2009. Helical extension of the neuronal SNARE complex into the membrane. Nature 460: 525-528.
- Chen, S. and Barbieri, J.T. 2009. Engineering botulinum neurotoxin to extend therapeutic intervention. Proc. Natl. Acad. Sci. USA 106: 9180-9184.

#### **CHROMOSOMAL LOCATION**

Genetic locus: SNAP47 (human) mapping to 1q42.13.

### **SOURCE**

SNAP47 (D-11) is a mouse monoclonal antibody raised against amino acids 290-442 mapping near the C-terminus of SNAP47 of human origin.

#### **PRODUCT**

Each vial contains 200  $\mu g \ lgG_1$  kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

SNAP47 (D-11) is available conjugated to agarose (sc-514428 AC), 500  $\mu$ g/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-514428 HRP), 200  $\mu$ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-514428 PE), fluorescein (sc-514428 FITC), Alexa Fluor\* 488 (sc-514428 AF488), Alexa Fluor\* 546 (sc-514428 AF546), Alexa Fluor\* 594 (sc-514428 AF594) or Alexa Fluor\* 647 (sc-514428 AF647), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor\* 680 (sc-514428 AF680) or Alexa Fluor\* 790 (sc-514428 AF790), 200  $\mu$ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

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### **APPLICATIONS**

SNAP47 (D-11) is recommended for detection of SNAP47 of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for SNAP47 siRNA (h): sc-88350, SNAP47 shRNA Plasmid (h): sc-88350-SH and SNAP47 shRNA (h) Lentiviral Particles: sc-88350-V.

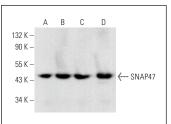
Molecular Weight of SNAP47 isoforms 1-4: 53/50/23/25 kDa.

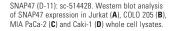
Positive Controls: Jurkat whole cell lysate: sc-2204, COLO 205 whole cell lysate: sc-364177 or Caki-1 cell lysate: sc-2224.

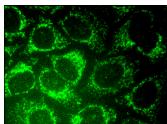
## **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG $\kappa$  BP-HRP: sc-516102 or m-lgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> Molecular Weight Standards: sc-2035, UltraCruz\* Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG $\kappa$  BP-FITC: sc-516140 or m-lgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz\* Mounting Medium: sc-24941 or UltraCruz\* Hard-set Mounting Medium: sc-359850.

#### DATA







SNAP47 (D-11): sc-514428. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization.

#### **SELECT PRODUCT CITATIONS**

 Jian, F., et al. 2025. Deacetylated SNAP47 recruits HOPS to facilitate autophagosome-lysosome fusion independent of STX17. Nat. Commun. 16: 543.

### **STORAGE**

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.