

Sorting Nexin 4

Cat.No. 392 003; Polyclonal rabbit antibody, 50 µg specific antibody (lyophilized)

Data Sheet

Reconstitution/ Storage	50 µg specific antibody, lyophilized. Affinity purified with the immunogen. Rabbit serum albumin was added for stabilization. For reconstitution add 50 µl H ₂ O to get a 1mg/ml solution in PBS. Then aliquot and store at -20°C until use.
Applications	WB: 1 : 1000 (AP staining) IP: not tested yet ICC: 1 : 500 IHC: not tested yet IHC-P/FFPE: not tested yet
Immunogen	Synthetic peptide corresponding to AA 1 to 21 from mouse SNX4 (UniProt Id: Q91YJ2)
Reactivity	Reacts with: rat (E9PU13), mouse (Q91YJ2). Other species not tested yet.
Specificity	Specific for SNX 4. (K.D. verified)
matching control	392-0P

TO BE USED IN VITRO / FOR RESEARCH ONLY
NOT TOXIC, NOT HAZARDOUS, NOT INFECTIOUS, NOT CONTAGIOUS

Sorting **Nexin 4 (SNX 4)** is involved in preventing the degradation of endocytosed transferrin receptors and their sorting into the Rab 11a recycling pathway. It may also form complexes with clathrin and dynein and binds several additional receptors for the epidermal growth factor, insulin and leptin.

Selected General References

Reggies/flotillins interact with Rab11a and SNX4 at the tubulovesicular recycling compartment and function in transferrin receptor and E-cadherin trafficking.

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SNX4 in complex with clathrin and dynein: implications for endosome movement.

Skänland SS, Wälchli S, Brech A, Sandvig K

PLoS one (2009) 4(6): e5935.

SNX4 coordinates endosomal sorting of TfnR with dynein-mediated transport into the endocytic recycling compartment.

Traer CJ, Rutherford AC, Palmer KJ, Wassmer T, Oakley J, Attar N, Carlton JG, Kremerskothen J, Stephens DJ, Cullen PJ
Nature cell biology (2007) 9(12): 1370-80.

A large family of endosome-localized proteins related to sorting nexin 1.

Teasdale RD, Loci D, Houghton F, Karlsson L, Gleeson PA

The Biochemical journal (2001) 358(Pt 1): 7-16.

Identification of a family of sorting nexin molecules and characterization of their association with receptors.

Haft CR, de la Luz Sierra M, Barr VA, Haft DH, Taylor SI

Molecular and cellular biology (1998) 18(12): 7278-87.