

**ATP6V1H Antibody (C-term)**  
**Affinity Purified Rabbit Polyclonal Antibody (Pab)**  
**Catalog # AP5182B**

**Specification**

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**ATP6V1H Antibody (C-term) - Product Information**

Application	WB,E
Primary Accession	<a href="#">O9UI12</a>
Other Accession	<a href="#">O9TVC1</a> , <a href="#">O8BVE3</a> , <a href="#">O46563</a>
Reactivity	Human, Mouse
Predicted	Bovine, Pig
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	55883
Antigen Region	400-426

**ATP6V1H Antibody (C-term) - Additional Information**

**Gene ID** 51606

**Other Names**

V-type proton ATPase subunit H, V-ATPase subunit H, Nef-binding protein 1, NBP1, Protein VMA13 homolog, V-ATPase 50/57 kDa subunits, Vacuolar proton pump subunit H, Vacuolar proton pump subunit SFD, ATP6V1H

**Target/Specificity**

This ATP6V1H antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 400-426 amino acids from the C-terminal region of human ATP6V1H.

**Dilution**

WB~~1:1000

E~~Use at an assay dependent concentration.

**Format**

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

**Storage**

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

ATP6V1H Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

**ATP6V1H Antibody (C-term) - Protein Information**

**Name** ATP6V1H

**Function** Subunit of the V1 complex of vacuolar(H<sup>+</sup>)-ATPase (V-ATPase), a multisubunit enzyme composed of a peripheral complex (V1) that hydrolyzes ATP and a membrane integral complex (V0) that translocates protons (PubMed:[33065002](#)). V-ATPase is responsible for acidifying and maintaining the pH of intracellular compartments and in some cell types, is targeted to the plasma membrane, where it is responsible for acidifying the extracellular environment (By similarity). Subunit H is essential for V-ATPase activity, but not for the assembly of the complex (By similarity). Involved in the endocytosis mediated by clathrin-coated pits, required for the formation of endosomes (PubMed:[12032142](#)).

**Cellular Location**

Cytoplasmic vesicle, clathrin-coated vesicle membrane {ECO:0000250|UniProtKB:O46563};  
Peripheral membrane protein

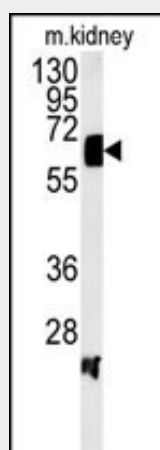
**Tissue Location**

Widely expressed..

**ATP6V1H Antibody (C-term) - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

**ATP6V1H Antibody (C-term) - Images**

Western blot analysis of ATP6V1H Antibody (C-term) (Cat. #AP5182b) in mouse kidney tissue lysates (35ug/lane).ATP6V1H (arrow) was detected using the purified Pab.

**ATP6V1H Antibody (C-term) - Background**

ATP6V1H encodes a component of vacuolar ATPase (V-ATPase), a multisubunit enzyme that mediates acidification of eukaryotic intracellular organelles. V-ATPase dependent organelle

acidification is necessary for such intracellular processes as protein sorting, zymogen activation, receptor-mediated endocytosis, and synaptic vesicle proton gradient generation. V-ATPase is composed of a cytosolic V1 domain and a transmembrane V0 domain. The V1 domain consists of three A and three B subunits, two G subunits plus the C, D, E, F, and H subunits. The V1 domain contains the ATP catalytic site. The V0 domain consists of five different subunits: a, c, c', c', and d. Additional isoforms of many of the V1 and V0 subunit proteins are encoded by multiple genes or alternatively spliced transcript variants. This gene encodes the regulatory H subunit of the V1 domain which is required for catalysis of ATP but not the assembly of V-ATPase.

**ATP6V1H Antibody (C-term) - References**

Fuster, D.G., et al. *Kidney Int.* 73(10):1151-1158(2008)  
Stove, V., et al. *J. Virol.* 79(17):11422-11433(2005)  
Morel, N. *Biol. Cell* 95(7):453-457(2003)