

V-ATPase B1 (C-19): sc-31466

BACKGROUND

Vacuolar-type H⁺-ATPase (V-ATPase) is a multisubunit enzyme responsible for acidification of eukaryotic intracellular organelles. V-ATPases pump protons against an electrochemical gradient, while F-ATPases reverse the process, thereby synthesizing ATP. A peripheral V₁ domain, which is responsible for ATP hydrolysis, and a integral V₀ domain, which is responsible for proton translocation, compose V-ATPase. Nine subunits (A–H) make up the V₁ domain and five subunits (a, d, c, c' and c'') make up the V₀ domain. Like F-ATPase, V-ATPase most likely operates through a rotary mechanism. The V-ATPase V₁ B subunit exists as two isoforms. In the inner ear, the V-ATPase B1 isoform functions in proton secretion and is required to maintain proper endolymph pH and normal auditory function. The gene encoding the human V-ATPase B1 isoform maps to chromosome 2q13.1. Mutations in this gene cause distal renal tubular acidosis associated with sensorineural deafness. The V-ATPase B2 isoform is expressed in kidney and is the only B isoform expressed in osteoclasts. The gene encoding the human V-ATPase B2 isoform maps to chromosome 8p22-p21.

REFERENCES

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- Lee, B.S., et al. 1996. Osteoclasts express the B2 isoform of vacuolar H⁺-ATPase intracellularly and on their plasma membranes. *Am. J. Physiol.* 270: 382-388.
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- Nishi, T., et al. 2002. The vacuolar H⁺-ATPases — nature's most versatile proton pumps. *Nat. Rev. Mol. Cell. Biol.* 3: 94-103.

CHROMOSOMAL LOCATION

Genetic locus: ATP6V1B1 (human) mapping to 2q13.1; Atp6v1b1 (mouse) mapping to 6 C3.

SOURCE

V-ATPase B1 (C-19) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of V-ATPase B1 of human origin.

STORAGE

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

PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-31466 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

V-ATPase B1 (C-19) is recommended for detection of V-ATPase B1 of human and, to a lesser extent, mouse and rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), 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); non cross-reactive with V-ATPase B2.

Suitable for use as control antibody for V-ATPase B1 siRNA (h): sc-36787, V-ATPase B1 shRNA Plasmid (h): sc-36787-SH and V-ATPase B1 shRNA (h) Lentiviral Particles: sc-36787-V.

Molecular Weight of V-ATPase B1: 56-58 kDa.

Positive Controls: mouse kidney extract: sc-2255 or rat kidney extract: sc-2394.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

RESEARCH USE

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

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.



Try **V-ATPase B1/2 (D-4): sc-271832** or **V-ATPase B1/2 (F-6): sc-55544**, our highly recommended monoclonal alternatives to V-ATPase B1 (C-19).