V-ATPase G1 (Q-20): sc-21224



The Power to Question

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 V $_1$ domain, which is responsible for ATP hydrolysis, and a integral V $_0$ domain, which is responsible for proton translocation, compose V-ATPase. Nine subunits (A-H) make up the V $_1$ domain and five subunits (a, d, c, c' and c") make up the V $_0$ domain. Like F-ATPase, V-ATPase most likely operates through a rotary mechanism. In yeast, the V-ATPase G subunit is a soluble subunit that shares homology with the F-ATPase G subunit and may be part of a connection stalk between V $_1$ and V $_0$. The G $_2$ isoform of the G subunit associates with the pore-forming a1C-subunit of L-type calcium channel and aids in proper membrane targeting of the calcium channel. The genes encoding the G $_1$ and G $_2$ V-ATPase subunits map to chromosomes 9q32 and 6p21.3, respectively.

REFERENCES

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- Neville, M.J., et al. 1999. A new member of the lg superfamily and a V-ATPase G subunit are among the predicted products of novel genes close to the TNF locus in the human MHC. J. Immunol. 162: 4745-4754.
- 3. Gao, T., et al. 2000. Association of L-type calcium channels with a vacuolar H+-ATPase $\rm G_2$ subunit. Biochem. Biophys. Res. Commun. 277: 611-616.
- 4. Nishi, T., et al. 2002. The vacuolar H+-ATPases—nature's most versatile proton pumps. Nat. Rev. Mol. Cell. Biol. 3: 94-103.
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CHROMOSOMAL LOCATION

Genetic locus: ATP6V1G1 (human) mapping to 9q32; Atp6v1g1 (mouse) mapping to 4 C1.

SOURCE

V-ATPase G1 (Q-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of V-ATPase G1 of human origin.

PRODUCT

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

Blocking peptide available for competition studies, sc-21224 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

STORAGE

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

APPLICATIONS

V-ATPase G1 (Q-20) is recommended for detection of V-ATPase G1 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ 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).

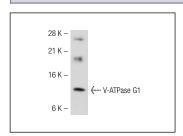
V-ATPase G1 (Q-20) is also recommended for detection of V-ATPase G1 in additional species, including canine, bovine, porcine and avian.

Suitable for use as control antibody for V-ATPase G1 siRNA (h): sc-36797, V-ATPase G1 siRNA (m): sc-36798, V-ATPase G1 shRNA Plasmid (h): sc-36797-SH, V-ATPase G1 shRNA Plasmid (m): sc-36798-SH, V-ATPase G1 shRNA (h) Lentiviral Particles: sc-36797-V and V-ATPase G1 shRNA (m) Lentiviral Particles: sc-36798-V.

Molecular Weight of V-ATPase G1: 13 kDa.

Positive Controls: rat kidney extract: sc-2394, MIA PaCa-2 cell lysate: sc-2285 or rat pancreas extract: sc-364806.

DATA



V-ATPase G1 (Q-20): sc-21224. Western blot analysis of V-ATPase G1 expression in rat kidney tissue extract.

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 G1 (D-5):** sc-25333, our highly recommended monoclonal alternative to V-ATPase G1 (0-20).

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