

# Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 (E-4): sc-376406

## BACKGROUND

The ubiquitously expressed sodium/potassium-ATPase (Na<sup>+</sup>/K<sup>+</sup>-ATPase) exists as a oligomeric plasma membrane complex that couples the hydrolysis of one molecule of ATP to the importation of three Na<sup>+</sup> ions and two K<sup>+</sup> ions against their respective electrochemical gradients. As a member of the P-type family of ion motives, Na<sup>+</sup>/K<sup>+</sup>-ATPase plays a critical role in maintaining cellular volume, resting membrane potential and Na<sup>+</sup>-coupled solute transport. Multiple isoforms of three subunits, α, β and γ, comprise the sNa<sup>+</sup>/K<sup>+</sup>-ATPase oligomer. The α subunit contains the binding sites for ATP and the cations; the glycosylated β subunit ensures correct folding and membrane insertion of the α subunits. The small γ subunit co-localizes with the α subunit in nephron segments, where it increases the affinity of Na<sup>+</sup>/K<sup>+</sup>-ATPase for ATP. The β subunit, but not the γ subunit, is essential for normal activity of Na<sup>+</sup>/K<sup>+</sup>-ATPase.

## REFERENCES

- Hardwicke, P.M., et al. 1981. A proteolipid associated with Na,K-ATPase is not essential for ATPase activity. *Biochem. Biophys. Res. Commun.* 102: 250-257.
- Ackermann, U., et al. 1990. Mutual dependence of Na,K-ATPase α- and β-subunits for correct post-translational processing and intracellular transport. *FEBS Lett.* 269: 105-108.
- McDonough, A.A., et al. 1990. The sodium pump needs its β subunit. *FASEB J.* 4: 1598-1605.
- Pedemonte, C.H., et al. 1990. Chemical modification as an approach to elucidation of sodium pump structure-function relations. *Am. J. Physiol.* 258: C1-C23.

## CHROMOSOMAL LOCATION

Genetic locus: ATP1B1 (human) mapping to 1q24.2.

## SOURCE

Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 (E-4) is a mouse monoclonal antibody raised against amino acids 41-155 of Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 of human origin.

## PRODUCT

Each vial contains 200 μg IgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 (E-4) is available conjugated to agarose (sc-376406 AC), 500 μg/0.25 ml agarose in 1 ml, for IP; to HRP (sc-376406 HRP), 200 μg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-376406 PE), fluorescein (sc-376406 FITC), Alexa Fluor® 488 (sc-376406 AF488), Alexa Fluor® 546 (sc-376406 AF546), Alexa Fluor® 594 (sc-376406 AF594) or Alexa Fluor® 647 (sc-376406 AF647), 200 μg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-376406 AF680) or Alexa Fluor® 790 (sc-376406 AF790), 200 μg/ml, for Near-Infrared (NIR) WB, IF and FCM.

## STORAGE

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

## APPLICATIONS

Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 (E-4) is recommended for detection of Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μg per 100-500 μg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (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 Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 siRNA (h): sc-36008, Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 shRNA Plasmid (h): sc-36008-SH and Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 shRNA (h) Lentiviral Particles: sc-36008-V.

Molecular Weight of Na<sup>+</sup>/K<sup>+</sup>-ATPase β1: 40-60 kDa.

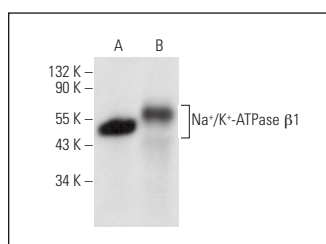
Positive Controls: human kidney extract: sc-363764, Caki-1 cell lysate: sc-2224 or human brain extract: sc-364375.

## RECOMMENDED SUPPORT REAGENTS

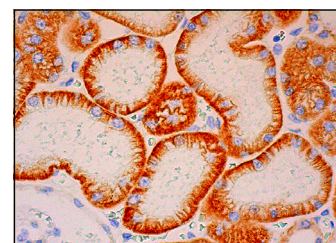
To ensure optimal results, the following support reagents are recommended:

- 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ 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-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.
- 4) Immunohistochemistry: use m-IgGκ BP-HRP: sc-516102 with DAB, 50X: sc-24982 and Immunohistomount: sc-45086, or Organo/Limonene Mount: sc-45087.

## DATA



Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 (E-4): sc-376406. Western blot analysis of Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 expression in human brain (A) and human kidney (B) tissue extracts.



Na<sup>+</sup>/K<sup>+</sup>-ATPase β1 (E-4): sc-376406. Immunoperoxidase staining of formalin fixed, paraffin-embedded human kidney tissue showing basolateral membrane staining of cells in tubules.

## SELECT PRODUCT CITATIONS

1. Ferencic, A., et al. 2020. Left ventricular hypertrophy is associated with overexpression of HSP60, TLR2, and TLR4 in the myocardium. *Scand. J. Clin. Lab. Invest.* 80: 236-246.

## RESEARCH USE

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

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