

# ATPIF1 (A-3): sc-271614

## BACKGROUND

Mitochondrial ATP synthases (ATPases) transduce the energy contained in membrane electrochemical proton gradients into the energy required for synthesis of high-energy phosphate bonds. ATPases contain two linked complexes:  $F_1$ , the hydrophilic catalytic core; and  $F_0$ , the membrane-embedded protein channel.  $F_1$  consists of three  $\alpha$  chains and three  $\beta$  chains, which are weakly homologous, as well as one  $\gamma$  chain, one  $\delta$  chain and one  $\epsilon$  chain.  $F_0$  consists of three subunits: a, b and c. A mitochondrial  $F_1$ -ATPase inhibitor protein, ATPIF1 (ATPase inhibitory factor 1), also known as IP, IF<sub>1</sub>, ATPi or ATPIP (ATPase inhibitor protein), binds to the C-terminal region of a  $\beta$  subunit of the  $F_1$ -ATPase at low pH values and, via interference of the  $\beta$  and  $\gamma$  subunit interaction, ATPIF1 regulates the activity of the  $F_1F_0$ -ATPase. This reversible ATPIF1 binding to  $F_1F_0$ -ATPase also occurs on the surface of endothelial cells.

## REFERENCES

1. Ichikawa, N., et al. 1999. Nucleotide sequence of cDNA coding the mitochondrial precursor protein of the ATPase inhibitor from humans. *Biosci. Biotechnol. Biochem.* 63: 2225-2227.
2. Cabezón, E., et al. 2001. The structure of bovine IF<sub>1</sub>, the regulatory subunit of mitochondrial F-ATPase. *EMBO J.* 20: 6990-6996.
3. Contessi, S., et al. 2005. Identification of a conserved calmodulin-binding motif in the sequence of  $F_1F_0$  ATPsynthase inhibitor protein. *J. Bioenerg. Biomembr.* 37: 317-326.
4. Cortés-Hernández, P., et al. 2005. The inhibitor protein of the  $F_1F_0$ -ATP synthase is associated to the external surface of endothelial cells. *Biochem. Biophys. Res. Commun.* 330: 844-849.

## CHROMOSOMAL LOCATION

Genetic locus: ATPIF1 (human) mapping to 1p35.3.

## SOURCE

ATPIF1 (A-3) is a mouse monoclonal antibody raised against amino acids 1-106 representing full length ATPIF1 of human origin.

## PRODUCT

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

ATPIF1 (A-3) is available conjugated to agarose (sc-271614 AC), 500  $\mu$ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-271614 HRP), 200  $\mu$ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-271614 PE), fluorescein (sc-271614 FITC), Alexa Fluor® 488 (sc-271614 AF488), Alexa Fluor® 546 (sc-271614 AF546), Alexa Fluor® 594 (sc-271614 AF594) or Alexa Fluor® 647 (sc-271614 AF647), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-271614 AF680) or Alexa Fluor® 790 (sc-271614 AF790), 200  $\mu$ g/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

ATPIF1 (A-3) is recommended for detection of ATPIF1 of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g 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 ATPIF1 siRNA (h): sc-78711, ATPIF1 shRNA Plasmid (h): sc-78711-SH and ATPIF1 shRNA (h) Lentiviral Particles: sc-78711-V.

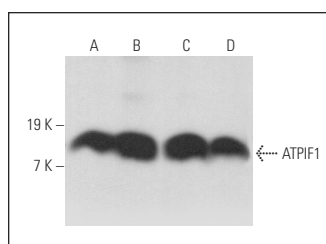
Molecular Weight of ATPIF1: 12 kDa.

Positive Controls: MCF7 whole cell lysate: sc-2206, HEK293T whole cell lysate: sc-45137 or U266 whole cell lysate: sc-364800.

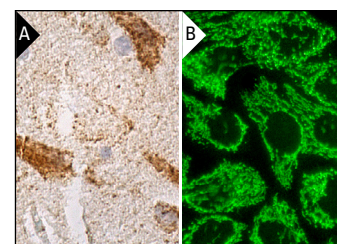
## RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  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-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  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-IgG $\kappa$  BP-HRP: sc-516102 with DAB, 50X: sc-24982 and Immunohistomount: sc-45086, or Organo/Limonene Mount: sc-45087.

## DATA



ATPIF1 (A-3): sc-271614. Western blot analysis of ATPIF1 expression in MCF7 (A), HEK293T (B), U266 (C) and HEK293 (D) whole cell lysates.



ATPIF1 (A-3): sc-271614. Immunoperoxidase staining of formalin fixed, paraffin-embedded human brain tissue showing cytoplasmic staining of neuronal cells (A). Immunofluorescence staining of formalin-fixed A-431 cells showing mitochondrial localization (B).

## SELECT PRODUCT CITATIONS

1. Elshaarawy, R.F.M., et al. 2020. Role of Pd(II)-chitooligosaccharides-Gboxin analog in oxidative phosphorylation inhibition and energy depletion: targeting mitochondrial dynamics. *Chem. Biol. Drug Des.* 96: 1148-1161.

## RESEARCH USE

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

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