

ATP5A (PT0255R) PT® Rabbit mAb

Catalog No: YM8161

Reactivity: Human; Mouse; Rat;

Applications: WB;IHC;IF;IP;ELISA

Target: ATP5A

Fields: >>Oxidative phosphorylation;>>Metabolic

pathways;>>Thermogenesis;>>Alzheimer disease;>>Parkinson

disease;>>Amyotrophic lateral sclerosis;>>Huntington disease;>>Prion disease;>>Pathways of neurodegeneration - multiple diseases;>>Chemical carcinogenesis - reactive oxygen species;>>Diabetic cardiomyopathy

Gene Name: ATP5A1

Protein Name: ATP synthase subunit alpha mitochondrial

P25705

Q03265

Human Gene Id: 498

Human Swiss Prot

No:

Mouse Gene Id: 11946

Mouse Swiss Prot

No:

Rat Gene Id: 65262

Rat Swiss Prot No: P15999

Specificity: endogenous

Formulation: PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA

Source : Monoclonal, rabbit, IgG, Kappa

Dilution: IHC 1:200-1:1000;WB 1:2000-1:10000;IF 1:200-1:1000;ELISA

1:5000-1:20000;IP 1:50-1:200;



Purification: Protein A

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight: 60kD

Observed Band: 55kD

Cell Pathway: Oxidative phosphorylation; Alzheimer's disease; Parkinson's disease; Huntington's

disease;

Background: This gene encodes a subunit of mitochondrial ATP synthase. Mitochondrial ATP

synthase catalyzes ATP synthesis, using an electrochemical gradient of protons across the inner membrane during oxidative phosphorylation. ATP synthase is composed of two linked multi-subunit complexes: the soluble catalytic core, F1, and the membrane-spanning component, Fo, comprising the proton channel. The catalytic portion of mitochondrial ATP synthase consists of 5 different subunits (alpha, beta, gamma, delta, and epsilon) assembled with a stoichiometry of 3 alpha, 3 beta, and a single representative of the other 3. The proton channel consists of three main subunits (a, b, c). This gene encodes the alpha subunit of the catalytic core. Alternatively spliced transcript variants encoding the different

isoforms have been identified. Pseudogenes of thi

Function: function:Mitochondrial membrane ATP synthase (F(1)F(0) ATP synthase or

Complex V) produces ATP from ADP in the presence of a proton gradient across the membrane which is generated by electron transport complexes of the respiratory chain. F-type ATPases consist of two structural domains, F(1) - containing the extramembraneous catalytic core, and F(0) - containing the membrane proton channel, linked together by a central stalk and a peripheral stalk. During catalysis, ATP synthesis in the catalytic domain of F(1) is coupled via a rotary mechanism of the central stalk subunits to proton translocation. Subunits alpha and beta form the catalytic core in F(1). Rotation of the central stalk against the surrounding alpha(3)beta(3) subunits leads to hydrolysis of ATP in three separate catalytic sites on the beta subunits. Subunit alpha does not bear

the catalytic high-affinity ATP-binding sites.,

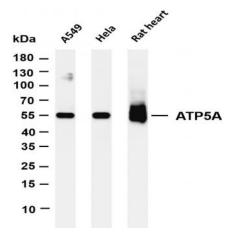
Subcellular Location:

Mitochondrion

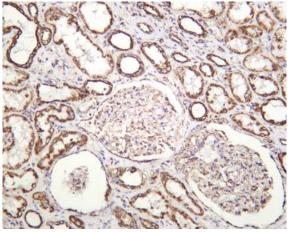
Expression: Fetal lung, heart, liver, gut and kidney. Expressed at higher levels in the fetal

brain, retina and spinal cord.

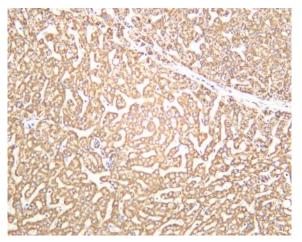
Products Images



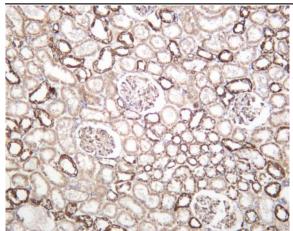
Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-ATP5A (PT0255R) antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: A549 Lane 2: Hela Lane 3: Rat heart Predicted band size: 60kDa Observed band size: 55kDa



Human kidney was stained with anti-ATP5A (PT0255R) rabbit antibody



Human liver was stained with anti-ATP5A (PT0255R) rabbit antibody



Rat kidney was stained with anti-ATP5A (PT0255R) rabbit antibody