

AT132 rabbit pAb

Catalog No: YT7465

Reactivity: Human; Mouse

Applications: WB

Target: AT132

Gene Name: ATP13A2 PARK9

Q9NQ11

Q9CTG6

Protein Name: AT132

Human Gene Id: 23400

Human Swiss Prot

No:

Mouse Gene Id: 74772

Mouse Swiss Prot

No:

Immunogen: Synthesized peptide derived from human AT132 AA range: 56-106

Specificity: This antibody detects endogenous levels of AT132 at Human/Mouse

Formulation: Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1 ?500-2000

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

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

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Molecularweight: 130kD

Background: This gene encodes a member of the P5 subfamily of ATPases which transports

inorganic cations as well as other substrates. Mutations in this gene are associated with Kufor-Rakeb syndrome (KRS), also referred to as Parkinson disease 9. Multiple transcript variants encoding different isoforms have been

found for this gene.[provided by RefSeq, Nov 2008],

Function: catalytic activity:ATP + H(2)O = ADP + phosphate., disease:Defects in ATP13A2

are the cause of Kufor-Rakeb syndrome (KRS) [MIM:606693]; also known as Parkinson disease-9. KRS is a rare hereditary disease with juvenile onset. In addition to typical signs of Parkinson disease, affected individuals show

symptoms of more widespread neurodegeneration, including

dementia., similarity: Belongs to the cation transport ATPase (P-type)

family., similarity: Belongs to the cation transport ATPase (P-type) family. Type V

subfamily.,

Subcellular Location:

Lysosome membrane; Multi-pass membrane protein. Late endosome membrane; Multi-pass membrane protein. Endosome, multivesicular body membrane; Multi-pass membrane protein. Cytoplasmic vesicle, autophagosome

membrane : Multi-pass membrane protein .

Expression: Expressed in brain; protein levels are markedly increased in brain from subjects

with Parkinson disease and subjects with dementia with Lewy bodies. Detected in pyramidal neurons located throughout the cingulate cortex (at protein level). In the substantia nigra, it is found in neuromelanin-positive dopaminergic neurons (at

protein level).

Sort: 2338

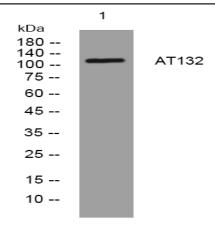
No4: 1

Host: Rabbit

Modifications: Unmodified

Products Images

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Western blot analysis of lysates from KB cells, primary antibody was diluted at 1:1000, 4° over night