

### ATP13A2 Antibody (monoclonal) (M06)

Mouse monoclonal antibody raised against a partial recombinant ATP13A2. Catalog # AT1230a

## **Specification**

## ATP13A2 Antibody (monoclonal) (M06) - Product Information

**Application** WB, E **Primary Accession 09N011** Other Accession NM 022089 Reactivity Human Host mouse Clonality **Monoclonal** Isotype IgG2a Kappa Calculated MW 128794

## ATP13A2 Antibody (monoclonal) (M06) - Additional Information

Gene ID 23400

#### **Other Names**

Probable cation-transporting ATPase 13A2, 363-, ATP13A2, PARK9

## Target/Specificity

ATP13A2 (NP\_071372, 68 a.a.  $\sim$  154 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

#### **Dilution**

WB~~1:500~1000

#### **Format**

Clear, colorless solution in phosphate buffered saline, pH 7.2.

## Storage

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

### **Precautions**

ATP13A2 Antibody (monoclonal) (M06) is for research use only and not for use in diagnostic or therapeutic procedures.

# ATP13A2 Antibody (monoclonal) (M06) - Protocols

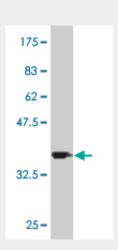
Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry

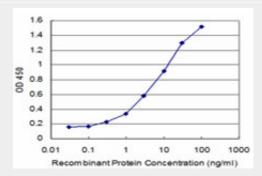


- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

## ATP13A2 Antibody (monoclonal) (M06) - Images



Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (35.31 KDa).



Detection limit for recombinant GST tagged ATP13A2 is approximately 0.1ng/ml as a capture antibody.

## ATP13A2 Antibody (monoclonal) (M06) - 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.

## ATP13A2 Antibody (monoclonal) (M06) - References

Mutational analysis of GIGYF2; ATP13A2 and GBA genes in Brazilian patients with early-onset Parkinson's disease. Dos Santos AV, et al. Neurosci Lett, 2010 Sep 14. PMID 20816920.Structural imaging in the presymptomatic stage of genetically determined parkinsonism. Reetz K, et al. Neurobiol Dis, 2010 Sep. PMID 20483373.Lack of association between ATP13A2 Ala746Thr variant and Parkinson's disease in Han population of mainland China. Fei QZ, et al. Neurosci Lett, 2010 May 14. PMID 20227461.A genome-wide association study in 19 633 Japanese subjects identified LHX3-QSOX2 and IGF1 as adult height loci. Okada Y, et al. Hum Mol Genet, 2010 Jun 1. PMID 20189936.A new variant of the ATP13A2 gene in Chinese patients with early-onset parkinsonism. Wang L, et al. Chin Med J (Engl), 2009 Dec 20. PMID 20137506.