# **Product Name: Aladin Rabbit Polyclonal Antibody**

Catalog #: APRab06753



#### **Summary**

**Production Name** Aladin Rabbit Polyclonal Antibody

**Description** Rabbit Polyclonal Antibody

HostRabbitApplicationWB,ELISAReactivityHuman,Rat

#### **Performance**

ConjugationUnconjugatedModificationUnmodified

**Isotype** IgG

Clonality Polyclonal Form Liquid

**Storage** Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% New type

preservative N.

**Purification** Affinity purification

#### **Immunogen**

Buffer

Gene Name AAAS

Alternative Names AAAS; ADRACALA; GL003; Aladin; Adracalin

**Gene ID** 8086.0

SwissProt ID Q9NRG9. Synthesized peptide derived from Aladin . at AA range: 360-440

## **Application**

**Dilution Ratio** WB 1:500-1:2000, ELISA 1:40000.Not yet tested in other applications.

Molecular Weight 59kDa

### **Background**

The protein encoded by this gene is a member of the WD-repeat family of regulatory proteins and may be involved in

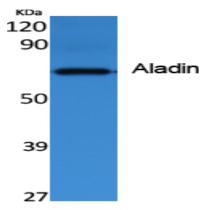
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normal development of the peripheral and central nervous system. The encoded protein is part of the nuclear pore complex and is anchored there by NDC1. Defects in this gene are a cause of achalasia-addisonianism-alacrima syndrome (AAAS), also called triple-A syndrome or Allgrove syndrome. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Mar 2010], disease:Defects in AAAS are the cause of achalasia-addisonianism-alacrima syndrome (AAAS) [MIM:231550]; also known as triple-A syndrome or Allgrove syndrome. AAAS is an autosomal recessive disorder characterized by adreno-corticotropic hormone (ACTH)-resistant adrenal failure, achalasia of the esophageal cardia and alacrima. The syndrome is associated with variable and progressive neurological impairment involving the central, peripheral, and autonomic nervous system. Other features such as palmoplantar hyperkeratosis, short stature, facial dysmorphy and osteoporosis may also be present.,function:Plays a role in the normal development of the peripheral and central nervous system.,similarity:Contains 4 WD repeats.,tissue specificity:Widely expressed. Particularly abundant expression is found in cerebellum, corpus callosum, adrenal gland, pituary gland, gatsrointestinal structures and fetal lung.,

#### **Research Area**

#### **Image Data**



Western Blot analysis of extracts from rat kidney, using Aladin Polyclonal Antibody.. Secondary antibody was diluted at 1:20000 cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003,Inventbiotech,MN,USA).

#### Note

For research use only.