## **ATXN1 Rabbit Polyclonal Antibody**



## **CAB6217**

**Product Information** 

Size:

20uL, 50uL, 100uL, 200uL

**Observed MW:** 

87kDa

Calculated MW:

86kDa

**Applications:** 

WB IF

Reactivity:

Human, Mouse, Rat

**Antibody Information** 

**Recommended dilutions:** WB 1:500 - 1:2000 IF 1:10 -

1:100

Source:

Rabbit

Isotype:

lgG

to amino acids 586-815 of human ATXN1 (NP\_001121636.1).

**Purification:** 

Affinity purification

**Protein Background** 

The autosomal dominant cerebellar ataxias (ADCA) are a heterogeneous group of neurodegenerative disorders characterized by progressive degeneration of the cerebellum, brain stem and spinal cord. Clinically, ADCA has been divided into three groups: ADCA types I-III. ADCAI is genetically heterogeneous, with five genetic loci, designated spinocerebellar ataxia (SCA) 1, 2, 3, 4 and 6, being assigned to five different chromosomes. ADCAII, which always presents with retinal degeneration (SCA7), and ADCAIII often referred to as the 'pure' cerebellar syndrome (SCA5), are most likely homogeneous disorders. Several SCA genes have been cloned and shown to contain CAG repeats in their coding regions. ADCA is caused by the expansion of the CAG repeats, producing an elongated polyglutamine tract in the corresponding protein. The expanded repeats are variable in size and unstable, usually increasing in size when transmitted to successive generations. The function of the ataxins is not known. This locus has been mapped to chromosome 6, and it has been determined that the diseased allele contains 40-83 CAG repeats, compared to 6-39 in the normal allele, and is associated with spinocerebellar ataxia type 1 (SCA1). At least two transcript variants encoding the same protein have been found for this gene.

Immunogen information

Gene ID:

6310

Uniprot P54253

Synonyms:

Immunogen:

Recombinant fusion protein containing a sequence corresponding

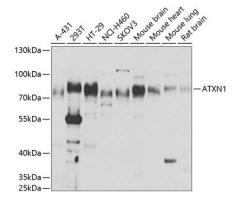
Storage:

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02%

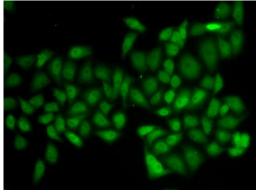
sodium azide, 50% glycerol, pH7.3.

ATXN1; ATX1; D6S504E; SCA1; ataxin-1

## **Product Images**



Western blot analysis of extracts of various cell lines, using ATXN1 antibody (CAB6217) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (CABS014) at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (CABM00020). Exposure time: 120s.



Immunofluorescence analysis of HeLa cells using ATXN1 antibody (CAB6217).