

## Ataxin-1 (phospho Ser776) Polyclonal Antibody

Catalog No: YP0536

**Reactivity:** Human; Mouse

**Applications:** WB;IHC;IF;ELISA

Target: Ataxin-1

**Fields:** >>Notch signaling pathway;>>Spinocerebellar ataxia;>>Pathways of

neurodegeneration - multiple diseases

Gene Name: ATXN1

Protein Name: Ataxin-1

Human Gene Id: 6310

**Human Swiss Prot** 

No:

Mouse Gene Id: 20238

**Mouse Swiss Prot** 

No:

Immunogen: The antiserum was produced against synthesized peptide derived from human

Ataxin 1 around the phosphorylation site of Ser776. AA range:742-791

Specificity: Phospho-Ataxin-1 (S776) Polyclonal Antibody detects endogenous levels of

Ataxin-1 protein only when phosphorylated at S776.

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

Source: Polyclonal, Rabbit, IgG

P54253

P54254

**Dilution:** WB 1:500 - 1:2000. IHC 1:100 - 1:300. IF 1:200 - 1:1000. ELISA: 1:10000. Not

yet tested in other applications.

**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)

Observed Band: 87kD

**Background:** ataxin 1(ATXN1) Homo sapiens 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

Function: alternative products:At least 2 isoforms are produced, disease:Defects in ATXN1

are the cause of spinocerebellar ataxia type 1 (SCA1) [MIM:164400]; also known as olivopontocerebellar atrophy I (OPCA I or OPCA1). Spinocerebellar ataxia is a clinically and genetically heterogeneous group of cerebellar disorders. Patients show progressive incoordination of gait and often poor coordination of hands, speech and eye movements, due to cerebellum degeneration with variable involvement of the brainstem and spinal cord. SCA1 belongs to the autosomal dominant cerebellar ataxias type I (ADCA I) which are characterized by cerebellar ataxia in combination with additional clinical features like optic atrophy,

ophthalmoplegia, bulbar and extrapyramidal signs, peripheral neuropathy and dementia. SCA1 is caused by expansion of a CAG repeat in the coding region of

ATXN1. Longer expansions result in earlier

Subcellular Location:

Cytoplasm . Nucleus . Colocalizes with USP7 in the nucleus. .

**Expression:** Widely expressed throughout the body.

Tag: orthogonal

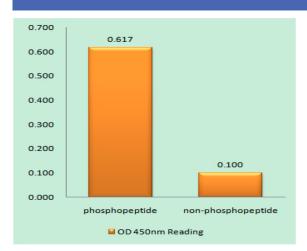
**Sort :** 2341

**No4:** 1

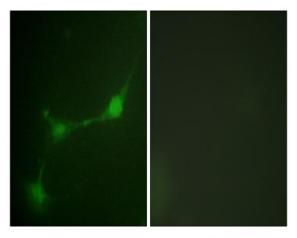
Host: Rabbit

Modifications : Phospho

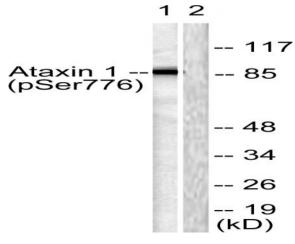
## **Products Images**



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using Ataxin 1 (Phospho-Ser776) Antibody

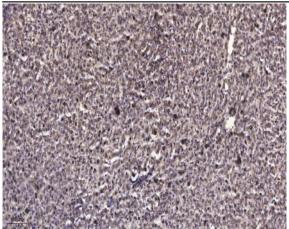


Immunofluorescence analysis of NIH/3T3 cells, using Ataxin 1 (Phospho-Ser776) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from HepG2 cells treated with Adriamycin 0.5uM 5h, using Ataxin 1 (Phospho-Ser776) Antibody. The lane on the right is blocked with the phospho peptide.





Immunohistochemical analysis of paraffin-embedded human liver cancer. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).