

## Datasheet for ABIN682819

## anti-Ataxin 1 antibody (pSer775) (Cy7)



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Overview		
Quantity:	100 μL	
Target:	Ataxin 1 (ATXN1)	
Binding Specificity:	pSer775	
Reactivity:	Rat	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This Ataxin 1 antibody is conjugated to Cy7	
Application:	Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded	
	Sections) (IF (p))	
Product Details		
Immunogen:	KLH conjugated synthetic phosphopeptide derived from human Ataxin-1 around the	
	phosphorylation site of Ser775	
Isotype:	IgG	
Cross-Reactivity:	Rat	
Predicted Reactivity:	Human,Mouse,Dog,Cow,Pig,Horse,Rabbit	
Purification:	Purified by Protein A.	
Target Details		
Target:	Ataxin 1 (ATXN1)	

## **Target Details**

Alternative Name:	ATXN1 (ATXN1 Products)		
Background:	Synonyms: ATXN1, ATX1, D6S504E, SCA1, Ataxin-1, Spinocerebellar ataxia type 1,		
	ATX1_HUMAN.		
	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-II		
	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		
	contains41-81 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. [provided by RefSeq].		
Gene ID:	6310		
Pathways:	Synaptic Membrane		
Application Details			
Application Notes:	IF(IHC-P) 1:50-200		
	IF(IHC-F) 1:50-200		
	IF(ICC) 1:50-200		
Restrictions:	For Research Use only		
Handling			
Format:	Liquid		
Concentration:	1 μg/μL		
Buffer:	Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol.		

## Handling

Preservative:	ProClin	
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only.	
Storage:	-20 °C	
Storage Comment:	Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles.	
Expiry Date:	12 months	