

**Superoxide Dismutase 1 (SOD-1) Antibody**  
**Rabbit Polyclonal Antibody**  
**Catalog # ABV11197****Specification**

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**Superoxide Dismutase 1 (SOD-1) Antibody - Product Information**

Application	WB
Primary Accession	<a href="#">P00441</a>
Other Accession	<a href="#">AAR21563</a>
Reactivity	Human, Mouse, Rat
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	15936

**Superoxide Dismutase 1 (SOD-1) Antibody - Additional Information****Gene ID** 6647

Positive Control	Western Blot: 3T3 cell lysate, rat kidney lysate
Application & Usage	Western blot: 1:200
<b>Other Names</b>	
Superoxide Dismutase 1, SODC	

**Target/Specificity**  
SOD1**Antibody Form**  
Liquid**Appearance**  
Colorless liquid**Formulation**  
100 µg or 30 µg (0.5 mg/ml) of antibody in PBS containing 0.01 % BSA, 0.01 % thimerosal, and 50 % glycerol, pH 7.2**Handling**  
The antibody solution should be gently mixed before use.**Reconstitution & Storage**  
-20 °C**Background Descriptions****Precautions**

Superoxide Dismutase 1 (SOD-1) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

## **Superoxide Dismutase 1 (SOD-1) Antibody - Protein Information**

**Name** SOD1 ([HGNC:11179](#))

### **Function**

Destroys radicals which are normally produced within the cells and which are toxic to biological systems.

### **Cellular Location**

Cytoplasm. Nucleus. Note=Predominantly cytoplasmic; the pathogenic variants ALS1 Arg-86 and Ala-94 gradually aggregates and accumulates in mitochondria.

## **Superoxide Dismutase 1 (SOD-1) Antibody - Protocols**

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

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

## **Superoxide Dismutase 1 (SOD-1) Antibody - Images**

## **Superoxide Dismutase 1 (SOD-1) Antibody - Background**

Superoxide Dismutase (SOD) is an oxidoreductase that catalyzes the reaction between superoxide anions and hydrogen to yield molecular oxygen and hydrogen peroxide. The enzyme protects the cell against dangerous levels of superoxide. It belongs to the Cu-Zn superoxide dismutase family. It binds copper and zinc ions. The encoded isozyme is a soluble cytoplasmic protein, acting as a homodimer to convert naturally-occurring but harmful superoxide radicals to molecular oxygen and hydrogen peroxide. The other isozyme is a mitochondrial protein. Mutations in this gene have been implicated as causes of familial amyotrophic lateral sclerosis. Rare transcript variants have been reported for this gene. SOD1 destroys radicals which are normally produced within the cells and which are toxic to biological systems. Defects in SOD1 are the cause of amyotrophic lateral sclerosis type 1 (ALS1). ALS1 is a familial form of amyotrophic lateral sclerosis, a neurodegenerative disorder affecting upper and lower motor neurons and resulting in fatal paralysis. Sensory abnormalities are absent. Death usually occurs within 2 to 5 years. The etiology of amyotrophic lateral sclerosis is likely to be multifactorial, involving both genetic and environmental factors. The disease is inherited in 5-10% of cases leading to familial forms.