# Product datasheet Anti-SOD2 Antibody (Clone#2B12B1) Catalog Number: M00349-3

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BOSTER BIOLOGICAL TECHNOLOGY

Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

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Product Name	Anti-SOD2 Antibody (Clone#2B12B1)
Gene Name	SOD2
Source	Mouse
Clonality	Monoclonal
Isotype	IgG2b
Species Reactivity	human, mouse
Tested Application	WB, IHC
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.
Immunogen	A synthetic peptide corresponding to a sequence at the C-terminus of human SOD2, different from the related mouse sequence by one amino acid, and from the related rat sequence by fo amino acids.
Concentration	500 ug/ml
Purification	protein G purified.
Observed MW	25 kDa
Dilution Ratios	Western blot (WB):  Immunohistochemistry (IHC):  (Boiling the paraffin sections in 10mM citrate buffer,pH6.0,or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must

## **Storage**

12 months from date of receipt, -20°C as supplied.

determined by end user.

### **Background Information**

SOD2(Superoxide Dismutase 2), also called IPO-B or MNSOD, is a mitochondrial matrix enzyme that scavenges oxygen radicals produced by the extensive oxidation-reduction and electron transport reactions occurring in mitochondria. This gene is a member of the iron/manganese superoxide dismutase family. Using a somatic cell hybrid panel containing different segments of chromosome 6, they demonstrated that SOD2 is located in the region 6q25.3-qter which, together with the FISH analysis, indicated that SOD2 is in the distal portion of 6q25. The SOD2 gene encodes an intramitochondrial free radical scavenging enzyme that is the first line of defense against superoxide produced as a byproduct of oxidative phosphorylation. Adenoassociated viral delivery of the human SOD2 gene resulted in suppression of optic nerve degeneration and rescue of retinal ganglion cells. The findings suggested that reactive oxygen species contributed to retinal cell death and optic nerve damage in mice with complex I deficiency, and that expression of SOD2 attenuated the disease process.

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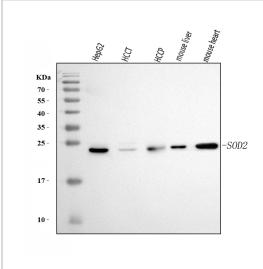
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## Reference

Anti-SOD2 Antibody (Clone#2B12B1)被引用在4文献中。

## **Selected Validation Data**



Western blot analysis of SOD2 using anti-SOD2 antibody (M00349-3). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: HEPG2 whole cell lysates,

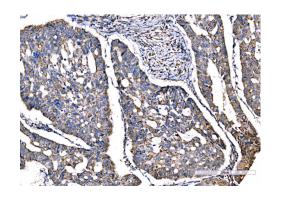
Lane 2: HCCP tissue lysates,

Lane 3: HCCP tissue lysates,

Lane 4: mouse liver tissue lysates,

Lane 5: mouse heart tissue lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with mouse anti-SOD2 antigen affinity purified monoclonal antibody (M00349-3) at a dilution of 1:1000 and probed with a goat anti-mouse IgG-HRP secondary antibody (Catalog # BA1050). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for SOD2 at approximately 25 kDa. The expected band size for SOD2 is at 25 kDa.



IHC analysis of SOD2 using anti-SOD2 antibody (M00349-3). SOD2 was detected in a paraffin-embedded section of human Adenocarcinoma of the right colon tissue. The tissue section was incubated with mouse anti-SOD2 Antibody (M00349-3) at a dilution of 1:200 and developed using HRP Conjugated mouse IgG Super Vision Assay Kit (Catalog # SV0001) with DAB (Catalog # AR1027) as the chromogen.