

SOD-2 Polyclonal Antibody

Catalog No: YT5575

Reactivity: Human; Mouse; Rat

Applications: WB;IHC;IF;ELISA

Target: SOD2

Fields: >>FoxO signaling pathway;>>Peroxisome;>>Longevity regulating

pathway;>>Longevity regulating pathway - multiple species;>>Huntington disease;>>Chemical carcinogenesis - reactive oxygen species;>>Lipid and

atherosclerosis

Gene Name: SOD2

Protein Name: Superoxide dismutase [Mn] mitochondrial

P04179

P09671

Human Gene Id: 6648

Human Swiss Prot

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No:

Mouse Gene Id: 20656

Mouse Swiss Prot

No:

Rat Gene ld: 24787

Rat Swiss Prot No: P07895

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

Internal region of human SOD2. AA range:91-140

Specificity: SOD-2 Polyclonal Antibody detects endogenous levels of SOD-2 protein.

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

Source: Polyclonal, Rabbit, IgG

1/3



No4:

1

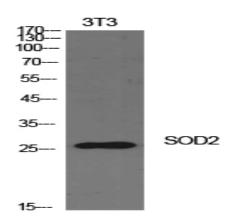
IHC: 100-300.WB 1:500 - 1:2000. ELISA: 1:10000.. IF 1:50-200 **Dilution: Purification:** The antibody was affinity-purified from rabbit antiserum by affinitychromatography using epitope-specific immunogen. Concentration: 1 mg/ml -15°C to -25°C/1 year(Do not lower than -25°C) **Storage Stability: Observed Band:** 24kD **Cell Pathway:** Huntington's disease; **Background:** This gene is a member of the iron/manganese superoxide dismutase family. It encodes a mitochondrial protein that forms a homotetramer and binds one manganese ion per subunit. This protein binds to the superoxide byproducts of oxidative phosphorylation and converts them to hydrogen peroxide and diatomic oxygen. Mutations in this gene have been associated with idiopathic cardiomyopathy (IDC), premature aging, sporadic motor neuron disease, and cancer. Alternative splicing of this gene results in multiple transcript variants. A related pseudogene has been identified on chromosome 1. [provided by RefSeq, Apr 2016], **Function:** catalytic activity: 2 superoxide + 2 H(+) = O(2) + H(2)O(2), cofactor: Binds 1 manganese ion per subunit., disease: Genetic variation in SOD2 is associated with susceptibility to diabetic nephropathy [MIM:612634]; also called susceptibility to microvascular complications of diabetes type 6 (MVCD6). Diabetic nephropathy is a kidney disease and resultant kidney function impairment due to the long standing effects of diabetes on the microvasculature (glomerulus) of the kidney. Features include increased urine protein and declining kidney function., function: Destroys radicals which are normally produced within the cells and which are toxic to biological systems., online information: Superoxide dismutase entry, online information: The Singapore human mutation and polymorphism database, PTM: Nitrated under oxidative stress. Nitration coupled with oxidation inhibits the catalytic activity., similarity: Belo Subcellular Mitochondrion matrix. Location: **Expression:** Brain, Colon, Heart, Liver, Lung, Mammary carcinoma, Tongue, Tag: orthogonal Sort: 1



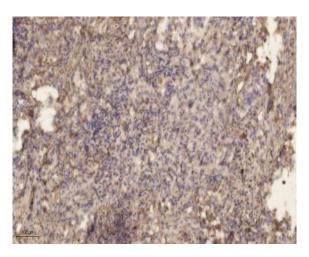
Host: Rabbit

Modifications: Unmodified

Products Images



Western Blot analysis of NIH-3T3 cells using SOD-2 Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000



Immunohistochemical analysis of paraffin-embedded human lung 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).