

Product datasheet

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ARG57160 anti-SOD1 antibody [6F1]

Package: 50 μl Store at: -20°C

Summary

Product Description Mouse Monoclonal antibody [6F1] recognizes SOD1

Tested Reactivity Hu

Tested Application FACS, WB

Host Mouse

Clonality Monoclonal

Clone 6F1

Isotype IgG1, kappa

Target Name SOD1
Species Human

Immunogen Recombinant fragment around aa. 1-154 of Human SOD1

Conjugation Un-conjugated

Alternate Names homodimer; EC 1.15.1.1; SOD; HEL-S-44; Superoxide dismutase [Cu-Zn]; ALS1; Superoxide dismutase 1;

IPOA; ALS; hSod1

Application Instructions

| Application table | Application | Dilution |
|-------------------|--|-----------------|
| | FACS | Assay-dependent |
| | WB | Assay-dependent |
| Application Note | * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist. | |

Properties

Form Liquid

Purification Purification with Protein A.

Buffer PBS (pH 7.4), 0.02% Sodium azide and 10% Glycerol.

Preservative 0.02% Sodium azide

Stabilizer 10% Glycerol

Concentration 1 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed before use.

Note For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Database links GeneID: 6647 Human

Swiss-port # P00441 Human

Gene Symbol SOD1

Gene Full Name superoxide dismutase 1, soluble

Background The protein encoded by this gene binds copper and zinc ions and is one of two isozymes responsible for

destroying free superoxide radicals in the body. The encoded isozyme is a soluble cytoplasmic protein, acting as a homodimer to convert naturally-occuring 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. [provided by RefSeq, Jul 2008]

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

systems. [UniProt]

Calculated Mw 16 kDa

PTM Unlike wild-type protein, the pathogenic variants ALS1 Arg-38, Arg-47, Arg-86 and Ala-94 are

polyubiquitinated by RNF19A leading to their proteasomal degradation. The pathogenic variants ALS1 Arg-86 and Ala-94 are ubiquitinated by MARCH5 leading to their proteasomal degradation. The ditryptophan cross-link at Trp-33 is responsible for the non-disulfide-linked homodimerization.

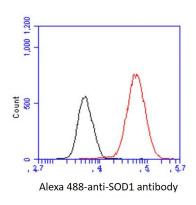
Such modification might only occur in extreme conditions and additional experimental evidence is required.

Palmitoylation helps nuclear targeting and decreases catalytic activity.

Succinylation, adjacent to copper catalytic site, probably inhibits activity. Desuccinylation by SIRT5

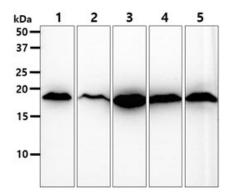
enhances activity.

Images



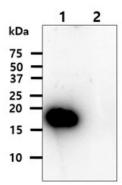
ARG57160 anti-SOD1 antibody [6F1] FACS image

Flow Cytometry: Hep3B cell line stained with ARG57160 anti-SOD1 antibody [6F1] at 2-5 μ g for 1x10^6 cells (red line). Secondary antibody: Goat anti-Mouse IgG Alexa fluor 488 conjugate. Isotype control antibody: Mouse IgG (black line).



ARG57160 anti-SOD1 antibody [6F1] WB image

Western blot: 40 μg of 1) A549, 2) A431, 3) 293T, 4) PC3, and 5) U87MG cell lysates stained with ARG57160 anti-SOD1 antibody [6F1] at 1:1000.



ARG57160 anti-SOD1 antibody [6F1] WB image

Western blot: 20 ng of 1) SOD1, and 2) SOD2 recombinant proteins stained with ARG57160 anti-SOD1 antibody [6F1] at 1:1000.