

Product datasheet for SC302253

SOD2 (NM 001024466) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: SOD2 (NM_001024466) Human Untagged Clone

Tag: Tag Free Symbol: SOD2

Synonyms: GClnc1; IPO-B; IPOB; Mn-SOD; MNSOD; MVCD6

Mammalian Cell None

Selection:

Vector: pCMV6-XL5

E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene ORF sequence for NM_001024466 edited

TGCAAAAAGTAA

Restriction Sites: Please inquire ACCN: NM 001024466

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: The ORF of this clone has been fully sequenced and found to be a perfect match to

NM 001124466.1.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).



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Reconstitution Method:

- 1. Centrifuge at 5,000xg for 5min.
- 2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
- 3. Close the tube and incubate for 10 minutes at room temperature.
- 4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
- 5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

RefSeq: <u>NM 001024466.1</u>, <u>NP 001019637.1</u>

 RefSeq Size:
 918 bp

 RefSeq ORF:
 552 bp

 Locus ID:
 6648

 UniProt ID:
 P04179

 Cytogenetics:
 6q25.3

Protein Families: Druggable Genome, Transcription Factors

Protein Pathways: Huntington's disease

Gene Summary: 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] Transcript Variant: This variant (3) lacks an alternate in-frame exon in the central coding region, and splices out a region of the 3' UTR, compared to variant 1. The encoded isoform (B)

is shorter than isoform A. Both variants 3 and 4 encode the same isoform (B).