

# **Product datasheet for TP720709M**

### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## SOD2 (NM 000636) Human Recombinant Protein

#### **Product data:**

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human superoxide dismutase 2, mitochondrial (SOD2),

nuclear gene encoding mitochondrial protein, transcript variant 1

Species: Human
Expression Host: E. coli

Expression cDNA Clone

or AA Sequence:

Lys25-Lys222

Tag: N-His

Predicted MW: 23.24 kDa
Concentration: lot specific

**Purity:** >95% as determined by SDS-PAGE and Coomassie blue staining

Buffer: Provided lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl

Endotoxin: Endotoxin level is < 0.1 ng/μg of protein (< 1 EU/μg)

Storage: Store at -80°C.

Stability: Stable for at least 6 months from date of receipt under proper storage and handling

conditions.

RefSeq: <u>NP 000627</u>

**Locus ID:** 6648

UniProt ID: <u>P04179</u>, <u>A0A384NL29</u>

RefSeq Size: 1593 Cytogenetics: 6q25.3 RefSeq ORF: 666

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





## SOD2 (NM\_000636) Human Recombinant Protein - TP720709M

**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]

**Protein Families:** Druggable Genome, Transcription Factors

**Protein Pathways:** Huntington's disease