

SOX2 Knockout 293T Cell Lysate, Homozygous

Catalog No.: RM02022

Basic Information

Catalog No.

RM02022

Category

Cell Lysate

Parental Cell line

293T

Genotype

Knockout

Gene Information

Gene Symbol

SOX2

Species

Human

Gene ID

6657

Swiss Prot

P48431

Synonyms

ANOP3; MCOPS3

Contact

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Background

This intronless gene encodes a member of the SRY-related HMG-box (SOX) family of transcription factors involved in the regulation of embryonic development and in the determination of cell fate. The product of this gene is required for stem-cell maintenance in the central nervous system, and also regulates gene expression in the stomach. Mutations in this gene have been associated with optic nerve hypoplasia and with syndromic microphthalmia, a severe form of structural eye malformation. This gene lies within an intron of another gene called SOX2 overlapping transcript (SOX2OT). [provided by RefSeq, Jul 2008]

Product Information

Description

SOX2 Knockout 293T Cell Line is engineered from 293T cell line with Gene-Editing technology.

Allele-1:14bp deletion in exon1

Allele-2:14bp deletion in exon1

Mammalian cells such as human, rat and mouse cells are normally diploid with two alleles. Homozygote: both alleles were knocked out, mRNA has no signal, no expression of proteins. Heterozygote: only one allele was knocked out, the mRNA transcript levels was decreased compared to wild type, and the protein expression levels was also lower than that of the wild type.

Packaging

1 vial parental cell Lysate and 1 vial knockout cell Lysate

Shipping Conditions

4°C

Amount

50μL, 2μg/μL.

Storage

Lysate is stable for 12 months when stored at -20°C. Minimizing freeze-thaw cycles.

Protocol

To be used as WB control. Lysate is supplied in 1× SDS sample buffer (2% SDS, 60 mM Tris-HCl pH 6.8, 10% Glycerol, 0.02% Bromophenol blue, 60 mM beta-mercaptoethanol). Lysate should be boiled for 3 - 5 minutes before loading onto gel.

Sequencing data

WT GCTGGCCCCGGCG*****AGCGGGTCGGGGT
Mut GCTGGCCCCGGCG***Deletion***AGCGGGTCGGGGT
Allele-1: 14bp deletion in exon1

WT GCTGGCCCCGGCG*****AGCGGGTCGGGGT
Mut GCTGGCCCCGGCG***Deletion***AGCGGGTCGGGGT
Allele-2: 14bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and SOX2 knockout (KO) 293T cells, using sanger sequencing.