

FOXA1 Knockdown 293T Cell Lysate, Heterozygous

Catalog No.: RM02568

Basic Information

Catalog No.

RM02568

Category

Cell Lysate

Parental Cell line

293T

Genotype

Knockdown

Gene Information

Gene Symbol

FOXA1

Species

Human

Gene ID

3169

Swiss Prot

P55317

Synonyms

HNF3A; TCF3A

Contact

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Background

This gene encodes a member of the forkhead class of DNA-binding proteins. These hepatocyte nuclear factors are transcriptional activators for liver-specific transcripts such as albumin and transthyretin, and they also interact with chromatin. Similar family members in mice have roles in the regulation of metabolism and in the differentiation of the pancreas and liver. [provided by RefSeq, Jul 2008]

Product Information

Description

FOXA1 Knockdown 293T Cell Line is engineered from 293T cell line with Gene-Editing technology.

Allele-1:137bp deletion in exon2

Allele-2:WT

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 AACATGACCCCGGC*****TGAGCCCGAGCGGC
Mut AACATGACCCCGGC***Deletion***TGAGCCCGAGCGGC
Allele-1: 137bp deletion in exon2

WT AACATGACCCCGGC*****TGAGCCCGAGCGGC
Mut AACATGACCCCGGC*****TGAGCCCGAGCGGC
Allele-2: WT

Genome sequence analysis of PCR products from parental (WT) and FOXA1 Knockdown (KD) 293T cells, using sanger sequencing.