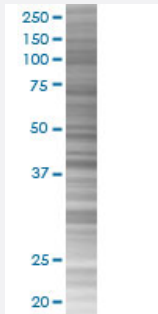


SHOX2 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # H00006474-T01

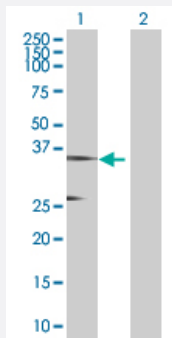
Size 100 uL

Applications



SDS-PAGE Gel

SHOX2 transfected lysate.



Western Blot

Lane 1: SHOX2 transfected lysate (39.16 KDa)

Lane 2: Non-transfected lysate.

Specification

Transfected Cell Line 293T

Plasmid pCMV-SHOX2 full-length

Host Human

Theoretical MW (kDa) 39.16

Quality Control Testing Transient overexpression cell lysate was tested with Anti-SHOX2 antibody ([H00006474-B01](#)) by Western Blots.
SDS-PAGE Gel
SHOX2 transfected lysate.
Western Blot
Lane 1: SHOX2 transfected lysate (39.16 KDa)
Lane 2: Non-transfected lysate.

Storage Buffer

1X Sample Buffer (50 mM Tris-HCl, 2% SDS, 10% glycerol, 300 mM 2-mercaptoethanol, 0.01% Bromophenol blue)

Storage Instruction

Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot

Gene Info — SHOX2

Entrez GeneID[6474](#)**GeneBank Accession#**[BC008829.2](#)**Protein Accession#**

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Gene Name

SHOX2

Gene Alias

OG12, OG12X, OGI2X, SHOT

Gene Description

short stature homeobox 2

Omim ID[602504](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

This gene is a member of the homeobox family of genes that encode proteins containing a 60-amino acid residue motif that represents a DNA binding domain. Homeobox genes have been characterized extensively as transcriptional regulators involved in pattern formation in both invertebrate and vertebrate species. Several human genetic disorders are caused by aberrations in human homeobox genes. This locus represents a pseudoautosomal homeobox gene that is thought to be responsible for idiopathic short stature, and it is implicated in the short stature phenotype of Turner syndrome patients. This gene is considered to be a candidate gene for Cornelia de Lange syndrome. Alternative splicing results in multiple transcript variants. [provided by RefSeq]

Other Designations

SHOX homologous gene on chromosome 3|short stature homeobox homolog