

Product datasheet for TP328253L

OriGene Technologies, Inc.

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SHOX2 (NM_001163678) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human short stature homeobox 2 (SHOX2), transcript variant 3, 1 mg

Species: Human Expression Host: HEK293T

Expression cDNA >RC228253 representing NM 001163678

Clone or AA Sequence:

Red=Cloning site Green=Tags(s)

SASAASVVAAAAAAKTTSKNSSIADLRLKAKKHAAALGL

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK
Predicted MW: 33.4 kDa

Concentration: >0.05 μg/μL as determined by microplate BCA method

Purity: > 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer: 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

Preparation: NULL or Add: Recombinant proteins was captured through anti-DDK affinity column followed by

conventional chromatography steps.

Note: For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and handling

conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 001157150

Locus ID: 6474



SHOX2 (NM_001163678) Human Recombinant Protein - TP328253L

 UniProt ID:
 O60902

 Cytogenetics:
 3q25.32

 RefSeq ORF:
 957

Synonyms: OG12; OG12X; SHOT

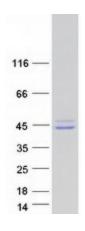
Summary: This gene is a member of the homeobox family of genes that encode proteins containing a 60-

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, Jul 2009]

Protein Families: Transcription Factors

Product images:



Coomassie blue staining of purified SHOX2 protein (Cat# [TP328253]). The protein was produced from HEK293T cells transfected with SHOX2 cDNA clone (Cat# [RC228253]) using MegaTran 2.0 (Cat# [TT210002]).