

# Product datasheet for TP328253M

## 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, 100 $\mu g$
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC228253 representing NM_001163678 <mark>Red</mark> =Cloning site Green=Tags(s)
	MEELTAFVSKSFDQKVKEKKEAITYREVLESGPLRGAKEPTGCTEAGRDDRSSPAVRAAGGGGGGGGGGGG GGGGGGGVGGGGAGGGAGGGRSPVRELDMGAAERSREPGSPRLTEVSPELKDRKEDAKGMEDEGQTKIKQ RRSRTNFTLEQLNELERLFDETHYPDAFMREELSQRLGLSEARVQVWFQNRRAKCRKQENQLHKGVLIGA ASQFEACRVAPYVNVGALRMPFQQVQAQLQLDSAVAHAHHHLHPHLAAHAPYMMFPAPPFGLPLATLAAD 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:	<u>NP 001157150</u>
Locus ID:	6474



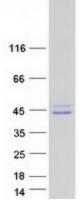
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	SHOX2 (NM_001163678) Human Recombinant Protein – TP328253M
UniProt ID:	<u>060902</u>
Cytogenetics:	3q25.32
RefSeq ORF:	957
Synonyms:	OG12; OG12X; SHOT
Summary:	This gene is a member of the homeobox family of genes that encode proteins containing a 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]).

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