

Product datasheet for RC212490L3

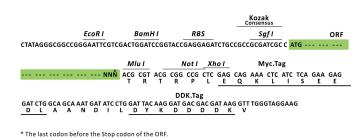
SHOX (NM_006883) Human Tagged Lenti ORF Clone

Product data:

OriGene Technologies, Inc.

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Expression Plasmids
SHOX (NM_006883) Human Tagged Lenti ORF Clone
Myc-DDK
SHOX
GCFX; PHOG; SHOXY; SS
Puromycin
pLenti-C-Myc-DDK-P2A-Puro (PS100092)
Chloramphenicol (34 ug/mL)
The ORF insert of this clone is exactly the same as(RC212490).
Sgfl-Mlul
Cloning sites used for ORF Shuttling: Sgf 1 ORF Mlu 1 GCG ATC GC C ATG // NNN ACG CGT



ACCN: ORF Size: NM_006883 675 bp



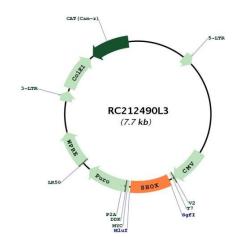
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ORIGENE SHOX (NM_006883) Human Tagged Lenti ORF Clone – RC212490L3
OTI Disclaimer:	The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <u>More info</u>
OTI Annotation:	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.
Components:	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
Reconstitution Method:	 Centrifuge at 5,000xg for 5min. Carefully open the tube and add 100ul of sterile water to dissolve the DNA. Close the tube and incubate for 10 minutes at room temperature. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.
RefSeq:	<u>NM 006883.2, NP 006874.1</u>
RefSeq Size:	1951 bp
RefSeq ORF:	678 bp
Locus ID:	6473
UniProt ID:	<u>015266</u>
Cytogenetics:	X;Y
Protein Families:	Transcription Factors
MW:	25.5 kDa
Gene Summary:	This gene belongs to the paired homeobox family and is located in the pseudoautosomal region 1 (PAR1) of X and Y chromosomes. Defects in this gene are associated with idiopathic growth retardation and in the short stature phenotype of Turner syndrome patients. This gene is highly conserved across species from mammals to fish to flies. Alternatively spliced transcript variants encoding different isoforms have been noted for this gene. [provided by RefSeq, Jul 2008]

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Product images:



Circular map for RC212490L3

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