

## Product datasheet for SC307123

### SOX5 (NM\_178010) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	SOX5 (NM_178010) Human Untagged Clone
Tag:	Tag Free
Symbol:	SOX5
Synonyms:	L-SOX5; L-SOX5B; L-SOX5F; LAMSHF
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>SC307123 representing NM_178010. Blue=Insert sequence Red=Cloning site Green=Tag(s)

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GCTCGTTT TAGTGAACCGTCAGAATTTTGT AATACGACTCACTATAGGGCGCCGGGAATTCGTCGACTG
GATCCGGTACCGAGGAGATCTGCCGCC CGGATCGCC
ATGCATGATGAAGTGGCAGCCACTGAACCTATCAGCTAAACCCAAGACCTCTGATGGCAAATCACCC
ACATCACCCACCTCTCCCATATGCCAGCTCTGAGAATAAACAGTGGGGCAGGCCCCCTCAAAGCCTCT
GTCCCAGCAGCGTTAGCTAGTCCTTCAGCCAGAGTTAGCACAATAGGTTACTTAAATGACCATGATGCT
GTCACCAAGGCAATCCAAGAAGCTCGGCAATGAAGGAGCAACTCCGACGGGAACAACAGGTGCTTGAT
GGGAAGGTGGCTGTTGTGAATAGTCTGGGTCTCAATAACTGCCGAACAGAAAAGAAAAACAACACTG
GAGAGTCTGACTCAGCAACTGGCAGTTAAACAGAATGAAGAAGGAAAATTTAGCCATGCAATGATGGAT
TTCAATCTGAGTGGAGATTCTGATGGAAGTGCTGGAGTCTCAGAGTCAAGAATTTATAGGGAATCCCGA
GGGCGTGGTAGCAATGAACCCACATAAAGCGTCCAATGAATGCCTTCATGGTGTGGGCTAAAGATGAA
CGGAGAAAGATCCTTCAAGCCTTCTGACATGCACAACCTCCAACATCAGCAAGATATTGGGATCTCGC
TGGAAAGCTATGACAAACCTAGAGAAACAGCCATATTAAGAGGAGCAAGCCCGTCTCAGCAAGCAGCAC
CTGGAGAAGTACCCTGACTATAAGTACAAGCCAGGCCAAAGCGCACCTGCCTGGTGGATGGCAAAAAG
CTGCGCATTGGTGAATACAAGGCAATCATGCGCAACAGGCGGCAGGAAATGCGGCAGTACTTCAATGTT
GGGCAACAAGCACAGATCCCCATTGCCACTGCTGGTGTGTGTACCCTGGAGCCATCGCCATGGCTGGG
ATGCCCTCCCCTCACCTGCCCTCGGAGCACTCAAGCGTGTCTAGCAGCCAGCCATGGGATGCTGTT
ATCCAGAGCACTTACGGTGTGAAAGGAGAGGAGCCACATATCAAAGAAGAGATACAGGCCGAGGACATC
AATGGAGAAATTTATGATGAGTACGACGAGGAAGAGGATGATCCAGATGTAGATTATGGGAGTGACAGT
GAAAACCATATTGCAGGACAAGCCAACTGA
ACGCGTACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT
TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC
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Restriction Sites: SgfI-MluI



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Plasmid Map:	□
ACCN:	NM_178010
Insert Size:	1134 bp
OTI Disclaimer:	Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).
OTI Annotation:	This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.
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:	<ol style="list-style-type: none"><li>1. Centrifuge at 5,000xg for 5min.</li><li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li><li>3. Close the tube and incubate for 10 minutes at room temperature.</li><li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li><li>5. 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.</li></ol>
RefSeq:	<a href="#">NM_178010.2</a>
RefSeq Size:	3133 bp
RefSeq ORF:	1134 bp
Locus ID:	6660
UniProt ID:	<a href="#">P35711</a>
Cytogenetics:	12p12.1
Protein Families:	Transcription Factors
MW:	42 kDa

**Gene Summary:**

This gene encodes a member of the SOX (SRY-related HMG-box) family of transcription factors involved in the regulation of embryonic development and in the determination of the cell fate. The encoded protein may act as a transcriptional regulator after forming a protein complex with other proteins. The encoded protein may play a role in chondrogenesis. A pseudogene of this gene is located on chromosome 8. Multiple transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq, Jul 2008]

Transcript Variant: This variant (3) contains an alternate first exon in place of much of the 5' UTR and coding sequence compared to variant 1. The resulting isoform (c) has a shorter and distinct N-terminus compared to isoform a. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.