

## Product datasheet for **MC207403**

### Sox2 (NM\_011443) Mouse Untagged Clone

#### Product data:

**Product Type:** Expression Plasmids  
**Product Name:** Sox2 (NM\_011443) Mouse Untagged Clone  
**Tag:** Tag Free  
**Symbol:** Sox2  
**Synonyms:** lc; lcc; Sox; Sox-2; ys; ysb  
**Mammalian Cell Selection:** Neomycin  
**Vector:** pCMV6-Entry (PS100001)  
**E. coli Selection:** Kanamycin (25 ug/mL)  
**Fully Sequenced ORF:** >MC207403 representing NM\_011443  
**Red**=Cloning site **Blue**=ORF **Orange**=Stop codon

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
GCC**CGATCGCC**

ATGTATAACATGATGGAGACGGAGCTGAAGCCGCCGGGCCCGCAGCAAGCTTCGGGGGGCGGCGGGAG  
GAGGCAACGCCACGGCGGCGGACCGGCCAACCAGAAGAACAGCCCGGACCGCTCAAGAGGCCCAT  
GAACGCCCTCATGGTATGGTCCCGGGGCGAGCGGCTAAGATGGCCAGGAGAACCCTAAGATGCACAAC  
TCGGAGATCAGCAAGCGCTGGGCGGGAGTGAAACTTTGTCCGAGACCGAGAAGCGGCCGTTTCATCG  
ACGAGGCCAAGCGCTGCGGCTCTGCACATGAAGGAGCACCCGATTATAAATACCGGCCGCGGGGAA  
AACCAAGACGCTCATGAAGAAGGATAAGTACACGCTTCCCGAGGCTTGCTGGCCCCGGCGGGAACAGC  
ATGGCGAGCGGGTTGGGTGGGCGCCGCGCTGGGTGCGGGCGTGAACCAGCGCATGGACAGCTACGCGC  
ACATGAACGGCTGGAGCAACGGCAGCTACAGCATGATGCAGGAGCAGCTGGGCTACCCGCAGCACCCGGG  
CCTCAACGCTCACGGCGCGGCACAGATGAACCGATGCACCGCTACGACGTACGCGCCCTGCAGTACAAC  
TCCATGACCAGCTCGCAGACCTACATGAACGGCTCGCCACCTACAGCATGTCTACTCGCAGCAGGGCA  
CCCCGGTATGGCGTGGGCTCCATGGGCTCTGTGGTCAAGTCCGAGGCCAGCTCCAGCCCCCGTGGT  
TACCTTCTCTCCACTCCAGGGCGCCCTGCCAGGCCGGGACCTCCGGGACATGATCAGCATGTACCTC  
CCCGCGCGGAGGTGCCGAGCCCGCTGCGCCAGTAGACTGCACATGGCCAGCACTACCAGAGCGGCC  
CGGTGCCCGGCACGGCCATTAACGGCACACTGCCCTGTGCACATG**TGA**

**ACGCGT**ACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCTGGATT  
ACAAGGATGACGACGATAAGGTTTAA

**Chromatograms:** [https://cdn.origene.com/chromatograms/ja1770\\_c05.zip](https://cdn.origene.com/chromatograms/ja1770_c05.zip)

**Restriction Sites:** SgfI-MluI



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ACCN:	NM_011443
Insert Size:	960 bp
OTI Disclaimer:	<p>Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at <a href="mailto:custsupport@origene.com">custsupport@origene.com</a> or by calling 301.340.3188 option 3 for pricing and delivery.</p> <p>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. <a href="#">More info</a></p>
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="#">BC057574</a> , <a href="#">AAH57574</a>
RefSeq Size:	2457 bp
RefSeq ORF:	960 bp
Locus ID:	20674
UniProt ID:	<a href="#">P48432</a>
Cytogenetics:	3 16.93 cM
Gene Summary:	<p>This intronless gene encodes a member of the SRY-related HMG-box (SOX) family of transcription factors involved in the regulation of embryonic development and in the determination of cell fate. The product of this gene is required for stem-cell maintenance in the central nervous system, and also regulates gene expression in the stomach. Mutations in a similar gene in human have been associated with optic nerve hypoplasia and with syndromic microphthalmia, a severe form of structural eye malformation. This gene lies within an intron of another gene called SOX2 overlapping transcript (Sox2ot). [provided by RefSeq, Sep 2015]</p>