

## Product datasheet for **SC314589**

### FOXP2 (NM\_148898) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	FOXP2 (NM_148898) Human Untagged Clone
Tag:	Tag Free
Symbol:	FOXP2
Synonyms:	CAGH44; SPCH1; TNRC10
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)



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**Fully Sequenced ORF:** >OriGene sequence for NM\_148898 edited  
 ATGATGCAGGAATCTGCGACAGAGACAATAAGCAACAGTTCATGAATCAAAATGGAATG  
 AGCACTCTAAGCAGCCAATTAGATGCTGGCAGCAGAGATGGAAGATCAAGTGGTGACACC  
 AGCTCTGAAGTAAACACAGTAGAACTGCTGCATCTGCAACAACAGCAGGCTCTCCAGGCA  
 GCAAGACAACCTCTTTTACAGCAGCAACAAGTGGATTGAAATCTCCTAAGAGCAGTGAT  
 AAACAGAGACCACTGCAGGAATTGCTTCCAGAAACAAAATTATGTATCTGTGGCCACTCT  
 TCTGGTGATGGGCATCCTCACAACACATTTGCAGTGCCTGTGTGTCAGTGGCCATGATGACT  
 CCCCAGGTGATCACCCCTCAGCAAAATGCAGCAGATCCTTCAGCAACAAGTCTGTCTCCT  
 CAGCAGCTACAAGCCCTTCTCCAACAACAGCAGGCTGTGATGCTGCAGCAGCAACAACCTA  
 CAAGAGTTTTACAAGAAACAGCAAGAGCAGTTACATCTTCAGCTTTTGCAGCAGCAGCAG  
 CAACAGCAGCAGCAGCAACAACAGCAGCAACAACAGCAGCAGCAACAACAACAACAG  
 CAGCAACAACAGCAGCAGCAGCAGCAACAGCAGCAGCAGCAGCAACAGCATCCTGGAAAG  
 CAAGCGAAAGAGCAGCAGCAGCAGCAGCAGCAACAGCAATTGGCAGCCAGCAGCTT  
 GTCTTCCAGCAGCAGCTTCTCCAGATGCAACAACCTCCAGCAGCAGCAGCATCTGCTCAGC  
 CTTCCAGCGTCAGGGACTCATCTCCATTCCACCTGGCCAGGCAGCACTTCTGTCCAAATCG  
 CTGCCTCAAGCTGGCTTAAAGTCTGCTGAGATTAGCAGTTATGGAAGAAGTGACTGGA  
 GTTCACAGTATGGAAGACAATGGCATTAAACATGGAGGGCTAGACCTCACTACTAACAAT  
 TCCTCCTCGACTACCTCCTCAACACTTCCAAAGCATCACCAATAACTCATCATTCC  
 ATAGTGAATGGACAGTCTTCAGTTCTAAGTGAAGACGAGACAGCTCGTCACATGAGGAG  
 ACTGGGGCCTCTCACACTCTCTATGGCCATGGAGTTTGCAAAATGGCCAGGCTGTGAAAGC  
 ATTTGTGAAGATTTTGGACAGTTTTTAAAGCACCTTAAACAATGAACACGCATTGGATGAC  
 CGAAGCACTGCTCAGTGTGAGTGCAAAATGCAGGTGGTGAACAGTTAGAAAATACAGCTT  
 TCTAAAGAACCGAACGTCCTTCAAGCAATGATGACCCACTTGACATGCGACCCCTCAGAG  
 CCCAAACCATCTCCCAAACCTCTAAATCTGGTGTCTAGTGTACCATGTGGAAGAATATG  
 TTGGAGACATCCCCACAGAGCTTACCTCAAACCCCTACCACACCAACGGCCCCAGTCACC  
 CCGATTACCCAGGGACCCTCAGTAATCACCCAGCCAGTGTGCCAATGTGGGAGCCATA  
 CGAAGGCGACATTAGACAAAATACAACATTCCCATGTCATCAGAAAATGCCCAAACCTAT  
 GAATTTTATAAAAATGCAGATGTCAGACCTCCATTTACTTATGCAACTCTCATAAGGCAG  
 GCTATCATGGAGTCATCTGACAGGCAGTTAACACTTAATGAAATTTACAGCTGGTTTACA  
 CGGACATTTGCTTACTTCAGGCGTAATGCAGCAACTTGGGAAGAATGCAGTACGTACATAAT  
 CTTAGCCTGCACAAGTGTGTTGTCGAGTAGAAAATGTTAAAGGAGCAGTATGGACTGTG  
 GATGAAGTAGAATACCAGAAGCGAAGGTACAAAAGATAACAGGAAGTCCAACTTAGTA  
 AAAAATATACCTACCAGTTTAGGCTATGGAGCAGCTCTTAATGCCAGTTTGCAGGCTGCC  
 TTGGCAGAGAGCAGTTTACCTTTGCTAAGTAATCCTGGACTGATAAATAATGCATCCAGT  
 GGCTACTGCAGGCCGTCCACGAAGACCTCAATGGTTCTCTGGATCACATTGACAGCAAT  
 GGAAACAGTAGTCCGGGCTGCTCACCTCAGCCGCACATACATTCAATCCACGTCAAGGAA  
 GAGCCAGTGATTGCAGAGGATGAAGACTGCCCAATGTCCTTAGTGACAACAGCTAATCAC  
 AGTCCAGAATTAGAAGACGACAGAGAGATTGAAGAAGAGCCTTTATCTGAAGATCTGGAA  
 TGA

**Restriction Sites:** Please inquire

**ACCN:** NM\_148898

**Insert Size:** 2200 bp

**OTI Disclaimer:** 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 [custsupport@origene.com](mailto:custsupport@origene.com) or by calling 301.340.3188 option 3 for pricing and delivery.

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. [More info](#)

**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:**

1. Centrifuge at 5,000xg for 5min.
2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
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.

**RefSeq:** [NM\\_148898.1](#), [NP\\_683696.1](#)

**RefSeq Size:** 2547 bp

**RefSeq ORF:** 2223 bp

**Locus ID:** 93986

**UniProt ID:** [O15409](#)

**Cytogenetics:** 7q31.1

**Domains:** FH

**Protein Families:** Transcription Factors

**Gene Summary:**

This gene encodes a member of the forkhead/winged-helix (FOX) family of transcription factors. It is expressed in fetal and adult brain as well as in several other organs such as the lung and gut. The protein product contains a FOX DNA-binding domain and a large polyglutamine tract and is an evolutionarily conserved transcription factor, which may bind directly to approximately 300 to 400 gene promoters in the human genome to regulate the expression of a variety of genes. This gene is required for proper development of speech and language regions of the brain during embryogenesis, and may be involved in a variety of biological pathways and cascades that may ultimately influence language development. Mutations in this gene cause speech-language disorder 1 (SPCH1), also known as autosomal dominant speech and language disorder with orofacial dyspraxia. Multiple alternative transcripts encoding different isoforms have been identified in this gene.[provided by RefSeq, Feb 2010]

Transcript Variant: This variant (2) encodes the longest isoform (II). 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.