

Product datasheet for SC317945

FOXC1 (NM_001453) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: FOXC1 (NM_001453) Human Untagged Clone

Tag: Tag Free Symbol: FOXC1

Synonyms: ARA; ASGD3; FKHL7; FREAC-3; FREAC3; IGDA; IHG1; IRID1; RIEG3

Mammalian Cell

Selection:

Neomycin

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

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Fully Sequenced ORF: >SC317945 representing NM_001453.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GCTCGTTTAGTGAACCGTCAGAATTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

ATGCAGGCGCGCTACTCCGTGTCCAGCCCCAACTCCCTGGGAGTGGTGCCCTACCTCGGCGGCGAGCAG AGCTACTACCGCGCGGCGGCGGCGGCGGCGGGGGGGCGGCTACACCGCCATGCCGGCCCCATGAGCGTG TACTCGCACCCTGCGCACGCCGAGCAGTACCCGGGCGGCATGGCCCGCGCCTACGGGCCCTACACGCCG CAGCCGCAGCCCAAGGACATGGTGAAGCCGCCCTATAGCTACATCGCGCTCATCACCATGGCCATCCAG AACGCCCCGGACAAGAAGATCACCCTGAACGGCATCTACCAGTTCATCATGGACCGCTTCCCCTTCTAC CGGGACAACAAGCAGGGCTGGCAGAACAGCATCCGCCACAACCTCTCGCTCAACGAGTGCTTCGTCAAG GTGCCGCGCGACGACAAGAAGCCGGGCAAGGGCAGCTACTGGACGCTGGACCCGGACTCCTACAACATG TTCGAGAACGGCAGCTTCCTGCGGCGGCGGCGCGCTTCAAGAAGAAGGACGCGGTGAAGGACAAGGAG CAGGCCGACGGCAACGCCCCGGTCCGCAGCCGCCGCCGTGCGCATCCAGGACATCAAGACCGAGAAC GGTACGTGCCCCTCGCCGCCCCAGCCCCTGTCCCCGGCCGCCGCCTGGGCAGCGGCAGCGCCGCCGC GTGCCCAAGATCGAGAGCCCCGACAGCAGCAGCAGCCTGTCCAGCGGGAGCAGCCCCCCGGGCAGC CCGCCGCCGCACCATAGCCAGGGCTTCAGCGTGGACAACATCATGACGTCGCTGCGGGGGTCGCCGCAG AGCGCGGCCGCGGAGCTCAGCTCCGGCCTTCTGGCCTCGGCGGCGCGTCCTCGCCGCGCGCGGATCGCA CCCCGCTGGCGCTCGGCCTACTCGCCCGGCCAGAGCTCCCTCTACAGCTCCCCCTGCAGCCAGACC ACCTACCACTGCAACCTGCAAGCCATGAGCCTGTACGCGGCCGGGCGAGCGCGGGGGCCACTTGCAGGGC GCGCCCGGGGGCGCGGCCGCCGTGGACGACCCCCTGCCCGACTACTCTCTGCCTCCGGTCACC AGCAGCAGCTCGTCCCTGAGTCACGGCGGCGGCGGCGGCGGCGGGGGGGAGGCCAGGAGGCCGGC CACTTGGCGAGCGCGGCGGCGGCGGCGGCGCCGCAGGCTACCCGGGCCAGCAGCAGAACTTCCACTCG GTGCGGGAGATGTTCGAGTCACAGAGGATCGGCTTGAACAACTCTCCAGTGAACGGGAATAGTAGCTGT CAAATGGCCTTCCCTTCCAGCCAGTCTCTGTACCGCACGTCCGGAGCTTTCGTCTACGACTGTAGCAAG **TTTTGA**

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC

Restriction Sites: Sgfl-Mlul

ACCN: NM_001453

Insert Size: 1662 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:

- 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: <u>NM 001453.2</u>

 RefSeq Size:
 3452 bp

 RefSeq ORF:
 1662 bp

 Locus ID:
 2296

 UniProt ID:
 Q12948

 Cytogenetics:
 6p25.3

Protein Families: Druggable Genome, Transcription Factors

MW: 56.8 kDa

Gene Summary: This gene belongs to the forkhead family of transcription factors which is characterized by a

distinct DNA-binding forkhead domain. The specific function of this gene has not yet been determined; however, it has been shown to play a role in the regulation of embryonic and ocular development. Mutations in this gene cause various glaucoma phenotypes including primary congenital glaucoma, autosomal dominant iridogoniodysgenesis anomaly, and

Axenfeld-Rieger anomaly. [provided by RefSeq, Jul 2008]