

Product datasheet for SC325534

OriGene Technologies, Inc.

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STRA6 (NM_001142620) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: STRA6 (NM_001142620) Human Untagged Clone

Tag: Tag Free Symbol: STRA6

Synonyms: MCOPCB8; MCOPS9; PP14296

Vector: pCMV6 series

Fully Sequenced ORF: >NCBI ORF sequence for NM_001142620, the custom clone sequence may differ by one or

more nucleotides

Restriction Sites: Please inquire ACCN: NM_001142620

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.

RefSeg: NM 001142620.1, NP 001136092.1

 RefSeq Size:
 1956 bp

 RefSeq ORF:
 480 bp

 Locus ID:
 64220

 UniProt ID:
 Q9BX79

 Cytogenetics:
 15q24.1

Protein Families: Transmembrane

Gene Summary: The protein encoded by this gene is a membrane protein involved in the metabolism of

retinol. The encoded protein acts as a receptor for retinol/retinol binding protein complexes.

This protein removes the retinol from the complex and transports it across the cell

membrane. Defects in this gene are a cause of syndromic microphthalmia type 9 (MCOPS9). Several transcript variants encoding a few different isoforms have been found for this gene.

[provided by RefSeq, Dec 2008]

Transcript Variant: This variant (5) differs in the 3' UTR and coding sequence compared to variant 8. The resulting isoform (c) has a shorter and distinct C-terminus compared to isoform

f.