

## **Product datasheet for SC332509**

## OriGene Technologies, Inc.

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## TSGA14 (CEP41) (NM\_001257159) Human Untagged Clone

## **Product data:**

**Product Type:** Expression Plasmids

**Product Name:** TSGA14 (CEP41) (NM\_001257159) Human Untagged Clone

Tag: Tag Free
Symbol: TSGA14

Synonyms: JBTS15; TSGA14

**Vector:** pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC332509 representing NM\_001257159.

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

GGTCACCTGCAAGGCAAACCCTGGAAGTAA

**Restriction Sites:** Sgfl-Mlul

**ACCN:** NM 001257159

**Insert Size:** 858 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).

**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 001257159.1

 RefSeq Size:
 6222 bp

 RefSeq ORF:
 858 bp

 Locus ID:
 95681

 UniProt ID:
 Q9BYV8

 Cytogenetics:
 7q32.2

**Protein Families:** Druggable Genome

MW: 31.6 kDa

**Gene Summary:** This gene encodes a centrosomal and microtubule-binding protein which is predicted to have

two coiled-coil domains and a rhodanese domain. In human retinal pigment epithelial cells the protein localized to centrioles and cilia. Mutations in this gene have been associated with

Joubert Syndrome 15; an autosomal recessive ciliopathy and neurological disorder.

Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2012] Transcript Variant: This variant (3) lacks two in-frame exons in the coding region, compared to variant 1, which results in a shorter isoform (3), compared to isoform 1. 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.