

## **Product datasheet for SC202023**

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## CEP290 (NM\_025114) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: CEP290 (NM\_025114) Human 3' UTR Clone

Symbol: CEP290

Synonyms: 3H11Ag; BBS14; CT87; JBTS5; LCA10; MKS4; NPHP6; POC3; rd16; SLSN6

**Mammalian Cell** 

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_025114

**Insert Size:** 198 bp

Insert Sequence: >SC202023 3'UTR clone of NM\_025114

The sequence shown below is from the reference sequence of NM\_025114. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GTACCTTTATACTTAGATTGGAATTCTTAATAAATAAAATTATATGAAATTTTCAACTTA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

**RefSeq:** <u>NM 025114.4</u>





## CEP290 (NM\_025114) Human 3' UTR Clone - SC202023

Summary: This gene encodes a protein with 13 putative coiled-coil domains, a region with homology to

SMC chromosome segregation ATPases, six KID motifs, three tropomyosin homology domains and an ATP/GTP binding site motif A. The protein is localized to the centrosome and cilia and has sites for N-glycosylation, tyrosine sulfation, phosphorylation, N-myristoylation,

and amidation. Mutations in this gene have been associated with Joubert syndrome and nephronophthisis and the presence of antibodies against this protein is associated with

several forms of cancer. [provided by RefSeq, Jul 2008]

**Locus ID:** 80184

**MW:** 7.9