

## **Product datasheet for SC206144**

## Pericentrin (PCNT) (NM\_006031) Human 3' UTR Clone

**Product data:** 

Product Type: 3' UTR Clones

Symbol: Pericentrin

Synonyms: KEN; MOPD2; PCN; PCNT2; PCNTB; PCTN2; SCKL4

Mammalian Cell Neomycin

Selection:

Vector: pMirTarget (PS100062)

ACCN: NM\_006031

Insert Size: 447 bp

Insert Sequence: >SC206144 3' UTR clone of NM\_006031

The sequence shown below is from the reference sequence of NM\_006031. The complete sequence of this clone may contain minor differences, such as SNPs. Red=Cloning site Blue=Stop Codon

CAATTGGCAGAGCTCAGAATTCAAGCGATCGC

ACAAAACAACTCAAAAAGGAATAAAAT

**ACGCGT**AAGCGGCCGCGGCATCTAGATTCGAAGAAAATGACCG

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).



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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.

Note: Plasmids are not sterile. For experiments where strict sterility is required, filtration with 0.22um

filter is required.

**RefSeq:** <u>NM\_006031.5</u>

Summary: The protein encoded by this gene binds to calmodulin and is expressed in the centrosome. It is

an integral component of the pericentriolar material (PCM). The protein contains a series of coiled-coil domains and a highly conserved PCM targeting motif called the PACT domain near its C-terminus. The protein interacts with the microtubule nucleation component gammatubulin and is likely important to normal functioning of the centrosomes, cytoskeleton, and cell-cycle progression. Mutations in this gene cause Seckel syndrome-4 and microcephalic osteodysplastic primordial dwarfism type II. Two transcript variants encoding different

isoforms have been found for this gene. [provided by RefSeq, Oct 2015]

**Locus ID:** 5116