

## Product datasheet for **SC332037**

### CEP57 (NM\_001243777) Human Untagged Clone

#### Product data:

**Product Type:** Expression Plasmids  
**Product Name:** CEP57 (NM\_001243777) Human Untagged Clone  
**Tag:** Tag Free  
**Symbol:** CEP57  
**Synonyms:** MVA2; PIG8; TSP57  
**Vector:** pCMV6-Entry (PS100001)  
**Fully Sequenced ORF:** >SC332037 representing NM\_001243777.  
Blue=Insert sequence Red=Cloning site Green=Tag(s)

```
ATGGCGGCGCGTCTGTCTCTGCGGCTTCTGGTTCTCACTTGTCGAACAGCTTTGCTGAGCCATCAAGG
TCTAATGGAAGCATGGTTCGGCATTCTTCATCTCCATATGTAGTATATCCTTCGGATAAGCCTTTCCTT
AATAGTGATCTACGACGCTCCCCAAGTAAGCCTACACTTGCCATCCAGAAAGCAACAGCAGAGCCATA
TTTTCTGCTCTTAAGAATCTTCAAGATAAGATTGACGCTTGGAACTTGAGAGGATTCAGGCAGAAGAA
AGTGTGAAAACCTTGCTAGAGAAAACAATTGAATATAAGAAAGTACTGGATGAACAGATACAAGAAAGG
GAGAATCAAAGAATGAGGAATCAAAGCACAATCAAGAAGTACATCTCAGTTGTTAGCTGCAGAAAAT
AAATGCAATCTATTAGAAAAACAATTGGAATACATGCGAAATATGATAAAGCATGCCGAAATGGAGAGG
ACATCTGTCTTAGAGAAAACAAGTTTCCCTAGAAAAGAGAACGACAACATGATCAAACACATGTTTCAGAGC
CAACTTGAAAAATTGGATCTTCTTGAACAGGAGTATAACAACTTACCACAATGCAGGCCCTTGCAGAA
AAAAAATGCAAGAGTTGGAAGCAAACTCCATGAAGAAGAACAGGAAAGGAAACGCATGCAAGCTAAG
GCAGCTGAGTTGCAGACTGGTCTAGAAAACAATAGACTTATCTTTGAAGATAAGGCAACTCCGTGTGTT
CCCAATGCAAGAAGAATTAAGAAAAAGAGTCAAAACCACCAGAAAAGTCCACAAGCCCTAGCCATGCC
GTGGTAGCCAATGTTTCAGCTTGTCTTGCATCTAATGAAGCAACACAGTAAAGCTTTGTGCAATGATCGA
GTCATCAACAGTATTCCTTTGGCAAAGCAAGTATCTTCACGAGGTGGTAAAAGTAAAGATTGTCAGTA
ACACCTCCCTCCCAACGGTATTAATGAGGAGTTGTCAGAAAGTCTTACAGACTTTACAGGATGAATTT
GGGCAAATGAGCTTTGATCACCAGCAGCTTGCAAACTTATCCAGGAGTCGCAACCGTTGAACTGAAA
GACAAGTTGGAGTGTGAATTGGAGGCATTAGTGGGAAGGATGGAAGCAAAAGCCAACCAATAACTAAA
GTTTCGAAAATACCAAGCCAGCTGGAGAAACAGAAGTTAGAGAAGCAGAAGAAGGAATTAAGGCTACC
AAAAAGACTCTTGATGAAGAAAGAAACAGCAGCAGCCGTTCTGGAATCACAGGGACCACAATAAGAAA
GATTTTATGAAACTGAGACCTGGAGAAAAAGGAGAAAAAATCTTCAGTTATTGAAGGACATGCAAAGC
ATACAGAATTCATTACAAAGCAGTAGTTTGTGTTGGGATTACTGA
```

**Restriction Sites:** SgfI-MluI  
**ACCN:** NM\_001243777  
**Insert Size:** 1425 bp



[View online »](#)

<b>OTI Disclaimer:</b>	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).
<b>Components:</b>	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).
<b>Reconstitution Method:</b>	<ol style="list-style-type: none"><li>1. Centrifuge at 5,000xg for 5min.</li><li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li><li>3. Close the tube and incubate for 10 minutes at room temperature.</li><li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li><li>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.</li></ol>
<b>RefSeq:</b>	<a href="#">NM_001243777.1</a>
<b>RefSeq Size:</b>	3114 bp
<b>RefSeq ORF:</b>	1425 bp
<b>Locus ID:</b>	9702
<b>UniProt ID:</b>	<a href="#">Q86XR8</a>
<b>Cytogenetics:</b>	11q21
<b>MW:</b>	54.2 kDa
<b>Gene Summary:</b>	<p>This gene encodes a cytoplasmic protein called Transloklin. This protein localizes to the centrosome and has a function in microtubular stabilization. The N-terminal half of this protein is required for its centrosome localization and for its multimerization, and the C-terminal half is required for nucleating, bundling and anchoring microtubules to the centrosomes. This protein specifically interacts with fibroblast growth factor 2 (FGF2), sorting nexin 6, Ran-binding protein M and the kinesins KIF3A and KIF3B, and thus mediates the nuclear translocation and mitogenic activity of the FGF2. It also interacts with cyclin D1 and controls nucleocytoplasmic distribution of the cyclin D1 in quiescent cells. This protein is crucial for maintaining correct chromosomal number during cell division. Mutations in this gene cause mosaic variegated aneuploidy syndrome, a rare autosomal recessive disorder. Multiple alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq, Aug 2011]</p> <p>Transcript Variant: This variant (3) lacks an in-frame exon in the 3' coding region, compared to variant 1. The resulting isoform (c) lacks an internal segment, compared to isoform a.</p>