

## Product datasheet for SC206071

## OriGene Technologies, Inc.

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## Phospholipase C epsilon 1 (PLCE1) (NM\_001165979) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: Phospholipase C epsilon 1 (PLCE1) (NM\_001165979) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: PLCE1

Synonyms: NPHS3; PLCE; PPLC

**ACCN:** NM\_001165979

**Insert Size:** 492 bp

Insert Sequence: >SC206071 3'UTR clone of NM\_001165979

The sequence shown below is from the reference sequence of NM\_001165979. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

AAAAAAAA

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:** NM 001165979.2





## Phospholipase C epsilon 1 (PLCE1) (NM\_001165979) Human 3' UTR Clone - SC206071

**Summary:** 

This gene encodes a phospholipase enzyme that catalyzes the hydrolysis of phosphatidylinositol-4,5-bisphosphate to generate two second messengers: inositol 1,4,5-triphosphate (IP3) and diacylglycerol (DAG). These second messengers subsequently regulate various processes affecting cell growth, differentiation, and gene expression. This enzyme is regulated by small monomeric GTPases of the Ras and Rho families and by heterotrimeric G proteins. In addition to its phospholipase C catalytic activity, this enzyme has an N-terminal domain with guanine nucleotide exchange (GEF) activity. Mutations in this gene cause early-onset nephrotic syndrome; characterized by proteinuria, edema, and diffuse mesangial sclerosis or focal and segmental glomerulosclerosis. Alternative splicing results in multiple transcript variants encoding distinct isoforms.[provided by RefSeq, Sep 2009]

**Locus ID:** 51196

**MW:** 19