

Product datasheet for SC201544

datashoot for SC201E4

SERPINB6 (NM_004568) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: SERPINB6 (NM_004568) Human 3' UTR Clone

Symbol: SERPINB6

Synonyms: CAP; DFNB91; MSTP057; PI-6; PI6; PTI; SPI3

Mammalian Cell

Selection:

Neomycin

Vector: pMirTarget (PS100062)

ACCN: NM_004568

Insert Size: 169 bp

Insert Sequence: >SC201544 3'UTR clone of NM_004568

The sequence shown below is from the reference sequence of NM_004568. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

 $\tt CTCTTCTGCGGCCGCTTTTCCTCTCGTGAGGACAGGGCAGTCTTGGTGTGCAGCCCCTCTCCTCTGTGCCCTGACACTCCACAGTGTGCCTGCAACCCCAAGTGGCCTTATCCGTGCAGTGGCAGTTCAGAAA$

TAAAGGCCCATTTGTGGGATGCCGCATTCA

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 004568.6</u>



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



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Summary: The protein encoded by this gene is a member of the serpin (serine proteinase inhibitor)

superfamily, and ovalbumin(ov)-serpin subfamily. It was originally discovered as a placental thrombin inhibitor. The mouse homolog was found to be expressed in the hair cells of the inner ear. Mutations in this gene are associated with nonsyndromic progressive hearing loss, suggesting that this serpin plays an important role in the inner ear in the protection against leakage of lysosomal content during stress, and that loss of this protection results in cell death and sensorineural hearing loss. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Sep 2010]

Locus ID: 5269 **MW:** 5.9