

Product datasheet for **SC320101**

RPL13 (NM_000977) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: RPL13 (NM_000977) Human Untagged Clone
Tag: Tag Free
Symbol: RPL13
Synonyms: BBC1; D16S44E; D16S444E; L13; SEMDIST
Mammalian Cell Selection: None
Vector: [pCMV6-XL5](#)
E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene sequence for NM_000977.2
 CTTTTCGCTCGGCTGTTTTCTGCGCAGGAGCCGAGGGCCGTAGGCAGCCATGGCGCCC
 AGCCGGAATGGCATGGTCTTGAAGCCCCACTTCCACAAGGACTGGCAGCGCGCGTGGCC
 ACGTGGTTCAACCAGCCGCCCCGTAAGATCCGCAGACGTAAGGCCCGCAAGCCAAGGCG
 CGCCGCATCGCCCCGCGCCCCGCGTGGGTCCCATCCGGCCCATCGTGGCTGCACCCAG
 GTTCGGTACCACGAAGGTGCGCGCCGCGCGCTTACGCTGGAGGAGCTCAGGGTG
 GCCGGCATTACAGAAGGTGGCCCGGACCATCGGCATTTCTGTGGATCCGAGGAGGCGG
 AACAAAGTCCACGGAGTCCCTGCAGGCCAACGTGCAGCGGCTGAAGGAGTACCGCTCCAAA
 CTCATCCTTTCCCAAGGAGCCCTCAGCCCCAAGAAGGGAGACAGTTCTGCTGAAGAA
 CTGAAACTGGCCACCCAGCTGACCGGACCGGTGATGCCCGTCCGGAACGTATATAAGAAG
 GAGAAAGCTCGAGTCATCACTGAGGAAGAGAAGAAATTTCAAAGCCTTCGCTAGTCTCCGT
 ATGGCCCGTGCCAACGCCCGGCTTTCCGCATACGGGCAAAAAGAGCCAAGGAAGCCGCA
 GAACAGGATGTTGAAAAGAAAAATAAAGCCCTCCTGGGGACTTGGAATCAGTCGGCAGT
 CATGCTGGGTCTCCACGTGGTGTGTTTCGTGGGAACAACCTGGGCCTGGGATGGGGCTTCA
 CTGCTGTGACTTCTCCTGCCAGGGGATTTGGGGCTTTCTTGAAGACAGTCCAAGCCCT
 GGATAATGCTTTACTTTCTGTGTTGAAGCACTGTTGGTTGTTGGTTAGTGACTGATGTA
 AAACGGTTTTCTGTGGGGAGGTTACAGAGGCTGACTTCAGAGTGGACTTGTGTTTTTTTC
 TTTTTAAAGAGGCAAGGTTGGGCTGGTGTCTCACAGCTGTAATCCCAGCACTTTGAGGTTG
 GCTGGGAGTTCAAGACCAGCCTGGCCAACATGTCAGAACTACTAAAAATAAAGAAATCAG
 CCATGAAAAAAAAAAAAAAAAA

Restriction Sites: Please inquire

ACCN: NM_000977



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OTI Disclaimer:	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).
OTI Annotation:	This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.
Components:	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).
Reconstitution Method:	<ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.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.
RefSeq:	NM_000977.2 , NP_000968.2
RefSeq Size:	1110 bp
RefSeq ORF:	636 bp
Locus ID:	6137
UniProt ID:	P26373
Cytogenetics:	17p11.2
Domains:	Ribosomal_L13e
Protein Families:	Druggable Genome
Protein Pathways:	Ribosome
Gene Summary:	<p>Ribosomes, the organelles that catalyze protein synthesis, consist of a small 40S subunit and a large 60S subunit. Together these subunits are composed of 4 RNA species and approximately 80 structurally distinct proteins. This gene encodes a ribosomal protein that is a component of the 60S subunit. The protein belongs to the L13E family of ribosomal proteins. It is located in the cytoplasm. This gene is expressed at significantly higher levels in benign breast lesions than in breast carcinomas. Alternatively spliced transcript variants encoding distinct isoforms have been found for this gene. As is typical for genes encoding ribosomal proteins, there are multiple processed pseudogenes of this gene dispersed through the genome. [provided by RefSeq, Jul 2011]</p> <p>Transcript Variant: This variant (1) is the predominant transcript. It encodes the longest isoform (1). Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.</p>