

Product datasheet for **SC118048**

UBE2G2 (NM_003343) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	UBE2G2 (NM_003343) Human Untagged Clone
Tag:	Tag Free
Symbol:	UBE2G2
Synonyms:	UBC7
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>OriGene ORF within SC118048 sequence for NM_003343 edited (data generated by NextGen Sequencing)

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ATGGCGGGGACCGCGCTCAAGAGGCTGATGGCCGAGTACAACAATTAACACTGAATCCT
CCGGAAGGAATTGTAGCAGGCCCATGAATGAAGAGAACTTTTTGAATGGGAGGCATTG
ATCATGGGCCAGAAAGACACCTGCTTTGAGTTTGGTGTTTTTCTGCCATCCTGAGTTT
CCAATTGATTACCCGTTAAGTCCCCAAAGATGAGATTTACCTGTGAGATGTTTCATCCC
AACATCTACCCTGATGGGAGAGTCTGCATTTCCATCCTCCACGCGCCAGGCGATGACCCC
ATGGGCTACGAGAGCAGCGGGAGCGGTGGAGTCTGTGCAGAGTGTGGAGAAGATCCTG
CTGTGGTGGTGGAGCATGCTGGCAGAGCCCAATGACGAAAGTGGAGCTAACGTGGATGCG
TCCAAAATGTGGCGCATGACCGGGAGCAGTTCTATAAGATTGCCAAGCAGATCGTCCAG
AAGTCTCTGGGACTGTGA
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Clone variation with respect to NM_003343.5



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5' Read Nucleotide Sequence:	>OriGene 5' read for NM_003343 unedited GGTTCAAATGTATACGACTCCTATAGGCGGCCGCGCAATTCGCACGAGGCTCGGCGCA GCTGTTGCGGGCCATGGCGGGGACCGCCTCAAGAGGCTGATGGCCGAGTACAAACAAT TAACACTGAATCCTCCGGAAGGAATTGTAGCAGGCCCATGAATGAAGAGAACTTTTTTG AATGGGAGGCATTGATCATGGGCCAGAAAGACACCTGCTTTGAGTTTGGTGTTCCTG CCATCCTGAGTTCCCACTTGATTACCCGTTAAGTCCCCAAAGATGAGATTTACCTGTG AGATGTTTCATCCCAACATCTACCCTGATGGGAGAGTCTGCATTTCCATCCTCCACGCGC CAGGCGATGACCCATGGGCTACGAGAGCAGCGGGAGCGGTGGAGTCTGTGCAGAGTG TGGAGAAGATCCTGCTGTCGGTGGTGAGCATGCTGGCAGAGCCCAATGACGAAAGTGGAG CTAACGTGGATGCGTCCAAAATGTGGCGGATGACCGGGAGCAGTTCTATAAGATTGCCA AGCAGATCGTCCAGAAGTCTCTGGGACTGTGAGACCTGGCCTCGCACAGGCGGCACACA CCGCCAAGCAGCTCAGCATTCTCCCCGGCACACTTAGTGACAGTGATGCTCTGTGCTGG TACCAAACAAGGCAGACTTGCAAGAACCATGGCATTTTTTTTTTTTCAAACCTTTCCT ACTTCAAACAGGCTTCTTCTGAAATGATGACTTAATGTGGAATATTGACAGCTTACTG CAGTTTTACAGTATTCTCACAAAGGGCTTCAGGTAGATTATCAGAGCTTTAGCACTAC CTCTNCCCGCTGAAACCAGCAGTTCATGGCCTTCTGTGGATTTCTCCTCCTGNAGTGT GAAGGGGGTT
Restriction Sites:	NotI-NotI
ACCN:	NM_003343
Insert Size:	3730 bp
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).
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_003343.4 , NP_003334.2
RefSeq Size:	2919 bp
RefSeq ORF:	498 bp
Locus ID:	7327
UniProt ID:	P60604
Cytogenetics:	21q22.3
Domains:	UBCc
Protein Families:	Druggable Genome

Protein Pathways: Parkinson's disease, Ubiquitin mediated proteolysis

Gene Summary: The modification of proteins with ubiquitin is an important cellular mechanism for targeting abnormal or short-lived proteins for degradation. Ubiquitination involves at least three classes of enzymes: ubiquitin-activating enzymes, or E1s, ubiquitin-conjugating enzymes, or E2s, and ubiquitin-protein ligases, or E3s. This gene encodes a member of the E2 ubiquitin-conjugating enzyme family. The encoded protein shares 100% sequence identity with the mouse counterpart. This gene is ubiquitously expressed, with high expression seen in adult muscle. Three alternatively spliced transcript variants encoding distinct isoforms have been found for this gene. [provided by RefSeq, Jan 2011]

Transcript Variant: This variant (1) 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.