

Product datasheet for **SC320881**

DCUN1D1 (NM_020640) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: DCUN1D1 (NM_020640) Human Untagged Clone
Tag: Tag Free
Symbol: DCUN1D1
Synonyms: DCNL1; DCUN1L1; RP42; SCCRO; SCRO; Tes3
Mammalian Cell Selection: Neomycin
Vector: pCMV6-AC (PS100020)
E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene sequence for NM_020640.2
GGAGGAGGAGGGAGAGGCTGGAGGACACCAACATGAACAAGTTGAAATCATCGCAGAA
GGATAAAGTTCGTCAGTTTATGATCTTCACACAATCTAGTGAAAAACAGCAGTAAGTTG
TCTTTCTCAAAATGACTGGAAGTTAGATGTTGCAACAGATAATTTTTTCCAAAATCCTGA
ACTTTATATACGAGAGAGTGTAAGGATCATTGGACAGGAAGAAGTTAGAACAGCTGTA
CAATAGATACAAAGACCCTCAAGATGAGAATAAAATTGGAATAGATGGCATAACAGCAGTT
CTGTGATGACCTGGCACTCGATCCAGCCAGCATTAGTGTGTTGATTATTGCATGGAAGTT
CAGAGCAGCAACACAGTGCAGTTCTCCAACAGGAGTTCATGGATGGCATGACAGAATT
AGGATGTGACAGCATAGAAAACTAAAGGCCAGATACCCAAGATGGAACAAGAATTGAA
AGAACCAGGACGATTTAAGGATTTTACCAGTTTACTTTAATTTTGCAAAGAATCCAGG
ACAAAAAGGATTAGATCTAGAAATGGCCATTGCCTACTGGAACCTTAGTGCTTAATGGAAG
ATTTAAATTCCTTAGACTTATGGAATAAATTTTTGTTGGAACATCATAAACGATCAATACC
AAAAGACACTTGGAAATCTCTTTTAGACTTCAGTACGATGATTGCAGATGACATGTCTAA
TTATGATGAAGAAGGAGCATGGCCTGTTCTTATTGATGACTTTGTGGAATTTGCACGCC
TCAAAATGCTGGGACAAAAAGTACAACAGTGTAGCACTAAAGGAACCTTCTAGAATGTAC
ATAGTCTGTACAATAAATACAACAGAAAATTGCACAGTCAATTTCTGCTGGCTGGACTGA
ACTGAAGATCAATCCTCACAATTCAGACTGAGGGTTGAGACAAAACCTTAAAGGATACATC
TTGGACCATATCGTATTTTCATTTCTAATGGTGGTTTGGGCTTGTCTTAGTCTGGC
CGCTCTAAACATTTATAATTCCAACATTGTGATTTTCATCTTATATCTGTGGACCATCCT
AGTTTATTCTCCATAAGTCTTAGAAGCTTTATGGTGATTATTTTGAGGTTTTTCATTCTC
GCATAAAGCACAATGCTGTCTTCATCAGAAAACAGTTGGCATAAGAATTAACATATGAA
CATCACAACAATTTATAAAAAAAAAAAAAAAAAA

Restriction Sites: Please inquire
ACCN: NM_020640



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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_020640.2 , NP_065691.2
RefSeq Size:	3192 bp
RefSeq ORF:	780 bp
Locus ID:	54165
UniProt ID:	Q96GG9
Cytogenetics:	3q26.33
Domains:	DUF298
Protein Families:	Druggable Genome
Gene Summary:	<p>Part of an E3 ubiquitin ligase complex for neddylation. Promotes neddylation of cullin components of E3 cullin-RING ubiquitin ligase complexes. Acts by binding to cullin-RBX1 complexes in the cytoplasm and promoting their nuclear translocation, enhancing recruitment of E2-NEDD8 (UBE2M-NEDD8) thioester to the complex, and optimizing the orientation of proteins in the complex to allow efficient transfer of NEDD8 from the E2 to the cullin substrates. Involved in the release of inhibitory effects of CAND1 on cullin-RING ligase E3 complex assembly and activity (PubMed:25349211, PubMed:28581483). Acts also as an oncogene facilitating malignant transformation and carcinogenic progression (By similarity). [UniProtKB/Swiss-Prot Function]</p> <p>Transcript Variant: This variant (1) encodes the longer 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>