

# Product datasheet for SC116267

### CD28 (NM\_006139) Human Untagged Clone

#### **Product data:**

#### OriGene Technologies, Inc.

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Product Type:	Expression Plasmids
Product Name:	CD28 (NM_006139) Human Untagged Clone
Tag:	Tag Free
Symbol:	CD28
Synonyms:	Тр44
Mammalian Cell Selection:	None
Vector:	pCMV6-XL5
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_006139, the custom clone sequence may differ by one or more nucleotides

ATGCTCAGGCTGCTCTTGGCTCTCAACTTATTCCCTTCAATTCAAGTAACAGGAAACAAGATTTTGGTGA AGCAGTCGCCCATGCTTGTAGCGTACGACAATGCGGTCAACCTTAGCTGCAAGTATTCCTACAATCTCTT CTCAAGGGAGTTCCGGGCATCCCTTCACAAAGGACTGGATAGTGCTGTGGAAGTCTGTGTTGTATATGGG AATTACTCCCAGCAGCTTCAGGTTTACTCAAAAACGGGGTTCAACTGTGATGGGAAATTGGGCAATGAAT CAGTGACATTCTACCTCCAGAATTTGTATGTTAACCAAACAGATATTTACTTCTGCAAAATTGAAGTTAT GTATCCTCCTCCTTACCTAGACAATGAGAAGAGCAATGGAACCATTATCCATGTGAAAGGGAAACACCTT TGTCCAAGTCCCCTATTTCCCGGACCTTCTAAGCCCTTTTGGGTGGTGGTGGTGGTGGTGGAGGAGCACGCTCTG CTTGCTATAGCTTGCTAGTAACAGTGGCCTTTATTATTTTCTGGGTGAGGAGTAAGAGGAAGCAGCCTCT GCACAGTGACTACATGAACATGACCCCCGCCGCCCCGGGCCCACCCGCAAGCATTACCAGCCCTATGCC CCACCACGCGACTTCGCAGCCTATCGCTCCTGA

<b>Restriction Sites:</b>	Notl-Notl
ACCN:	NM_006139
Insert Size:	3600 bp

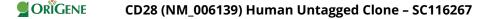


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## **CD28** (NM\_006139) Human Untagged Clone – SC116267

OTI Disclaimer:	Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at <u>custsupport@origene.com</u> or by calling 301.340.3188 option 3 for pricing and delivery. The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <u>More info</u>
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> <li>Centrifuge at 5,000xg for 5min.</li> <li>Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li> <li>Close the tube and incubate for 10 minutes at room temperature.</li> <li>Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li> <li>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.</li> </ol>
RefSeq:	<u>NM 006139.1, NP 006130.1</u>
RefSeq Size:	3804 bp
RefSeq ORF:	663 bp
Locus ID:	940
UniProt ID:	<u>P10747</u>
Cytogenetics:	2q33.2
Domains:	IGv
Protein Families:	Druggable Genome, Transmembrane
Protein Pathways:	Allograft rejection, Autoimmune thyroid disease, Cell adhesion molecules (CAMs), Graft- versus-host disease, Systemic lupus erythematosus, T cell receptor signaling pathway, Type I diabetes mellitus, Viral myocarditis

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Gene Summary:The protein encoded by this gene is essential for T-cell proliferation and survival, cytokine<br/>production, and T-helper type-2 development. Several alternatively spliced transcript variants<br/>encoding different isoforms have been found for this gene.[provided by RefSeq, Jul 2011]<br/>Transcript Variant: This variant (1) represents the predominant transcript, and encodes the<br/>longest isoform (1). Sequence Note: This RefSeq record was created from transcript and<br/>genomic sequence data to make the sequence consistent with the reference genome<br/>assembly. The genomic coordinates used for the transcript record were based on transcript<br/>alignments.

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