

Product datasheet for **SC310584**

PSME3 (NM_176863) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	PSME3 (NM_176863) Human Untagged Clone
Tag:	Tag Free
Symbol:	PSME3
Synonyms:	HEL-S-283; Ki; PA28-gamma; PA28G; PA28gamma; REG-GAMMA
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL4</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>OriGene ORF sequence for NM_176863 edited

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ATGGCCTCGTTGCTGAAGGTGGATCAGGAAGTGAAGCTCAAGGTTGATTCTTTCAGGGAG  
CGGATCACAAGTGAAGCAGAAGACTTGGTGGCAAATTTTTCCCAAAGAAGTTATTAGAA  
CTTGATAGTTTTCTGAAGGAACCAATCTTAAACATCCATGACCTAACTCAGATCCACTCT  
GACATGAATCTCCAGTCCCTGACCCCATCTTCTCACCAATAGCCATGATGGACTGGAT  
GGTCCCCTTATAAGAAGCGAAGGTTGGATGAGTGTGAAGAAGCCTTCCAAGGAACCAAG  
GTGTTTGTGATGCCCAATGGGATGCTGAAAAGCAACCAGCAGCTGGTGGACATTATTGAG  
AAAGTGAACCTGAGATCCGGCTGTTGATTGAGAAATGTAACACGCCTTCAGGCAAAGGT  
CCTCATATATGTTTTGACCTCCAGGTCAAATGTGGGTACAGCTCCTGATTCCCAGGATA  
GAAGATGGAACAACCTTTGGGGTGTCCATTGAGGAGAAACAGTTGCAGAGCTAAGAACT  
GTTGAGAGTGAAGCTGCATCTTATCTGGACCAGATTTCTAGATATTATATTACAAGAGCC  
AAATTGGTTTCTAAAATAGCTAAATATCCCATGTGGAGGACTATCGCCGCACCGTGACA  
GAGATTGATGAGAAAGAATATATCAGCCTTCGGCTCATATATCAGAGCTGAGGAATCAA  
TATGTCACTCTACATGACATGATCCTGAAAAATATCGAGAAGATCAAACGGCCCCGGAGC  
AGCAATGCAGAGACTCTGTACTGA
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5' Read Nucleotide Sequence:	<p>>OriGene 5' read for NM_176863 unedited TGCCTTTGTATACGACTCCTATAGGGCGGCCGCGTAATTCGGCACCAGGAGCGAGCGAGC AGTGAGTAAGCCAGCAAGGGCGGTCCGGTCCCAGGTCAGCCGAGATTTCTCAGGTCCCT CCGGCCCTCCCTGGAGTCCACAGCGCTCCGGTGTCCAGAGGATCGGACACGGCCCGG CCCGGCCATGGCTCGTTGCTGAAGGTGGATCAGGAAGTGAAGCTCAAGGTTGATCTTT CAGGGAGCGGATCACAAGTGAGGCAGAAGACTTGGTGGCAAATTTTTCCCAAAGAAGTT ATTAGAATTGATAGTTTTCTGAAGGAACCAATCTTAAACATCCATGACCTAACTCAGAT CCACTCTGACATGAATCTCCAGTCCCTGACCCATTCTTCTACCAATAGCCATGATGG ACTGGATGGTCCCACTTATAAGAAGCGAAGGTTGGATGAGTGTGAAGAAGCCTTCCAAGG AACCAAGTGTTTGTGATGCCAATGGGATGCTGAAAAGCAACCAGCAGCTGGTGGACAT TATTGAGAAAGTAAACCTGAGATCCGGCTGTTGATTGAGAAATGTAACACGCCTTCAGG CAAAGTCTCATATATGTTTTGACCTCCAGGTCAAAATGTGGGTACAGCTCCTGATTCC CAGGATAGAAGATGGAACAACCTTGGGGTGTCCATTGAGGAAACAGTTGCAGAGCT AAGAACTGTTGAGAGTGAAGCTGCATCTTATCTGGACCAGATTTCTAGATATTATATTAC AAGAGCCAAATTGTTTTCTAAATAGCTAAATATCCCATGTGGAGGACTATCGCCGACC GTGACAGAGATTGATGAGAAAGAATATATCAGCCCTTCGCTCATATATCAGAGCTGAGA TCAATATGTCACCTCTACATGAACATGATCCTGAAAAATTATCGGAGAGGATC</p>
3' Read Nucleotide Sequence:	<p>>Forward primer walk for NM_176863 unedited TGTTCGAAGAGAACAGGAAAATCCCAGGATAGAAGATGGAACAACCTTTGGGGTGTCCAT TCAGGAGGAAACAGTTGCAGAGCTAAGAAGTGTGAGAGTGAAGTGCATCTTATCTGGA CCAGATTTCTAGATATTATTACAAGAGCCAAATTTGGTTTCTAAATAGCTAAATATCC CCATGTGGAGGACTATCGCCGACCGTGACAGAGATTGATGAGAAAGAATATATCAGCCT TCGGCTCATCATATCAGAGCTGAGGAATCAATATGTCACCTCTACATGACATGATCCTGAA AAATATCGAGAAGATCAAACGGCCCGGAGCAGCAATGCAGAGACTCTGTACTGAGGCCA GGGCCAGGGCCAGGGGACTCTGTGAGTCTGGCTCAAGACCAGCATTGCCTTGGTTTGTTA CATGACTATCGTGATGGGAAACTGGCTGGAAATAGTAATCACACCTCTCTGTTTTTGT TAGAGTCTAATGAACTCTCATCTAGTTCTGTGATGTGTTTACCTCTTTTTTCAGGCCTC AGGAACTTTCTATTTCCCTTCCCTAATACCCACACCCAACTGTCGTAATTTCTGGAGA ACTCCAGGTTTGTGTGCAGGATGTTGGCACAATAACCTGTGTTTTTCATTCTCCCC TCTCTCCCTCCTGTGCTTTCGCTTTATGTTTTCTTCCGTTTGATAATTAGTTGGTTAA AGCTGAGGGAACCCGGAAGGAAAGTGTAGGTGTTTTTTAGGAACTAGGGTGGCGGGGG ACGAACTTCTCTTCTCATGAGGTTACTGTTTCTTCTCTGTTGGGCAATTGGATCC TCCCCAGTTGCCCTGGTGTGACTTAGGCTTCCAATCTGTGTACATCCCACTTTGAA TCTTGATCGGTGGACAAGAAAATAACCTTAGGCCTTTCAGTC</p>
Restriction Sites:	Please inquire
ACCN:	NM_176863
Insert Size:	2500 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).
OTI Annotation:	The open reading frame of this TrueClone was fully sequenced and found to be a perfect match to the protein associated to this reference.
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:

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_176863.1](#), [NP_789839.1](#)

RefSeq Size: 3228 bp

RefSeq ORF: 804 bp

Locus ID: 10197

UniProt ID: [P61289](#)

Cytogenetics: 17q21.31

Protein Families: Stem cell - Pluripotency

Protein Pathways: Antigen processing and presentation, Proteasome

Gene Summary: The 26S proteasome is a multicatalytic proteinase complex with a highly ordered structure composed of 2 complexes, a 20S core and a 19S regulator. The 20S core is composed of 4 rings of 28 non-identical subunits; 2 rings are composed of 7 alpha subunits and 2 rings are composed of 7 beta subunits. The 19S regulator is composed of a base, which contains 6 ATPase subunits and 2 non-ATPase subunits, and a lid, which contains up to 10 non-ATPase subunits. Proteasomes are distributed throughout eukaryotic cells at a high concentration and cleave peptides in an ATP/ubiquitin-dependent process in a non-lysosomal pathway. An essential function of a modified proteasome, the immunoproteasome, is the processing of class I MHC peptides. The immunoproteasome contains an alternate regulator, referred to as the 11S regulator or PA28, that replaces the 19S regulator. Three subunits (alpha, beta and gamma) of the 11S regulator have been identified. This gene encodes the gamma subunit of the 11S regulator. Six gamma subunits combine to form a homohexameric ring. Alternate splicing results in multiple transcript variants. [provided by RefSeq, May 2012]
Transcript Variant: This variant (2), uses an alternate in-frame splice site, compared to variant 1. The resulting isoform (2) is longer than 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.