

## Product datasheet for **SC331737**

### Hsp47 (SERPINH1) (NM\_001207014) Human Untagged Clone

#### Product data:

**Product Type:** Expression Plasmids  
**Product Name:** Hsp47 (SERPINH1) (NM\_001207014) Human Untagged Clone  
**Tag:** Tag Free  
**Symbol:** Hsp47  
**Synonyms:** AsTP3; CBP1; CBP2; gp46; HSP47; OI10; PIG14; PPROM; RA-A47; SERPINH2  
**Vector:** pCMV6-Entry (PS100001)  
**Fully Sequenced ORF:** >SC331737 representing NM\_001207014.  
 Blue=Insert sequence Red=Cloning site Green=Tag(s)

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ATGCGCTCCCTCCTGCTTCTCAGCGCCTTCTGCGCTCCTGGAGGCGGCCCTGGCCGCCGAGGTGAAGAAA
CCTGCAGCCGAGCAGCTCCTGGCACTGCGGAGAAGTTGAGCCCCAAGGCGGCCACGCTTGCCGAGCGC
AGCGCCGGCCTGGCCTTACGCTTGTACCAGGCCATGGCCAAGGACCAGGCAGTGGAGAACATCCTGGTG
TCACCCGTGGTGGTGGCCTCGTCGCTAGGGCTCGTGTGCTGGGCGGCAAGGCGACCACGGCGTCGCAG
GCCAAGGCAGTGTGAGCGCCGAGCAGCTGCGCGACGAGGAGGTGCACGCCGGCTGGGCGAGCTGCTG
CGTCACTCAGCAACTCCACGGCGCGCAACGTGACCTGGAAGCTGGGCGAGCCGACTGTACGGACCCAGC
TCAGTGAGCTTCGCTGATGACTTCGTGCGCAGCAGCAAGCAGCACTACAAGTGCAGCACTCCAAGATC
AACTTCCGCGACAAGCGCAGCGCCTGCACTCCATCAACGAGTGGGCGCGCAGACCACCGACGGCAAG
CTGCCCCGAGGTACCAAGGACGTGGAGCGCACGGACGGCGCCCTGCTAGTCAACGCCATGTTCTTCAAG
CCACACTGGGATGAGAAATTCACCACAAGATGGTGGACAACCGTGGCTTCATGGTGACTCGGTCCTAT
ACCGTGGGTGTCATGATGATGCACCGGACAGGCCCTTACAAGTACTACGACGACGAGAAGGAAAAGCTG
CAAATCGTGGAGATGCCCTGGCCACAAGCTCTCCAGCCTCATCATCCTCATGCCCATCACGTGGAG
CCTCTCGAGCGCCTTAAAAGCTGCTAACCAAAGAGCAGCTGAAGATCTGGATGGGAAGATGCAGAAG
AAGGCTGTTGCCATCTCCTTGCCCAAGGGTGTGGTGGAGGTGACCCATGACCTGCAGAAACACCTGGCT
GGGCTGGCCTGACTGAGGCCATTGACAAGAACAAGGCCGACTTGTACGCATGTCAGGCAAGAAGGAC
CTGTACCTGGCCAGCGTGTCCACGCCACCGCCTTTGAGTTGGACACAGATGGCAACCCCTTTGACCAG
GACATCTACGGGCGCGAGGAGCTGCGCAGCCCCAAGCTGTTCTACGCCGACCACCCCTTCTCTTCTTA
GTGCGGGACACCCAAAGCGGCTCCCTGCTATTTCATTGGGCGCCTGGTCCGGCCTAAGGGTGACAAGATG
CGAGACGAGTTATAG
  
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**Restriction Sites:** SgfI-MluI  
**ACCN:** NM\_001207014  
**Insert Size:** 1257 bp



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<b>OTI Disclaimer:</b>	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).
<b>Components:</b>	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).
<b>Reconstitution Method:</b>	<ol style="list-style-type: none"><li>1. Centrifuge at 5,000xg for 5min.</li><li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li><li>3. Close the tube and incubate for 10 minutes at room temperature.</li><li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li><li>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.</li></ol>
<b>RefSeq:</b>	<a href="#">NM_001207014.1</a>
<b>RefSeq Size:</b>	2333 bp
<b>RefSeq ORF:</b>	1257 bp
<b>Locus ID:</b>	871
<b>UniProt ID:</b>	<a href="#">P50454</a>
<b>Cytogenetics:</b>	11q13.5
<b>Protein Families:</b>	Druggable Genome
<b>MW:</b>	46.4 kDa
<b>Gene Summary:</b>	<p>This gene encodes a member of the serpin superfamily of serine proteinase inhibitors. The encoded protein is localized to the endoplasmic reticulum and plays a role in collagen biosynthesis as a collagen-specific molecular chaperone. Autoantibodies to the encoded protein have been found in patients with rheumatoid arthritis. Expression of this gene may be a marker for cancer, and nucleotide polymorphisms in this gene may be associated with preterm birth caused by preterm premature rupture of membranes. Alternatively spliced transcript variants have been observed for this gene, and a pseudogene of this gene is located on the short arm of chromosome 9. [provided by RefSeq, May 2011]</p> <p>Transcript Variant: This variant (1) represents the longer transcript. Variants 1 and 2 encode the same protein. 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>