

Product datasheet for **SC102514**

HARS2 (NM_012208) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	HARS2 (NM_012208) Human Untagged Clone
Tag:	Tag Free
Symbol:	HARS2
Synonyms:	HARSL; HARSR; HisRS; HO3; PRLTS2
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)



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Fully Sequenced ORF: >OriGene ORF within SC102514 sequence for NM_012208 edited (data generated by NextGen Sequencing)

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ATGCCCCGTGCTCGGACTTCTTCCCAGGAGGGCCTGGGCTTCGCTGCTCAGCCAGCTCCTG
CGACCCGCCTGCGCTTCGTGCACCGGGCGGTCCGTTGCCAAAGCCAGGTTGCAGAGGCA
GTGTTAACATCCCAACTGAAAGCACATCAAGAGAAACCAATTTTATTATCAAGACCCCA
AAGGGTACCAGGGATCTTAGTCTCAGCATATGGTTGTGAGGGAGAAAATCTTGATTTG
GTTATCAGCTGCTTTAAACGTCATGGAGCAAAGGGGATGGACACCCAGCATTGAGCTG
AAGGAAACCCTGACTGAGAAGTATGGGGAGGACTCTGGGCTCATGTATGATCTGAAGGAT
CAAGGTGGAGAGCTGTTGTCCTCCGCTATGACCTTACTGTTCCCTTTGCTCGTTATCTG
GCCATGAATAAGGTGAAGAAGATGAAACGTTATCATGTTGAAAGGTGTGGCGGCGAGAG
AGCCCAACCATAGTCCAAGGCCGTTATAGGGAGTTCTGCCAGTGTGATTTTGACATTGCT
GGTCAGTTTGACCCTATGATCCCCGATGCAGAGTGTGTTGAAGATCATGTGTGAAATCCTA
AGTGGATTGCAGTTGGGAGACTTCTCATTAAAGTAAATGACCGGCGGATTGTGGATGGG
ATGTTTGTCTGTGGTGTTCCTGAAAGCAAGTCCGTCGCATCTGCTCCTCCATAGAT
AAACTAGACAAGATGGCTTGAAAGATGTGAGACATGAGATGGTGGTGAAGAAAGCCCTG
GCTCCTGAGGTGGCTGATCGAATTGGGGACTATGTCCAGTGTGATGGTGGGGTATCCCTA
GTAGAGCAAATGTTTCAGGATCCCAGACTATCCCAGAACAAGCAGGCCCTGGAGGGCCTG
GGAGACCTAAAGCTGCTATTTGAATACCTGACTTTATTTGGAATTGCTGATAAGATCTCC
TTTGACCTCAGCCTGGCTCGGGCCTAGACTACTATACAGGAGTGATCTATGAAGCAGTG
CTGCTGCAGACCCCAACTCAGGCTGGGGAGGAGCCCTGAATGTGGGCAGTGTGGCTGCT
GGTGGGCGCTATGATGGGCTGGTGGCATGTTTGACCCCAAGGGCCACAAGGTGCCATGT
GTGGGACTCAGCATTGGGGTTGAGCGAATCTTCTACATTGTGGAGCAGAGGATGAAGACC
AAAGGTGAGAAGGTGCGGACTACAGAGACTCAAGTGTGTTGTGGCCACCCACAGAAGAAC
TTTCTCCAAGAACGGTTGAAGCTTATTGCAGAGCTTTGGGATTCTGGAATCAAGGCAGAG
ATGCTATACAAGAACAACCCCAAATTAACCCAGCTGCACTATTGTGAGAGCACAGGC
ATTCCACTGGTGGTCATTATTGGTGAAGAACTGAAAGAAGGGGTGATCAAGATCCGT
TCAGTGGCCAGCAGAGAGGAGGTGGCCATTAACGGGAAAATTTGTGGCTGAAATTCAG
AAGCGACTGTCTGAGTCTTGA
    
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Clone variation with respect to NM_012208.2
327 a=>g

5' Read Nucleotide Sequence:

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>OriGene 5' read for NM_012208 unedited
GCATTTTGAATACGACTCACTATAGGGCGGCCGGAATTCGGCACGAGGCTGGCTACTA
AGGGAACCTGGGAGGATCCCACCTCAGCCTTCGTGACTAGTGAGGTGCGCAAACGCCGA
GTTTTCCCTGGTGC GCGGGTCCGCCTTTGCAGTGCCCTCCACCCTTCCCTGGTGTCTGAC
CCGCCTCCTCCAGGCCTTTTGTTCCTGTCCCGAAAGCCGGCGTCTGCCGCGCGATG
CCCCTGCTCGGACTTCTCCAGGAGGGCCTGGGCTTCGCTGCTCAGCCAGCTCCTGCGA
CCGCCCTGCGCTTCGTGCACCGGGCGGTCCGTTGCCAAAGCCAGGTTGCAGAGGCAAGT
TTAACATCCCAACTGAAAGCACATCAAGAGAAACCAATTTTATTATCAAGACCCCAAAG
GGTACCAGGGATCTTAGTCTCAGCATATGGTTGTGAGGGAGAAAATCTTGATTTGGTT
ATCAGCTGCTTTAAACGTCATGGAGCAAAGGGGATGGACACCCAGCATTGAGCTGAAG
GAAACCCTGACTGAGAAGTATGGGGAGGACTCTGGGCTCATGTATGATCTGAAGGATCAA
GGTGGAGAGCTGTTGCCCTCCGCTATGACCTTACTGTTCCCTTTGCTCGTTATCTGGCC
ATGAATAAGGTGAAGAAGATGAAACGTTATCATGTTGAAAGGTGTGGCGGCGAGAGAGC
CCAACCATAGTCCAAGGCCGTTATAGGGAGTTCTGCCAGTGTGATTNTGACATTGCTGGT
CAGTTTGACCCTATGATCCCCGATGCAGAGTGGTTGAAGATCATGTGTGAAATCCTAAGT
GGATTGCAGTTGGGAGACTNTCTCATTAAANGGTAATGACCNGCNGATNGTGNATGGGGAT
GTTGCTGTCTGTGTGCTGAAAGCNAAGTNCCTGGCCATCTGCTCTCCTAGATAACTA
GACAGATGCTGGGAAGATG
    
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3' Read Nucleotide Sequence:	>OriGene 3' read for NM_012208 unedited NAAATTTTGGACTNTAGNCCCAGCGNCCCCTTCNANNGATCGGTTTTTTTTTTTTTTTT TTTGCACTTAATATTTTATTTAACATGTTATCCAACCACTTTCTAAAACAGCCTCCATGG CACGGTATGGAATAGAGCAGTGATTCTGGCACAGGAAATGGTCTCCAAAATCTCAAATA TCAGATCTCTACCTTGCAAGTCATCTATTAAGTTCAATGTTCAAGATTATCAAGCATA AGCCCCACTCTTCTACTGTAAGCAATGGATGGGCTAGAGAGATGTCAAATCAACATGGGT ATGCCATGCCCTTAGGATAGCATCTGTCCTCAAAGCTGAAAGCAGCCATTGCCCTCTT TGCTGTCTTCCAAAACTCCCAGACAGCCAATGGCTGATTGTGGACAAATTCACCAGA ATCTGGTTCCGTAGCTACAGTAGAAGGCTAAGTATCTGGTTTCTGATCTCAATGGTTCAG AATCCTGAGCCTGAGGTGACTCTCCAGAGCCTTGCTCACAGCAACAATCTCACTACCTC TTGACAACGTTCCGTGGCACTTCAAATTTGCCTCTGCAGCTAGAGCATGTCTTTTACA TCCAGCCATCTGCACCCATGCAGAATAGCTGCTCACACTAGCCCAGGAGTCTCAGGGTC TTCTACAGTTCTGCACCCAGGAAGAATGGAGGAGGCAGAAAGTACTTGTACCCTTCTG GCTAGTTGTTGTGAAGTCCATCACTCAATTCAGAGGAATTCAGTTCTAGAGAAAACCTT TTCTACAAGAGCAGCAGATGGGAATCANGCAAGGATCAAGACTCAGACAGTCGTTCTG AATTTTCAGCCACAAAATTTNCCCCTTAATGGCCCACTNCTCTCTGCTGGCCACTGNAACG GAT
Restriction Sites:	NotI-NotI
ACCN:	NM_012208
Insert Size:	2550 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_012208.2 , NP_036340.1
RefSeq Size:	2508 bp
RefSeq ORF:	1521 bp
Locus ID:	23438
UniProt ID:	P49590
Cytogenetics:	5q31.3
Domains:	tRNA-synt_2b, HGTP_anticonodon
Protein Pathways:	Aminoacyl-tRNA biosynthesis

Gene Summary:

Aminoacyl-tRNA synthetases are a class of enzymes that charge tRNAs with their cognate amino acids. The protein encoded by this gene is an enzyme belonging to the class II family of aminoacyl-tRNA synthetases. Functioning in the synthesis of histidyl-transfer RNA, the enzyme plays an accessory role in the regulation of protein biosynthesis. The gene is located in a head-to-head orientation with HARS on chromosome five, where the homologous genes likely share a bidirectional promoter. Mutations in this gene are associated with the pathogenesis of Perrault syndrome, which involves ovarian dysgenesis and sensorineural hearing loss. Alternative splicing results in multiple transcript variants of this gene. [provided by RefSeq, Jul 2013]

Transcript Variant: This variant (1) represents the longest transcript and encodes the longest isoform (1).