

Product datasheet for **SC107929**

Collagen IX (COL9A1) (NM_078485) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	Collagen IX (COL9A1) (NM_078485) Human Untagged Clone
Tag:	Tag Free
Symbol:	Collagen IX
Synonyms:	DJ149L1.1.2; EDM6; MED; STL4
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL4</u>
E. coli Selection:	Ampicillin (100 ug/mL)



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Fully Sequenced ORF: >OriGene ORF within SC107929 sequence for NM_078485 edited (data generated by NextGen Sequencing)

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ATGGCCTGGACTGCGCGGGACCGCGGGGCCCTGGGGCTGCTGCTGTTGGGGCTCTGCTTG
TGCGCGGCTCAAAGAGGTCCCCGGGTGAGCAGGGTCCTCCCGGGCTCCGGGCCCCCT
GGAGTTCCAGGCATCGATGGCATCGACGGTGACCGAGGTCTAAGGGCCCCCGGGCCCC
CCGGTCTGCAGGTGAACCGGAAAGCCAGGAGCTCCAGGCAAGCCTGGCACACCTGGC
GTTGATGGATTAACAGGACCTGATGGATCCCCTGGCTCCATTGGGTCAAAGGGACAAAA
GGAGAACCTGGTGTGCCTGGATCGCGTGGATTTCCAGGCCGTGGTATTCTGGACCCCT
GGTCTCTGCGGACAGCAGGACTCCCTGGAGAGCTTGGCCGTGTAGGACCTGTTGGTGAC
CCTGGGAGAAGAGGACCCTGGCCCCCTGGCCCCCAGGACCCAGAGGAACAATTGGC
TTTCATGATGGAGATCCATTGTGTCCCAATGCCTGTCCACCAGGTGCTCAGGATATCCA
GGCCTACCAGGCATGAGGGGTCAAAAGGGGCTAAAGGAGAAATTGGTGAACCAGGAAGA
CAAGGACACAAGGGTGAAGAAGGTGACCAGGGAGAACTCGGAGAAGTTGGAGCTCAAGGA
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CAAGGACAGCGAGGACCTCCAGGAGAAGCAGGTCCCAAAGGAGATAGAGGGGCTGAAGGT
GCTAGAGGAATTCTGGTCTCCCTGGGCCCAAAGGAGACACGGGTTTGCCAGGTGTGGAT
GGCCGTGATGGGATCCCTGGAATGCCTGGAACAAAGGGTGAACCAGGAAAACCTGGGCC
CCTGGTGTGAGGATTGAGGGGTTACCAGGTGTACCTGGAATTCTGGTGCAGGAGGGT
GTTGCTGGTGAAGGGTAGCACAGGTGCTCCAGGGAAGCCTGGTGCAGATGGGAAATTCA
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CCTGGCAGTAGAGGAGAATTAGGACCAGTGGGATCCCCAGGCCTACCAGGTAACCTGGT
TCTCTGGTAGCCCTGGCCTCCCTGGCTTGCCTGGGCCCTGGACTTCTGGAATGAAA
GGTGACAGGGGTGTAGTCGGTGAACCGGGTCCAAGGGTGAACAGGGTGCCTCTGGTGAA
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TCTGCAGGTAATCTGGGGAACCTGGCTTGAAGGGCCTGAGGGAAGTCGGGGGCTTCT
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ACCGGCCTGCCTGGTGTCCAGGGCCCTCCGGGTAGAGCACCAGCAGATCAGCACATTAAG
CAGGTTTGCATGAGATCATACAAGAACATTTTGTGAGATGGCTGCCAGTCTTAAGCGT
CCAGACTCAGGTGCCACTGGGCTTCTGGAAGGCCTGGCCCTCCTGGTCCCCCGGCCCT
CCTGGAGAGAATGGTTTCCAGGCCAGATGGGAATTCGTGGCCTTCCGGGATTAAGGGG
CCCCCTGGTGCTTGGTTTGGGGGACCTAAAGGTGACTTGGGAGAAAAGGGGGAGCGT
GGCCCTCCAGGAAGAGGTCCCAACGGTTTGCCTGGAGCTATAGGTCTCCAGGTGACCCA
GGCCCTGCCAGCTATGGCAGAAATGGCCGAGACGGTGAGCGAGGCCCCCCAGGGGTGGCA
GGAATTCCTGGAGTGCCTGGACCCCGGGACCTCCTGGGCTTCCCGGTTTCTGTGAGCCA
GCCTCTGCACCATGCAGGCTGGTCAGCGAGCATTTAACAAGGGCCTGACCCCTGA
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Clone variation with respect to NM_078485.3
1133 a=>g

5' Read Nucleotide Sequence:	<p>>OriGene 5' read for NM_078485 unedited TATACGACTCACTATAGGGCGGCCGCGATTTCGGCACGAGGGCCACCTATGCTAGTGGCGG GTCTGGAAGCCTAGAGGGGAACCAAGGCTGCAGAGCCGGGCAAGGGATTAGCGCGGGCG GCGGGCATGGCCTGGACTGCGCGGGACCGGGGCCCTGGGGCTGCTGCTGTTGGGGCTC TGCTTGTGCGCGGCTCAAAGAGGTCCCCGGGTGAGCAGGGTCTCCCGGGCCTCCGGGC CCCCCTGGAGTTCAGGCATCGATGGCATCGACGGTGACCGAGGTCTAAGCCCCCCCC GGGCCCCCGGGTCTGCAGGTGAACCGGAAAGCCAGGAGCTCCAGGCAAGCCTGGCAC ACCTGGCGCTGATGGATTAAACAGGACCTGATGGATCCCCTGGCTCCATTGGGTCAAAGGG ACAAAAAGGAGAACCTGGTGTGCCTGGATCGCGTGGATTTCAGGCCGTGGTATTCTCGG ACCCCCTGGTCTCTGGGACAGCAGGACTCCCTGGAGAGCTTGCCGTGTAGGACCTGG TTGGTGACCTGGGAGAAGAGGACCACCTGGNCCCCCTGGCCNCCAGGACCCAGAGGA ACAATTGGCTTTCATGATGGGAGATCCANTTGTGTCCCAATGCCTGTCCACCANGTCGCT CAGGATATCCAGGCTACNCGATGGGGGGTATAAGGGGCTAAAGGAGAAATTGGGTG ACCANGAAAGACAGGACACAAGGTGGAAGAAAGTGACCAGGGAGACTCCGAGAAGTTGGA GCTCAAGGACCTCCAGAGCCAGGGTTTGGCGAGCATCACGGCNATATTGGGGACAAGGG GAAAAAGGTGCTCGGGCTTTAAAGGGGGAACCTGGGCCTTAGGTCTTCTGGGGCCCTGG GGATCAAGGACGCGAGGACCTCCGAAAACAGGCCAAGGAAATAAAGGGCTTGAGGGCT TAAGAA</p>
3' Read Nucleotide Sequence:	<p>>OriGene 3' read for NM_078485 unedited GGTCCGACTATGNACCGGNCGCATNCTANGATCGNGTTTTTTTTTTTTTTTTTTTTTTT GTTAAGTAATGTTAATGTATCTAAATAAGCTCAGTAACCTCTTTTCATTGAACATGATT ATTTATAAGAAAACATGCAAGAGATTAAGAAACCATATTTATTTCTTTTGGTACAACC AGAGATGTTACCTCAAACCTTTAAAATAAATAATCTCATCAATAAAATTTATGAATGCCA GAGTATTAATAATTTCTTTGCTTTCACATAGAAGCGCTCAAAATATTAACAGGTTTGG GGTAAAGTACATACAGTTCAACAAAAGTTCAAGGTGAAGAAGGAAACCAATACATGACGT AAAGAAATCATTCGGGAGATGTAGCCCTTTCTTATCACTCCAGAGACAGCACATCGATG AGATTTTAAGCACCAATCCATTGAATGTAGTGGGGTTTCTTCATAACTAGTCTAGTAAT CCTCTATAACTGAAACAATAATCTGAGAACAACCTTCAATGAAAATAACTACTGCAA CTGAGGTAACCAACAGTTGTTTTTTTATTGGCACAGTGGCCACGATAAGACACAT AGGGTTAGCACACTAAAGCTCAAGATCCTGGGAACAGGAGTATAAAATTGTTCAAGGGAG GTGGTTGTTTTCTTTTTTTTTTTTAACTGATGACTCTGCTGTCTTCCCTCCCAGGGAA AGGAAAAAAAACCTACTGAATTAGCCCCCTCTATATGTGATTGGCCAGTAGGACTTC TGTAATCCTACTGGAGGGAAAACATTGTAAGCAGGCCGAAAGGAAACATCTTTGCCCCCA CTTTGGAAGGGGTTTTTAAACCAAGCTTCTTACCACAGGGTGGTTTATGGCCAACGGC TTGCGGCAGTAAACCTTTT</p>
Restriction Sites:	NotI-NotI
ACCN:	NM_078485
Insert Size:	2600 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:

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_078485.2](#), [NP_511040.2](#)

RefSeq Size: 2985 bp

RefSeq ORF: 2037 bp

Locus ID: 1297

UniProt ID: [P20849](#)

Cytogenetics: 6q13

Domains: Collagen

Protein Families: Druggable Genome

Gene Summary: This gene encodes one of the three alpha chains of type IX collagen, which is a minor (5-20%) collagen component of hyaline cartilage. Type IX collagen is usually found in tissues containing type II collagen, a fibrillar collagen. Studies in knockout mice have shown that synthesis of the alpha 1 chain is essential for assembly of type IX collagen molecules, a heterotrimeric molecule, and that lack of type IX collagen is associated with early onset osteoarthritis. Mutations in this gene are associated with osteoarthritis in humans, with multiple epiphyseal dysplasia, 6, a form of chondrodysplasia, and with Stickler syndrome, a disease characterized by ophthalmic, orofacial, articular, and auditory defects. Two transcript variants that encode different isoforms have been identified for this gene. [provided by RefSeq, Jul 2008]

Transcript Variant: This variant (2) differs in the 5' UTR and lacks several exons in the 5' coding region, compared to variant 1. These differences cause translation initiation at a unique AUG and an isoform (2, also known as the short form) with a shorter N-terminus, compared to isoform 1.