

## **Product datasheet for SC210741**

## Collagen IX (COL9A1) (NM\_001851) Human 3' UTR Clone

**Product data:** 

Product Type: 3' UTR Clones

Symbol: Collagen IX

Synonyms: DJ149L1.1.2; EDM6; MED; STL4

Mammalian Cell Neomycin

Selection:

Vector: pMirTarget (PS100062)

ACCN: NM\_001851

Insert Size: 801 bp

Insert Sequence: >SC210741 3'UTR clone of NM\_001851

The sequence shown below is from the reference sequence of NM\_001851. The complete sequence of

this clone may contain minor differences, such as  $\ensuremath{\mathsf{SNPs}}\xspace.$ 

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCCGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

ATCTCTGGTTGTACCAAAGAAGAAATAAATATGGTTTCTTAA

**ACGCGT**AAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul



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OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences, e.g., single nucleotide polymorphisms

(SNPs).

Components: The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

Note: Plasmids are not sterile. For experiments where strict sterility is required, filtration with 0.22um

filter is required.

**RefSeq:** <u>NM\_001851.6</u>

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 I 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]

**Locus ID:** 1297

**MW:** 30.7