

Product datasheet for SC208582

RDH12 (NM 152443) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: RDH12 (NM_152443) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: RDH12

Synonyms: LCA13; RP53; SDR7C2

ACCN: NM_152443

Insert Size: 633 bp

Insert Sequence: >SC208582 3'UTR clone of NM_152443

The sequence shown below is from the reference sequence of NM_152443. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

ACGCGTAAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

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.



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RDH12 (NM_152443) Human 3' UTR Clone - SC208582

RefSeq: <u>NM 152443.3</u>

Summary: The protein encoded by this gene is an NADPH-dependent retinal reductase whose highest

activity is toward 9-cis and all-trans-retinol. The encoded enzyme also plays a role in the metabolism of short-chain aldehydes but does not exhibit steroid dehydrogenase activity. Defects in this gene are a cause of Leber congenital amaurosis type 13 and Retinitis

Pigmentosa 53. [provided by RefSeq, Sep 2015]

Locus ID: 145226 MW: 23.3