

## **Product datasheet for SC206805**

## RRBP1 (NM 001042576) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: RRBP1 (NM 001042576) Human 3' UTR Clone

Symbol: RRBP1

**Synonyms:** ES/130; ES130; hES; p180; RRp

**Mammalian Cell** 

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_001042576

**Insert Size:** 533 bp

Insert Sequence: >SC206805 3'UTR clone of NM\_001042576

The sequence shown below is from the reference sequence of NM\_001042576. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

 ${\sf TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC}$ 

GTGTGACCTCCTGGGTGTTGATGCCATTAAAACCAACGTTGGTGCCCGGT

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).



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## RRBP1 (NM\_001042576) Human 3' UTR Clone - SC206805

**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.

**RefSeq:** <u>NM 001042576.2</u>

**Summary:** This gene encodes a ribosome-binding protein of the endoplasmic reticulum (ER) membrane.

Studies suggest that this gene plays a role in ER proliferation, secretory pathways and secretory cell differentiation, and mediation of ER-microtubule interactions. Alternative splicing has been observed and protein isoforms are characterized by regions of N-terminal decapeptide and C-terminal heptad repeats. Splicing of the tandem repeats results in variations in ribosome-binding affinity and secretory function. The full-length nature of variants which differ in repeat length has not been determined. Pseudogenes of this gene have been identified on chromosomes 3 and 7, and RRBP1 has been excluded as a candidate

gene in the cause of Alagille syndrome, the result of a mutation in a nearby gene on

chromosome 20p12. [provided by RefSeq, Apr 2012]

**Locus ID:** 6238

MW: 19.4