

Product datasheet for SC210066

HNRPH2 (HNRNPH2) (NM_019597) Human 3' UTR Clone

Product data:

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product Type:3' UTR ClonesProduct Name:HNRPH2 (HNRNPH2) (NM_019597) Human 3' UTR CloneSymbol:HNRPH2Symonyms:FTP3; hnRNPH'; HNRPH; HNRPH2; MRXSB; NRPH2Mammalian Cell Selection:NeomycinVector:pMirTarget (PS100062)ACCN:NM_019597Insert Size:825 bpInsert Sequence:Soc210066 3' UTR clone of NM_019597 The sequence shown below is from the reference sequence of NM_019597. The complete sequence of this clone may contain minor differences, such as SNPs. BluesStop Codon Red=Cloning siteGCCAATTGGCAGCCCCCCAGGACTCCCCCCCCCCCCCCC		
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	Restriction Sites:	Sgfl-Mlul



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	HNRPH2 (HNRNPH2) (NM_019597) Human 3' UTR Clone – SC210066
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.
RefSeq:	<u>NM 019597.5</u>
Summary:	This gene belongs to the subfamily of ubiquitously expressed heterogeneous nuclear ribonucleoproteins (hnRNPs). The hnRNPs are RNA binding proteins and they complex with heterogeneous nuclear RNA (hnRNA). These proteins are associated with pre-mRNAs in the nucleus and appear to influence pre-mRNA processing and other aspects of mRNA metabolism and transport. While all of the hnRNPs are present in the nucleus some seem to shuttle between the nucleus and the cytoplasm. The hnRNP proteins have distinct nucleic acid binding properties. The protein encoded by this gene has three repeats of quasi-RRM domains that binds to RNAs. It is very similar to the family member HNRPH1. This gene is thought to be involved in Fabray disease and X-linked agammaglobulinemia phenotype. Alternative splicing results in multiple transcript variants encoding the same protein. Read-through transcription between this locus and the ribosomal protein L36a gene has been observed. [provided by RefSeq, Jan 2011]
Locus ID:	3188
MW:	32.7

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