

Product datasheet for SC200359

IL1RAPL1 (NM_014271) Human 3' UTR Clone

Product data:

OriGene Technologies, Inc.

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Product Type:	3' UTR Clones
Product Name:	IL1RAPL1 (NM_014271) Human 3' UTR Clone
Symbol:	IL1RAPL1
Synonyms:	IL-1-RAPL-1; IL-1RAPL-1; IL1R8; IL1RAPL; IL1RAPL-1; MRX10; MRX21; MRX34; OPHN4; TIGIRR-2
Mammalian Cell Selection:	Neomycin
Vector:	pMirTarget (PS100062)
ACCN:	NM_014271
Insert Size:	928 bp
Insert Sequence:	<pre>>SC200359 3'UTR clone of NM_014271 The sequence shown below is from the reference sequence of NM_014271. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site GGCAAGTTGGACGCCCGCAAGATCCCGCGAGATTCTCATTAAGGCCAAGAAGGGCCGGAAAGATCGCCGTG</pre>
	GGCAAGTTGGACGCCCGCAAGATCCGCGAGGATTCTATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC GAGACCAGTATATCCAGTGTGATATGGTGACAGAAAAGCAAGGGACATCCCGTCCTGGGAGGTTGAGT GGAATCTGCAGTCCAGTGCCTGGAACTAAATCCTCGACTGCTGCTGTTAAAAAAACATGCATTAGAATCT CTAGAACACGAGGAAAAACAGGGTCTTGTACATATGTTTTTTGGAATTTCTTTGTAGCATCAGTGTCCT CCTGTTTTACCATGTCTTTTACCATTTACATTTTTTGACTTTGTTTTATATGTCGTTGGAATTGTAAAT TTACATTTTTTTAAAGAAGAGACTGATGTGTAGATAGAAAACCCTTTTTTTGCATCATTAGTTTAGTT TTAGAATGGGTTTTTATTTTTTTTTT
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.
RefSeq:	<u>NM 014271.4</u>
Summary:	The protein encoded by this gene is a member of the interleukin 1 receptor family and is similar to the interleukin 1 accessory proteins. This protein has an N-terminal signal peptide, three extracellular immunoglobulin Ig-like domains, a transmembrane domain, an intracellular Toll/IL-1R domain, and a long C-terminal tail which interacts with multiple signalling molecules. This gene is located at a region on chromosome X that is associated with a non-syndromic form of X-linked intellectual disability. Deletions and mutations in this gene were found in patients with intellectual disability. This gene is expressed at a high level in post-natal brain structures involved in the hippocampal memory system, which suggests a specialized role in the physiological processes underlying memory and learning abilities, and plays a role in synapse formation and stabilization. [provided by RefSeq, Jul 2017]
Locus ID:	11141
MW:	35.7

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