

Product datasheet for SC205140

PRAME (NM 206953) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: PRAME (NM_206953) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: PRAME

Synonyms: CT130; MAPE; OIP-4; OIP4

ACCN: NM_206953

Insert Size: 396 bp

Insert Sequence: >SC205140 3'UTR clone of NM_206953

The sequence shown below is from the reference sequence of NM_206953. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

ATCCTGTGCCCCTGTTTCATGCCTAATTAGCTGGGTGCACATATCAAATGCTTCATTCTGCATACTTGG
ACACTAAAGCCAGGATGTGCATGCATCTTGAAGCAACAAAGCAGCCACAGTTTCAGACAAATGTTCAGT
GTGAGTGAGGAAAACATGTTCAGTGAGGAAAAAACATTCAGACAAATGTTCAGTGAGGAAAAAAAGGGG
AAGTTGGGGGTAGGCAGATGTTGACTTGAGGAGTTAATGTGATCTTTGGGGAAGATACATCTTATAGAGT
TAGAAATAGAATCTGAATTTCTAAAGGGAGATTCTGGCTTGGGAAGTACATGTAGGAGTTAATCCCTGT

GTAGACTGTTGTAAAGAAACTGTTGAAAATAAAGAGAAGCAATGTGAAGCA

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.

RefSeg: NM 206953.3



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Summary: This gene encodes an antigen that is preferentially expressed in human melanomas and that

is recognized by cytolytic T lymphocytes. It is not expressed in normal tissues, except testis. The encoded protein acts as a repressor of retinoic acid receptor, and likely confers a growth advantage to cancer cells via this function. Alternative splicing results in multiple transcript

variants. [provided by RefSeq, Apr 2014]

Locus ID: 23532 MW: 15.2