

Product datasheet for **SC331146**

ZMYM2 (NM_001190964) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: ZMYM2 (NM_001190964) Human Untagged Clone
Tag: Tag Free
Symbol: ZMYM2
Synonyms: FIM; MYM; RAMP; SCLL; ZNF198
Vector: pCMV6-Entry (PS100001)
Fully Sequenced ORF: >SC331146 representing NM_001190964.
Blue=Insert sequence Red=Cloning site Green=Tag(s)

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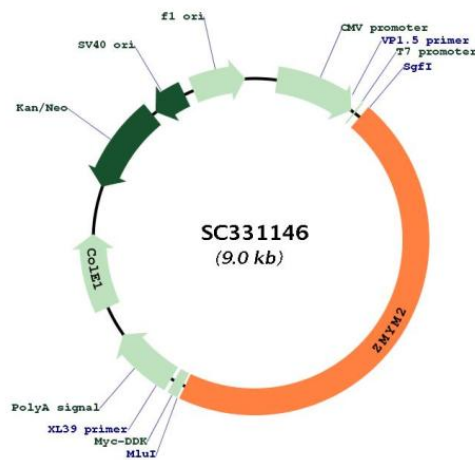
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Restriction Sites:

SgfI-MluI

Plasmid Map:



| | |
|-------------------------------|---|
| ACCN: | NM_001190964 |
| Insert Size: | 4134 bp |
| OTI Disclaimer: | Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP). |
| Components: | The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water). |
| Reconstitution Method: | <ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C. |
| RefSeq: | NM_001190964.1 |
| RefSeq Size: | 10138 bp |
| RefSeq ORF: | 4134 bp |
| Locus ID: | 7750 |
| UniProt ID: | Q9UBW7 |
| Cytogenetics: | 13q12.11 |
| Protein Families: | Druggable Genome, ES Cell Differentiation/IPS, Transcription Factors |
| MW: | 154.9 kDa |
| Gene Summary: | <p>The protein encoded by this gene is a zinc finger protein that may act as a transcription factor. The encoded protein may be part of a BHC histone deacetylase complex. Translocation of this gene with the fibroblast growth factor receptor-1 gene (FGFR1) results in a fusion gene, which may be a cause of stem cell leukemia lymphoma syndrome (SCLL). Several transcript variants encoding the same protein have been found for this gene. [provided by RefSeq, Jul 2010]</p> <p>Transcript Variant: This variant (3) differs in the 5' UTR compared to variant 1. Variants 1-4 encode the same protein. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.</p> |