

Product datasheet for MC226132

Rgr (NM_001301694) Mouse Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Rgr (NM_001301694) Mouse Untagged Clone

Tag: Tag Free

Symbol: Rgr

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Cell Selection: Neomycin

Fully Sequenced ORF: >MC226132 representing NM_001301694

Red=Cloning site Blue=ORF Orange=Stop codon

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC

GCCGCGATCGCC

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT

ACAAGGATGACGACGATAAGGTTTAA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001301694

Insert Size: 552 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).

OTI Annotation: Clone contains native stop codon, and expresses the complete ORF without any c-terminal

tag.



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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:

- 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: <u>NM 001301694.1</u>, <u>NP 001288623.1</u>

RefSeq Size: 4135 bp
RefSeq ORF: 552 bp
Locus ID: 57811
Cytogenetics: 14 B

Gene Summary: The gene is a member of the opsin family of G-protein coupled receptors. The encoded

protein is expressed in the retina, and acts as a photoisomerase to catalyze the conversion of all-trans-retinal to 11-cis-retinal. Disruption of a similar gene in human is associated with autosomal recessive (arRP) and autosomal dominant retinitis pigmentosa (adRP). Alternative

splicing results in multiple transcript variants. [provided by RefSeq, Sep 2014]

Transcript Variant: This variant (3) uses an alternate splice site and lacks an alternate exon in the 5' coding region, and uses an alternate downstream start codon compared to variant 1. The resulting protein (isoform 2) has a distinct N-terminus and is shorter compared to

isoform 1. Variants 2 and 3 encode the same isoform (2).