

Product datasheet for **RG230663**

PHKA1 (NM_001172436) Human Tagged ORF Clone

Product data:

| | |
|---------------------------|--|
| Product Type: | Expression Plasmids |
| Product Name: | PHKA1 (NM_001172436) Human Tagged ORF Clone |
| Tag: | TurboGFP |
| Symbol: | PHKA1 |
| Synonyms: | PHKA |
| Mammalian Cell Selection: | Neomycin |
| Vector: | pCMV6-AC-GFP (PS100010) |
| E. coli Selection: | Ampicillin (100 ug/mL) |
| ORF Nucleotide Sequence: | >RG230663 representing NM_001172436 Red=Cloning site Blue=ORF Green=Tags(s) |

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GCC**CGATCGCC**

ATGAGGAGCCGGAGTAACTCCGGGGTCCGGCTGGACGGCTACGCTCGACTGGTGCACAGACCATCCTGT
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AGAATGAAACTCAGTGGACGACCCTACAGACACATGGGAGTGCTTGGAACTTCAAACTCTATGACATTC
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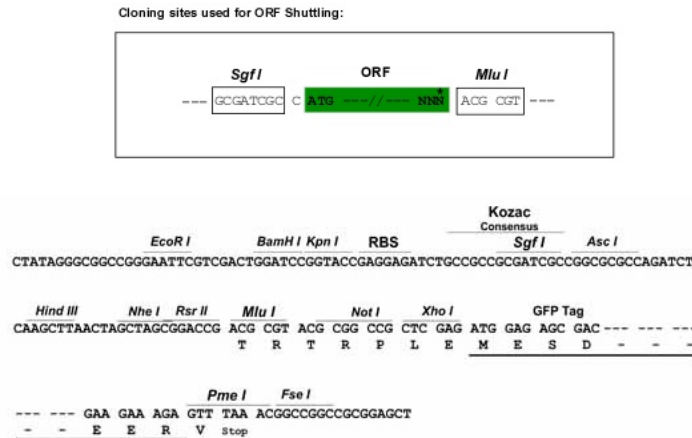
ACGCGTACGCGGCCGCTCGAG - GFP Tag - GTTTAA

Protein Sequence: >RG230663 representing NM_001172436
 Red=Cloning site Green=Tags(s)

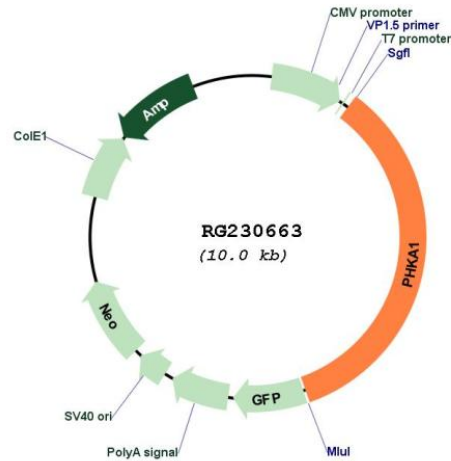
MRSRSNSGVRLDGYARLVQQTILCHQNPVTGLLPASYDQKDAWVRDNVYSILAVWGLGLAYRKNADRDDE
 KAKAYELEQSVVKLMRGLLHCMIRQVDKVESFKYSQSTKDSLHAKYNTKTCATVVGDDQWGHQLDQTSV
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 QEIIITKLQGRYGCCRFLRDGYKTPKEDPNRLYYEPAELKLFENIECEWPLFWTYFILDGVFSGNAEQVQE
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 RMKLSGRPYRHMVGLGTSKLYDIRKTIFFTPQFIDQQQFYALDNKMIVEMLRDLSYLCRWRMTGQP
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 DYDDNYDYLESGNWMNDYDSTSHDVHMYLPTKLFQASRPSFNLLDSPHPRQENQVPSVRVEIHLPRDQSG
 EVDFKALVLQKETSSELEQADILYMLYTMKGPDWNTL YNERSATVRELL TELYGVGVEIRHWGLIRYI
 SGILRKKVEALDEACTDLLSHQKHLTVGLPPEPREKTISAPLPYEAL TQLIDEASEGDMSSILTQEIMV
 YLAMYMRTQPLFAEMFRLRIGLI IQVMATELAHSLRCSAEAEGLMNLSPSAMKNLLHHILSGKEFGV
 ERSVPRDTSNVPASISIEIGAVGATKERTGIMQLKSEIKQSPGTSMT PSSGSP SAYDQSSKDSRQG
 QWQRRRLD GALNRVPVGFYQVWVVLQKCHGLSVEGFVLP SSTS TREMPGEIKFSVHVESV LNRVPQPE
 YRQLLVEAILVLTMLADIEIHSIGSIIAVEKIVHIANDLFLQE QKTLGADDTMLAKDPASGICTLLYDSA
 PSGRFGTMYLSKAAATYVQEFLPHSICAMQ

TRTRPLE - GFP Tag - V

Restriction Sites: SgfI-MluI
 Cloning Scheme:



Plasmid Map:



ACCN: NM_001172436

ORF Size: 3453 bp

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

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: [NM_001172436.1](#), [NP_001165907.1](#)

RefSeq Size: 5961 bp

RefSeq ORF: 3456 bp

Locus ID: 5255

UniProt ID: [P46020](#)

Cytogenetics: Xq13.1

| | |
|--------------------------|---|
| Protein Families: | Druggable Genome |
| Protein Pathways: | Calcium signaling pathway, Insulin signaling pathway |
| Gene Summary: | <p>Phosphorylase kinase is a polymer of 16 subunits, four each of alpha, beta, gamma and delta. The alpha subunit includes the skeletal muscle and hepatic isoforms, and the skeletal muscle isoform is encoded by this gene. The beta subunit is the same in both the muscle and hepatic isoforms, and encoded by one gene. The gamma subunit also includes the skeletal muscle and hepatic isoforms, which are encoded by two different genes. The delta subunit is a calmodulin and can be encoded by three different genes. The gamma subunits contain the active site of the enzyme, whereas the alpha and beta subunits have regulatory functions controlled by phosphorylation. The delta subunit mediates the dependence of the enzyme on calcium concentration. Mutations in this gene cause glycogen storage disease type 9D, also known as X-linked muscle glycogenosis. Alternatively spliced transcript variants encoding different isoforms have been identified in this gene. A pseudogene has been found on chromosome 1.[provided by RefSeq, Feb 2010]</p> |