

## Product datasheet for **RN216827**

### **Nrg1 (NM\_001271120) Rat Untagged Clone**

#### **Product data:**

Product Type:	Expression Plasmids
Product Name:	Nrg1 (NM_001271120) Rat Untagged Clone
Tag:	Tag Free
Symbol:	Nrg1
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Cell Selection:	Neomycin



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**Fully Sequenced ORF:** >RN216827 representing NM\_001271120  
 Red=Cloning site Blue=ORF Orange=Stop codon

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
 GCC**CGGATCGCC**

ATGGAGATTTATTCCCCAGACATGTCTGAGGTAGCTGGCGGGAGGTCCTCCAGCCCTCCACTCAGCTGA  
 GTGCAGCCCATCTCTTGTATGGCTTCCGGCAGCGGAGGAACATATACCAGACACCCACACAGAAGATGA  
 GAGAAGCCCTGGACTCCTGGGCCTGGCGGTGCCCTGTGTGTGCCTGGAAGCTGAGCGCTGAGAGGG  
 TGTCTCAACTCCGAGAAGATCTGCATTGTTCCATTCTGGCTTGCCTAGTCAGCCTCTGCCTCTGCATTG  
 CTGGCCTGAAGTGGGATTTGTGGACAAGATTTGAATACGACTCTCTACCCACCTTGACCCTGGGGG  
 GTTAGGCCAGGACCCTGTGATTTCTCTGGATCCAAGTCTGCCCCAGCCATTTTGGTATCATCTGAGGCA  
 TACTTACCTGTCTAAGGCTCAGTCTGAAGCTGGGGCTCATGTTACAGTACAAGGTGACCATGCTG  
 CTGTGGCCTCTGAACCTCAGCAGTACCGACCCGGAAGAACCCTGTCTGCTTTTCTCCCTTTCACTC  
 TACTGCACCGCCCTTCCCTTCTCCAGCTCGGACCCTGAGGTGAGAACACCCAAGTCAGGAAGTCAAGCA  
 CAAACAACAGAACTAACTGCAAAGTCTCTAACTTTCCACATCGACATCCAGACTGGGACCAGCC  
 ATCTCATAAAGTGCAGGAGAAGGAGAAAATTTCTGTGTGAATGGGGGGAGTGCTTCCAGGTGAAGGA  
 CCTGTCAAACCCGTCAAGATACTTGTGCAAGTGCCAACCTGGATTCACTGGAGCAAGATGTACTGAGAAT  
 GTACCCATGAAAGTCAAACCAAGAAAAGCGGAGGAACTCTACCAGAAGAGGGTGTGACAATTACTG  
 GCATCTGTATCGCCCTGTGGTGGTCCGGCATCATGTGTGTGGTGGCCTACTGCAAACCAAGAAGCAGCG  
 GCAGAAGCTTCATGATCGGCTTCGGCAGAGTCTTCGGTCAGAACGGAGCAACCTGGTGAACATAGCGAAT  
 GGGCCTACCACCCAAACCCACCGCCAGAGAACGTGCAGCTGGTGAATCAATACGTATCTAAAAACGTCA  
 TCTCCAGTGAGCATATTGTTGAGAGAGAAGTGGAGACTTCTTTTCCACCAGTCAATACACTTCCACAGC  
 CCATCACTCCACGACTGTCAACCCAGACTCCTAGTCACAGCTGGAGTAATGGGCACACGGAGAGCATCATT  
 TCAGAAAGCAACTCCGTAATCATGATGTCTTCGGTAGAGAACAGCAGGCACAGCAGTCCCGCCGGGGCC  
 CACGAGGACGTCTTCATGGCCTGGGAGGCCCTCGTGATAACAGCTTCTCAGGCATGCCAGAGAAAACCC  
 TGACTCTACAGAGACTCTCTCATAGCGAAAGACATAACCTTATAGCTGAGCTAAGGAGAAAACAAGGCT  
 TACAGATCCAAATGCATGCAGATCCAGCTGTCAGCAACTCATCTAGACCCTCTTCCATTACCCATTTGG  
 GCTTCATTCT**TAA**

**ACGCGT**ACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT  
 ACAAGGATGACGACGATAAGGTTTAA

- Restriction Sites:** Sgfl-Mlul
- ACCN:** NM\_001271120
- Insert Size:** 1554 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.
- 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\\_001271120.1](#), [NP\\_001258049.1](#)

**RefSeq Size:** 3842 bp

**RefSeq ORF:** 1554 bp

**Locus ID:** 112400

**Cytogenetics:** 16q12.3

**Gene Summary:** ligand for ErbB3 and ErbB4 receptors; gene produces many different alternative splicing isoforms; involved in neural and organ development [RGD, Feb 2006]  
Transcript Variant: This variant (3) lacks two alternate coding exons and contains two other alternate coding exons compared to variant 1. The resulting isoform (3) contains an alternate internal segment and has a shorter and distinct C-terminus compared to isoform 1. Sequence Note: The RefSeq transcript and protein were derived from genomic sequence to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on alignments.