

Product datasheet for **RG235942**

NDUFB9 (NM_001278646) Human Tagged ORF Clone

Product data:

Product Type: Expression Plasmids
Product Name: NDUFB9 (NM_001278646) Human Tagged ORF Clone
Tag: TurboGFP
Symbol: NDUFB9
Synonyms: B22; CI-B22; LYRM3; MC1DN24; UQOR22
Mammalian Cell Selection: Neomycin
Vector: pCMV6-AC-GFP (PS100010)
E. coli Selection: Ampicillin (100 ug/mL)
ORF Nucleotide Sequence: >RG235942 representing NM_001278646.
 Blue=ORF Red=Cloning site Green=Tag(s)

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GCTCGTTTAGTGAACCGTCAGAATTTGTAAACGACTCACTATAGGGCGCCGGGAATTCGTGACTG
GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC
ATGAGAGCCCGTTTGAAGAACAATAAGAATGAAAAGGATATGGCGAAGGCCACCCAGCTGCTGAAGGAG
GCCGAGGAAGAATTCTGGTACCGTCAGCATCCACAGCCATACATCTTCCCTGACTCTCTGGGGGCACC
TCCTATGAGAGATACGATTGCTACAAGTCCCAGAATGGTGCTTAGATGACTGGCATCCTTCTGAGAAG
GCAATGTATCCTGATTACTTTGCCAAGAGAGAACAGTGAAGAACTGCGGAGGGAAAGCTGGGAACGA
GAGGTTAAGCAGCTGCAGGAGGAAACGCCACCTGGTGGTCCTTAACTGAAGCTTGGCCCTGCCCGA
AAGGAAGGTGATTTGCCCCACTGTGGTGTATATTGTGACCAGACCCCGGAGCGGCCCATG
ACGCGTACGCGGCCGCTCGAG - GFP Tag - GTTTAAAC
```

Protein Sequence: >Peptide sequence encoded by RG235942
 Blue=ORF Red=Cloning site Green=Tag(s)

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MRARFEEHKNEKMAKATQLLKEAEEEFWYRQHPQPYIFPDSPPGTSYERYDCYKVPWCLDDWHPSEK
AMYPDYFAKREQWKKLRRESWEREVKQLQEETPPGGPLTEALPPARKEGDLPLWWYIVTRPRRPM
TRTRPLEMESDESGLPAMEIECRITGTLNGVEFELVGGGEGTPEQGRMTNKMSTKGALTFSPYLLSHV
MGYGFYHFGTYPSGYENPFLHAINNGGYNTRIEKYEDGGVLHVSFSYRYEAGRVIGDFKVMGTGFPE
SVIFTDKIIRSNAIVEHLHPMGDNDLDGSFTRTFSLRDGGYSSVVDSMHFKSAIHPSILQNGGPMFA
FRRVEEDHSNTELGIVEYQHAFKTPDADAGEERV
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Restriction Sites: Sgfl-Mlul



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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).
RefSeq:	NM_001278646.1 , NP_001265575.1
RefSeq Size:	733 bp
RefSeq ORF:	411 bp
Locus ID:	4715
Cytogenetics:	8q24.13
Protein Pathways:	Alzheimer's disease, Huntington's disease, Metabolic pathways, Oxidative phosphorylation, Parkinson's disease
MW:	17 kDa
Gene Summary:	The protein encoded by this gene is a subunit of the mitochondrial oxidative phosphorylation complex I (nicotinamide adenine dinucleotide: ubiquinone oxidoreductase). Complex I is localized to the inner mitochondrial membrane and functions to dehydrogenate nicotinamide adenine dinucleotide and to shuttle electrons to coenzyme Q. Complex I deficiency is the most common defect found in oxidative phosphorylation disorders and results in a range of conditions, including lethal neonatal disease, hypertrophic cardiomyopathy, liver disease, and adult-onset neurodegenerative disorders. Pseudogenes of this gene are found on chromosomes five, seven and eight. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2015]