

## Product datasheet for RC205164L4V

## OriGene Technologies, Inc.

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## DDX3Y (NM\_004660) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

Product Type: Lentiviral Particles

**Product Name:** DDX3Y (NM\_004660) Human Tagged ORF Clone Lentiviral Particle

Symbol: DDX3Y
Synonyms: DBY

Mammalian Cell Puromycin

Selection:

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**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_004660 **ORF Size:** 1980 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC205164).

Sequence:

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.

**RefSeg:** NM 004660.3

 RefSeq Size:
 4488 bp

 RefSeq ORF:
 1983 bp

 Locus ID:
 8653

 UniProt ID:
 O15523

 Cytogenetics:
 Yq11.221

**Domains:** DEAD, helicase\_C

**Protein Pathways:** RIG-I-like receptor signaling pathway





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**MW:** 73.2 kDa

**Gene Summary:** 

The protein encoded by this gene is a member of the DEAD-box RNA helicase family, characterized by nine conserved motifs, included the conserved Asp-Glu-Ala-Asp (DEAD) motif. These motifs are thought to be involved in ATP binding, hydrolysis, RNA binding, and in the formation of intramolecular interactions. This protein shares high similarity to DDX3X, on the X chromosome, but a deletion of this gene is not complemented by DDX3X. Mutations in this gene result in male infertility, a reduction in germ cell numbers, and can result in Sertolicell only sydrome. Pseudogenes sharing similarity to both this gene and the DDX3X paralog are found on chromosome 4 and the X chromosome. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Oct 2014]