

## Product datasheet for RC213364L4V

## OriGene Technologies, Inc.

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## Tau (MAPT) (NM\_016841) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Tau (MAPT) (NM\_016841) Human Tagged ORF Clone Lentiviral Particle

Symbol: Tau

Synonyms: DDPAC; FTDP-17; MAPTL; MSTD; MTBT1; MTBT2; PPND; PPP1R103; TAU; tau-40

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_016841 **ORF Size:** 1056 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC213364).

OTI Disclaimer:

Sequence:

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 016841.1

 RefSeq Size:
 2529 bp

 RefSeq ORF:
 1059 bp

 Locus ID:
 4137

 UniProt ID:
 P10636

 Cytogenetics:
 17q21.31

**Domains:** tubulin-binding

**Protein Families:** Druggable Genome





## Tau (MAPT) (NM\_016841) Human Tagged ORF Clone Lentiviral Particle - RC213364L4V

**Protein Pathways:** Alzheimer's disease, MAPK signaling pathway

MW: 36.6 kDa

**Gene Summary:** This gene encodes the microtubule-associated protein tau (MAPT) whose transcript

undergoes complex, regulated alternative splicing, giving rise to several mRNA species. MAPT

transcripts are differentially expressed in the nervous system, depending on stage of neuronal maturation and neuron type. MAPT gene mutations have been associated with

several neurodegenerative disorders such as Alzheimer's disease, Pick's disease,

frontotemporal dementia, cortico-basal degeneration and progressive supranuclear palsy.

[provided by RefSeq, Jul 2008]