

## Product datasheet for RC214812L4V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Tyrosine Hydroxylase (TH) (NM\_199292) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Tyrosine Hydroxylase (TH) (NM\_199292) Human Tagged ORF Clone Lentiviral Particle

Symbol: TH

Synonyms: DYT5b; DYT14; TYH

**Mammalian Cell** 

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_199292 **ORF Size:** 1584 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC214812).

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 199292.1

RefSeq Size: 1921 bp
RefSeq ORF: 1587 bp
Locus ID: 7054
UniProt ID: P07101
Cytogenetics: 11p15.5

**Protein Families:** Druggable Genome

**Protein Pathways:** Metabolic pathways, Parkinson's disease, Tyrosine metabolism





## Tyrosine Hydroxylase (TH) (NM\_199292) Human Tagged ORF Clone Lentiviral Particle – RC214812L4V

MW: 58.4 kDa

**Gene Summary:** 

The protein encoded by this gene is involved in the conversion of tyrosine to dopamine. It is the rate-limiting enzyme in the synthesis of catecholamines, hence plays a key role in the physiology of adrenergic neurons. Mutations in this gene have been associated with autosomal recessive Segawa syndrome. Alternatively spliced transcript variants encoding different isoforms have been noted for this gene. [provided by RefSeq, Jul 2008]