

## Product datasheet for RC202300L4V

## 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

## POLR1D (NM\_152705) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** POLR1D (NM\_152705) Human Tagged ORF Clone Lentiviral Particle

Symbol: POLR1D

Synonyms: AC19; POLR1C; RPA9; RPA16; RPAC2; RPC16; RPO1-3; TCS2

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_152705

ORF Size: 366 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC202300).

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 152705.1

 RefSeq Size:
 2043 bp

 RefSeq ORF:
 369 bp

 Locus ID:
 51082

 UniProt ID:
 P0DPB5

 Cytogenetics:
 13q12.2

**Protein Families:** Stem cell - Pluripotency, Transcription Factors





## POLR1D (NM\_152705) Human Tagged ORF Clone Lentiviral Particle - RC202300L4V

**Protein Pathways:** Cytosolic DNA-sensing pathway, Metabolic pathways, Purine metabolism, Pyrimidine

metabolism, RNA polymerase

MW: 14.3 kDa

**Gene Summary:** The protein encoded by this gene is a component of the RNA polymerase I and RNA

polymerase III complexes, which function in the synthesis of ribosomal RNA precursors and small RNAs, respectively. Mutations in this gene are a cause of Treacher Collins syndrome (TCS), a craniofacial development disorder. Alternative splicing results in multiple transcript

variants. [provided by RefSeq, Apr 2011]