

Product datasheet for SC329817

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POLR1D (NM_001206559) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: POLR1D (NM_001206559) Human Untagged Clone

Tag: Tag Free
Symbol: POLR1D

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

Vector: pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC329817 representing NM_001206559.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

CGGCGGTGA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001206559

Insert Size: 285 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

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).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: NM 001206559.1





Cytogenetics:

POLR1D (NM_001206559) Human Untagged Clone - SC329817

RefSeq Size: 2210 bp

 RefSeq ORF:
 285 bp

 Locus ID:
 51082

 UniProt ID:
 PODPB5

Protein Families: Stem cell - Pluripotency, Transcription Factors

13q12.2

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

metabolism, RNA polymerase

MW: 11 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]

Transcript Variant: This variant (3) differs in both UTRs and in the coding region, compared to variant 1. The encoded isoform (3) shares identity with isoform 2 but is distinct and shorter,

compared to isoform 1.