

Product datasheet for SC332927

CFC1 (NM 001270420) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: CFC1 (NM_001270420) Human Untagged Clone

Tag: Tag Free Symbol: CFC1

Synonyms: CFC1B; CRYPTIC; DTGA2; HTX2

Vector: pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC332927 representing NM_001270420.

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

TTTAATTTTCTATGTTGTAAATAA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001270420

Insert Size: 576 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).



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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: <u>NM 001270420.1</u>

 RefSeq Size:
 1562 bp

 RefSeq ORF:
 576 bp

 Locus ID:
 55997

 UniProt ID:
 POCG37

 Cytogenetics:
 2q21.1

 MW:
 20.3 kDa

Gene Summary: This gene encodes a member of the epidermal growth factor (EGF)- Cripto, Frl-1, and Cryptic

(CFC) family, which are involved in signalling during embryonic development. Proteins in this family share a variant EGF-like motif, a conserved cysteine-rich domain, and a C-terminal hydrophobic region. The protein encoded by this gene is necessary for patterning the left-

right embryonic axis. Mutations in this gene are associated with defects in organ development, including autosomal visceral heterotaxy and congenital heart disease.

Alternatively spliced transcript variants encoding multiple isoforms have been observed for

this gene. [provided by RefSeq, Jul 2012]

Transcript Variant: This variant (2) lacks an exon in the coding region, which results in a frameshift, compared to variant 1. The encoded isoform (2) has a distinct C-terminus and is

shorter than isoform 1.