

Product datasheet for SC333638

OriGene Technologies, Inc.

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Calcipressin 1 (RCAN1) (NM_001285392) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Calcipressin 1 (RCAN1) (NM 001285392) Human Untagged Clone

Tag: Tag Free

Symbol: Calcipressin 1

Synonyms: ADAPT78; CSP1; DSC1; DSCR1; MCIP1; RCN1

Vector: pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC333638 representing NM_001285392.

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

CTCAGCTGA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001285392

Insert Size: 354 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 001285392.2

 RefSeq Size:
 2327 bp

 RefSeq ORF:
 354 bp

 Locus ID:
 1827

 UniProt ID:
 P53805

 Cytogenetics:
 21q22.12

Protein Families: Transcription Factors

MW: 13.3 kDa

Gene Summary: The protein encoded by this gene interacts with calcineurin A and inhibits calcineurin-

dependent signaling pathways, possibly affecting central nervous system development. This gene is located in the minimal candidate region for the Down syndrome phenotype, and is overexpressed in the brain of Down syndrome fetuses. Chronic overexpression of this gene may lead to neurofibrillary tangles such as those associated with Alzheimer disease. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Nov 2013] Transcript Variant: This variant (6) uses an alternate promoter and 5' exon compared to variant 1. The resulting isoform (b) has a shorter N-terminus compared to isoform a.