

Product datasheet for **SC207227**

DISC1 (NM_001164551) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones
Product Name: DISC1 (NM_001164551) Human 3' UTR Clone
Vector: pMirTarget (PS100062)
Symbol: DISC1
Synonyms: C1orf136; SCZD9
ACCN: NM_001164551
Insert Size: 569 bp
Insert Sequence: >SC207227 3'UTR clone of NM_001164551
The sequence shown below is from the reference sequence of NM_001164551. The complete sequence of this clone may contain minor differences, such as SNPs.
Blue=Stop Codon **Red**=Cloning site

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GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC
GCCTTGGCCGTGGGGCCACTCAGCAGTGAAGTACTTGTATTGTCACCATTTTCCCCTCATGTTTCTTC
CTACCTCTCAGCTCCCACATAGTGAAGAGCTGGGAGGAGCTTAGCTGTAGAGACCCTTGGCTGCCTTTC
CTGGGAAGGGCCCTTTGTTGTTGGGAAGTGGGATTCTTCCAGAGTGAGAGATGGCCCATGTGGGTC
ATGCATCTCTAGAAAGAAGACTTTTCTGCCACTTGACAGAGAGAGAACCTTCTCAAGTGCAGCTGCATC
CCAGGTCTGAAGCATCTCTTGGGAACTACCCTTCTAAGAAGGTGGGGTAATTTCTCTTGTGAGGC
AGCTTGGTGTCTGTGATCATATTAAGCCTGTAAGATAAACAGATGGTCCCCTCTGACCGAGCGATGAA
AATCCATCCTCCTAACTCTGCCCTCCAGCTCCTGAGCAACATTTGCTATTTGTGTACCCCAAGGCT
CTTCTGAGATTAGTAATATATAGAGAGCAGTTGCTGCTTCTAATCCAGTGGCTTGTGGTCTGGCTGCA
TGTTTGAATAACTTAGA
ACGCGTAAGCGGCCGCGCATCTAGATTGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA
CGAGATTCGATTCCACCGCCGCTTCTATGAAAGG
```

Restriction Sites: SgfI-MluI

OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences, e.g., single nucleotide polymorphisms (SNPs).

Components: The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 µg dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.



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RefSeq: [NM_001164551.2](#)

Summary: This gene encodes a protein with multiple coiled coil motifs which is located in the nucleus, cytoplasm and mitochondria. The protein is involved in neurite outgrowth and cortical development through its interaction with other proteins. This gene is disrupted in a t(1;11) (q42.1;q14.3) translocation which segregates with schizophrenia and related psychiatric disorders in a large Scottish family. Alternate transcriptional splice variants, encoding different isoforms, have been characterized. [provided by RefSeq, Jul 2008]

Locus ID: 27185

MW: 21.4