

Product datasheet for **SC324955**

NDE1 (NM_001143979) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	NDE1 (NM_001143979) Human Untagged Clone
Tag:	Tag Free
Symbol:	NDE1
Synonyms:	HOM-TES-87; LIS4; MHAC; NDE; NUDE; NUDE1
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>SC324955 representing NM_001143979. Blue=Insert sequence Red=Cloning site Green=Tag(s)

```
GCTCGTTTAGTGAACCGTCAGAATTTTGTAAACGACTCACTATAGGGCGCCGGGAATTCGTCGACTG
GATCCGGTACCGAGGAGATCTGCCGCCCGGATCGCC
ATGGAGGACTCCGGAAAGACTTTCAGCTCCGAGGAGGAAGAAGCTAACTATTGGAAAGATCTGGCGATG
ACCTACAAACAGAGGGCAGAAAAACGCAAGAGGAACTCCGAGAATTCAGGAGGGAAGCCGAGAATAT
GAAGCTGAATTGGAGACGCAGCTGCAACAAATTGAAACCAGGAACAGAGACCTCCTGTCCGAAAATAAC
CGCCTTCGCATGGAGCTGGAAACCATCAAGGAGAAGTTTGAAGTGCAGCACTCTGAAGGCTACCGGCAG
ATCTCAGCCTTGGAGGATGACCTCGCGCAGACCAAGCCATTAAGACCAATTGCAGAAAATACATCAGA
GAGCTGGAGCAAGCAAATGACGACCTGGAAAGAGCCAAGCGGCCACGATCATGTCTCTCGAAGACTTT
GAGCAGCGCTTGAATCAGGCCATCGAAAGAAATGCCTTCCTGGAAAGTGAAGTGAATGATAAGAAAGAGAAT
CTCCTGGAATCTGTTAGAGACTGAAGGATGAAGCCAGAGATTTGCGGCAGGAAGTGGCCGTGCAGCAG
AAGCAGGAGAAACCCAGGACCCCATGCCAGCTCAGTGGAAAGCTGAGAGGACAGACACAGCTGTGCAG
GCCACGGGCTCCGTGCCGTCCACGCCATTGCTCACCGAGGACCCAGCTCAAGTTTAAACACACCTGGG
AGCTTCAGACGTGGCCTGGACGACTCCACCGGGGGACCCCTCACACCTGCGGCCCGGATATCAGCC
CTCAACATTGTGGGAGACCTACTGCGGAAAGTCCGGGCACTGGAGTCCAACTCGCTTCTGCCGGAAC
CTCGTGTACGATCAGTCCCCAAACCGAACAGGTGGCCAGCCTCTGGCGGAGCAGCAAGAACAGAGAT
GGCGGGGAGAGACGGCCAAGCAGCACCAGCGTGCCTTTGGGTGATAAGGGGTTGGACACGAGTTGCCGC
TGGTTGTCCAAATCAACAACAGGTCGTCAGCTCCTGCTGA
ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT
TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC
```

Restriction Sites:	Sgfl-MluI
ACCN:	NM_001143979
Insert Size:	1008 bp



[View online »](#)

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).
OTI Annotation:	This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.
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:	<ol style="list-style-type: none"> 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_001143979.1</u>
RefSeq Size:	3936 bp
RefSeq ORF:	1008 bp
Locus ID:	54820
UniProt ID:	<u>Q9NXR1</u>
Cytogenetics:	16p13.11
MW:	37.7 kDa
Gene Summary:	<p>This gene encodes a member of the nuclear distribution E (NudE) family of proteins. The encoded protein is localized at the centrosome and interacts with other centrosome components as part of a multiprotein complex that regulates dynein function. This protein plays an essential role in microtubule organization, mitosis and neuronal migration. Mutations in this gene cause lissencephaly 4, a disorder characterized by lissencephaly, severe brain atrophy, microcephaly, and severe cognitive disability. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2012]</p> <p>Transcript Variant: This variant (1) represents the longer transcript. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.</p>