

Product datasheet for **TR305083**

DCTN3 Human shRNA Plasmid Kit (Locus ID 11258)

Product data:

Product Type:	shRNA Plasmids
Product Name:	DCTN3 Human shRNA Plasmid Kit (Locus ID 11258)
Locus ID:	11258
Synonyms:	DCTN-22; DCTN22
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	DCTN3 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 11258). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	NM_001281425 , NM_001281426 , NM_001281427 , NM_007234 , NM_024348 , NM_024348.1 , NM_024348.2 , NM_024348.3 , NM_007234.1 , NM_007234.2 , NM_007234.3 , NM_007234.4 , NM_001281426.1 , NM_001281425.1 , NM_001281427.1 , BC003004 , BC003004.1 , BC107697 , BC107697.1 , BC000319 , NM_024348.4 , NM_007234.5 , NM_001281426.2 , NM_001281427.2
UniProt ID:	O75935
Summary:	This gene encodes the smallest subunit of dynactin, a macromolecular complex consisting of 10 subunits ranging in size from 22 to 150 kD. Dynactin binds to both microtubules and cytoplasmic dynein. It is involved in a diverse array of cellular functions, including ER-to-Golgi transport, the centripetal movement of lysosomes and endosomes, spindle formation, cytokinesis, chromosome movement, nuclear positioning, and axonogenesis. This subunit, like most other dynactin subunits, exists only as a part of the dynactin complex. It is primarily an alpha-helical protein with very little coiled coil, and binds directly to the largest subunit (p150) of dynactin. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2013]
shRNA Design:	These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact techsupport@origene.com . If you need a special design or shRNA sequence, please utilize our custom shRNA service .



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**Performance
Guaranteed:**

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).