

DATASHEET

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TUBA3A siRNA Catalogue No.:abx938545

siRNA to inhibit TUBA3A expression using RNA interference.

This product is provided as two 5 nmol vials (10 nmol), three 5 nmol vials (15 nmol) or 2x three 5 nmol vials (30 nmol) of lyophilized siRNA oligo duplexes. Each vial contains slightly different sequences to ensure full knockout of the gene. The duplexes can be transfected individually or pooled together to achieve knockdown of the target gene, which is most commonly assessed by qPCR or western blot. The siRNA oligos are also chemically modified (2'-OMe) for increased stability and enhanced knockdown in vitro and in vivo.

expression.Storage:Shipped at 4 °C. Store at -20 °C for up to one year.Swiss Prot:Q68FR8GeneID:500319Gene Symbol:TUBA3ADirections for use:1. Transfect with 100 nM siRNA 48 to 72 hours prior to cell lysis. 2. Before resuspending, briefly centrifuge the tube to ensure the lyophilized siRNA is at the bottom of the tube. 3. Resuspend the siRNA oligos to an appropriate concentration with DEPC water. Each vial is suitable for 250 transfections in a 24 well plate (20 pmol for each well).Quality Control:Oligonucleotide synthesis is monitored base by base through trityl analysis to ensure appropriate coupling efficiency. The oligo is subsequently purified by affinity-solid phase extraction. The annealed RNA duplex is further analyzed by mass spectrometry to verify the exact composition of the duplex.	Target:	TUBA3A
Tested Applications:RNAPurity:> 97%Form:LyophilizedForm:UpophilizedSpecificity:TUBA3A siRNA (Rat) is a target-specific 19-23 nt siRNA oligo duplexes designed to knock down gene expression.Storage:Obipped at 4 °C. Store at -20 °C for up to one year.Storage:Q68FR8GenelD:500319TUBA3ATUBA3ADirections for use:1. Transfect with 100 nM siRNA 48 to 72 hours prior to cell lysis. 2. Before resuspending, briefly centrifuge the tube to ensure the lyophilized siRNA is at the bottom of the tube. 3. Resuspend the siRNA oligos to an appropriate concentration with DEPC water. Each vial is suitable for 250 transfections in a 24 well plate (20 pmol for each well).Quality Contror:Oligonucleotide synthesis is monitored base by base through trily analysis to ensure appropriate coupling efficiency. The oligo is subsequently purified by affinity-solid phase extraction. The annealed RNA duplex is further analyzed by mass spectrometry to verify the exact composition of the duplex.	Reactivity:	Rat
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Note: This product is for research use only.