

Product Data Sheet

FOXN3 siRNA (Human)

Source	Reactivity	Appl	ications		
Synthetic	н	RNA	i		
Description siRNA to inhibit FOXN3 expression using RNA interference					
FOXN	FOXN3 siRNA (Human) is a target-specific 19-23 nt siRNA oligo duplexes designed to				
knock	down gene expressio	on.			
Lyoph	ilized powder				
Gene Symbol FOXN3					
ames C14or	f116; CHES1; Forkhea	ad box protein N3; Ch	eckpoint suppressor 1		
1112	(Human)				
00040	09 (Human)				
> 97%					
ol Oligor	Oligonucleotide synthesis is monitored base by base through trityl analysis to ensur				
appro	priate coupling efficie	ency. The oligo is subs	equently purified by affinity-solid		
phase	extraction. The anne	aled RNA duplex is fu	rther analyzed by mass		
spectr	rometry to verify the	exact composition of	the duplex. Each lot is compared to		
the pr	evious lot by mass sp	ectrometry to ensure	maximum lot-to-lot consistency.		
onents We offers pre-designed sets of 3 different target-specific siRNA oligo duplexes of					
huma	n FOXN3 gene. Each v	vial contains 5 nmol of	f lyophilized siRNA. The duplexes		
can be	e transfected individu	ally or pooled togethe	er to achieve knockdown of the		
target gene, which is most commonly assessed by qPCR or western blot.					
Com	ponent	15 nm	ol 30 nmol		
FOXN	N3 siRNA (Human) - A	5 nmo	l x 1 5 nmol x 2		
FOXN	N3 siRNA (Human) - B	5 nmo	l x 1 5 nmol x 2		
	Synthetic siRNA FOXN knock Lyoph FOXN TOU OOO40 > 97% Oligor appro phase spectr the pr We of huma can be target Com	SyntheticHsiRNA to inhibit FOXN3 exp FOXN3 siRNA (Human) is a knock down gene expression Lyophilized powderFOXN3FOXN3amesC14orf116; CHES1; Forkhead 1112 (Human) O00409 (Human) > 97%OlOligonucleotide synthesis is appropriate coupling efficient phase extraction. The anneed spectrometry to verify the the previous lot by mass spectrometry to verify the the pr	Synthetic H RNA siRNA to inhibit FOXN3 expression using RNA int FOXN3 siRNA (Human) is a target-specific 19-23 in knock down gene expression. Lyophilized powder FOXN3 FOXN3 Enersion C14orf116; CHES1; Forkhead box protein N3; Chein 1112 (Human) O00409 (Human) > 97% Oligonucleotide synthesis is monitored base by the appropriate coupling efficiency. The oligo is subserphase extraction. The annealed RNA duplex is fur spectrometry to verify the exact composition of the previous lot by mass spectrometry to ensure We offers pre-designed sets of 3 different target human FOXN3 gene. Each vial contains 5 nmol or can be transfected individually or pooled together target gene, which is most commonly assessed by Example. Component 15 nm FOXN3 siRNA (Human) - A 5 nmol		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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FOXN3 siRNA (Human) - C	5 nmol x 1	5 nmol x 2
Negative Control	2.5 nmol x 1	2.5 nmol x 2
 DEPC Water	1 ml x 1	1 ml x 2

Directions for Use

We recommends transfection with 10 nM - 100 nM siRNA 48 to 72 hours prior to cell lysis. Before resuspending, briefly centrifuge the tube to ensure the lyophilized siRNA is at the bottom of the tube. Resuspend the siRNA oligos to an appropriate concentration with DEPC water. For example, resuspend one tube of 5 nmol siRNA oligo in 250 μ l of DEPC water to get a final concentration of 20 μ M.

Plate	Final volume	Final concentration	siRNA (20 μM)	Lipofectamin
	of medium	of siRNA		2000
		100 nM	0.5 μl	0.25 μl
96-well	100 µl	50 nM	0.25 μl	0.25 μl
		10 nM	0.05 μl	0.25 μl
		100 nM	2.5 μl	1 µl
24-well	500 μl	50 nM	1.25 μl	1 µl
		10 nM	0.25 μl	1 µl
		100 nM	5 μl	2 µl
12-well	1 ml	50 nM	2.5 μl	2 µl
		10 nM	0.5 μl	2 µl
		100 nM	10 µl	5 µl
6-well	2 ml	50 nM	5 μΙ	5 µl
		10 nM	1 µl	5 µl

Storage/Stability

Shipped at 4 °C. Store at -20 °C for one year.

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