

Product Data Sheet

ACRV1 siRNA (Human)

Catalog #	Source	Reactivity	Applications		
CRH0044	Synthetic	н	RNAi		
Description	Description siRNA to inhibit ACRV1 expression using RNA interference				
Specificity	ACRV1	ACRV1 siRNA (Human) is a target-specific 19-23 nt siRNA oligo duplexes designed to			
	knock	knock down gene expression.			
Form	Lyophi	Lyophilized powder			
Gene Symbol	ACRV1	ACRV1			
Alternative Na	mes Acrosc	Acrosomal protein SP-10; Acrosomal vesicle protein 1			
Entrez Gene 56 (Human)					
SwissProt P26436 (Human)					
Purity > 97%					
Quality Control Oligonucleotide synthesis is monitored base by base through trityl analysis t			ugh trityl analysis to ensure		
	approj	appropriate coupling efficiency. The oligo is subsequently purified by affinity-solid			
	phase	phase extraction. The annealed RNA duplex is further analyzed by mass			
	spectr	spectrometry to verify the exact composition of the duplex. Each lot is compared to			
	the pro	evious lot by mass sp	ectrometry to ensure maximur	n lot-to-lot consistency.	
Components We offers pre-designed sets of 3 different target-specific siRNA oligo dup			iRNA oligo duplexes of		
	humar	human ACRV1 gene. Each vial contains 5 nmol of lyophilized siRNA. The duplexes			
	can be	can be transfected individually or pooled together to achieve knockdown of the			
	target	target gene, which is most commonly assessed by qPCR or western blot.			
	Com	ponent	15 nmol	30 nmol	
	ACRV	'1 siRNA (Human) - A	5 nmol x 1	5 nmol x 2	
	ACRV	'1 siRNA (Human) - B	5 nmol x 1	5 nmol x 2	
Alternative Nat Entrez Gene SwissProt Purity Quality Control	mes Acroso 56 (Hu P2643 > 97% Oligon approp phase spectro the pro We off humar can be target Comp ACRV	omal protein SP-10; A iman) 6 (Human) nucleotide synthesis is priate coupling efficie extraction. The annea ometry to verify the e evious lot by mass spe fers pre-designed sets n ACRV1 gene. Each v e transfected individua gene, which is most o ponent '1 siRNA (Human) - A	monitored base by base thround ncy. The oligo is subsequently aled RNA duplex is further ana exact composition of the duple ectrometry to ensure maximum of 3 different target-specific sial contains 5 nmol of lyophilized ally or pooled together to achie commonly assessed by qPCR or 15 nmol 5 nmol x 1	purified by affinity-solid lyzed by mass x. Each lot is compared m lot-to-lot consistency iRNA oligo duplexes of ed siRNA. The duplexes eve knockdown of the r western blot. 30 nmol 5 nmol x 2	

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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ACRV1 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 μl	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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