

Product Data Sheet

MYBPH siRNA (Rat)

Source	Reactivity	Applications		
Synthetic	R	RNAi		
siRNA	siRNA to inhibit MYBPH expression using RNA interference			
MYBP	MYBPH siRNA (Rat) is a target-specific 19-23 nt siRNA oligo duplexes designed to			
knock	down gene expressio	n.		
Form Lyophilized pov				
Gene Symbol MYBPH				
mes Myosi	Myosin-binding protein H; MyBP-H; H-protein			
83708	83708 (Rat)			
O8859	O88599 (Rat)			
> 97%	> 97%			
l Oligor	Oligonucleotide synthesis is monitored base by base through trityl analysis to ens			
appropriate coupling efficiency. The oligo is subsequently purified by affinity-so			ourified by affinity-solid	
phase	phase extraction. The annealed RNA duplex is further analyzed by mass			
spectr	ometry to verify the e	exact composition of the duplex	. Each lot is compared to	
the pr	evious lot by mass spe	ectrometry to ensure maximum	lot-to-lot consistency.	
We of	fers pre-designed sets	of 3 different target-specific sil	RNA oligo duplexes of rat	
MYBP	MYBPH gene. Each vial contains 5 nmol of lyophilized siRNA. The duplexes can be			
transf	transfected individually or pooled together to achieve knockdown of the target			
			n blot.	
Com	ponent	15 nmol	30 nmol	
			5 nmol x 2	
MYB				
	Synthetic siRNA MYBP knock Lyoph MYBP Myosi 83708 08859 > 97% Oligor appro phase spectr the pr We of MYBP transf gene, Com	SyntheticRsiRNA to inhibit MYBPH expMYBPH siRNA (Rat) is a targknock down gene expressioLyophilized powderMYBPHmesMyosin-binding protein H; N83708 (Rat)088599 (Rat)> 97%Oligonucleotide synthesis is appropriate coupling efficie phase extraction. The annea spectrometry to verify the e the previous lot by mass spec We offers pre-designed sets MYBPH gene. Each vial cont transfected individually or p	SyntheticRRNAisiRNA to inhibit MYBPH expression using RNA interferenceMYBPH siRNA (Rat) is a target-specific 19-23 nt siRNA oligoknock down gene expression.Lyophilized powderMYBPHmesMyosin-binding protein H; MyBP-H; H-protein83708 (Rat)O88599 (Rat)> 97%Oligonucleotide synthesis is monitored base by base througappropriate coupling efficiency. The oligo is subsequently pphase extraction. The annealed RNA duplex is further analyspectrometry to verify the exact composition of the duplexthe previous lot by mass spectrometry to ensure maximumWe offers pre-designed sets of 3 different target-specific silMYBPH gene. Each vial contains 5 nmol of lyophilized siRNAtransfected individually or pooled together to achieve knowgene, which is most commonly assessed by qPCR or westerComponent15 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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DEPC Water	1 ml x 1	1 ml x 2
Negative Control	2.5 nmol x 1	2.5 nmol x 2
MYBPH siRNA (Rat) - C	5 nmol x 1	5 nmol 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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