

Product Data Sheet

CYBA siRNA (Human)

rce	Reactivity	Applications		
hetic	н	RNAi		
siRNA	to inhibit CYBA expr	ession using RNA interference		
CYBA :	siRNA (Human) is a t	arget-specific 19-23 nt siRNA oligo duplexes designed to		
knock	down gene expressi	on.		
Lyoph	ilized powder			
CYBA	СҮВА			
Cytocł	Cytochrome b-245 light chain; Cytochrome b(558) alpha chain; Cytochrome b558			
subun	it alpha; Neutrophil	cytochrome b 22 kDa polypeptide; Superoxide-generating		
NADP	H oxidase light chain	subunit; p22 phagocyte B-cytochrome; p22-phox;		
p22ph	юх			
1535 (1535 (Human)			
P1349	P13498 (Human)			
> 97%				
Oligor	Oligonucleotide synthesis is monitored base by base through trityl analysis to ensure			
appro	priate coupling effici	ency. The oligo is subsequently purified by affinity-solid		
phase	extraction. The anne	ealed RNA duplex is further analyzed by mass		
spectr	ometry to verify the	exact composition of the duplex. Each lot is compared to		
the pr	evious lot by mass sp	pectrometry to ensure maximum lot-to-lot consistency.		
We of	We offers pre-designed sets of 3 different target-specific siRNA oligo duplexes of			
humai	n CYBA gene. Each vi	al contains 5 nmol of lyophilized siRNA. The duplexes can		
be tra	nsfected individually	or pooled together to achieve knockdown of the target		
gene,	which is most comm	only assessed by qPCR or western blot.		
	CYBA knock Lyoph CYBA Cytocl subun NADP p22ph 1535 (P1349 > 97% Oligor appro phase spectr the pr We of huma be tra	theticHsiRNA to inhibit CYBA exprCYBA siRNA (Human) is a takknock down gene expressiaLyophilized powderCYBACytochrome b-245 light chsubunit alpha; NeutrophilaNADPH oxidase light chainp22phox1535 (Human)P13498 (Human)> 97%Oligonucleotide synthesis iappropriate coupling efficiaphase extraction. The annespectrometry to verify thethe previous lot by mass spWe offers pre-designed sethuman CYBA gene. Each vibe transfected individually		

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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Component	15 nmol	30 nmol
CYBA siRNA (Human) - A	5 nmol x 1	5 nmol x 2
CYBA siRNA (Human) - B	5 nmol x 1	5 nmol x 2
CYBA 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 μΙ
12-well	1 ml	50 nM	2.5 μl	2 μΙ
		10 nM	0.5 μl	2 μΙ
6-well	2 ml	100 nM	10 µl	5 µl
		50 nM	5 μl	5 μΙ

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10 nM

1 µl

5 µl

Storage/Stability

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

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