

Product Data Sheet

UBA1 siRNA (Human)

e Reactivity	Applications		
etic H	RNAi		
escription siRNA to inhibit UBA1 expression using RNA interference			
UBA1 siRNA (Human) is a target-specific 19-23 nt siRNA oligo duplexes designed to			
knock down gene expression.			
Lyophilized powder			
UBA1			
ternative Names A1S9T; UBE1; Ubiquitin-like modifier-activating enzyme 1; Protein A1S9;		otein A1S9;	
Ubiquitin-activating enzyme E1			
7317 (Human)			
P22314 (Human)			
> 97%			
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. Each lot is compared to			
the previous lot by mass spectrometry to ensure maximum lot-to-lot consistency.			
We offers pre-designed sets of 3 different target-specific siRNA oligo duplexes of			
human UBA1 gene. Each vial contains 5 nmol of lyophilized siRNA. The duplexes can			
be transfected individually or po	oled together to achieve kno	ckdown of the target	
gene, which is most commonly assessed by qPCR or western blot. Component 15 nmol 30 nmol			
			UBA1 siRNA (Human) - A
è	siRNA to inhibit UBA1 expression UBA1 siRNA (Human) is a target- knock down gene expression. Lyophilized powder UBA1 A1S9T; UBE1; Ubiquitin-like mod Ubiquitin-activating enzyme E1 7317 (Human) P22314 (Human) > 97% Oligonucleotide synthesis is mor appropriate coupling efficiency. phase extraction. The annealed spectrometry to verify the exact the previous lot by mass spectro We offers pre-designed sets of 3 human UBA1 gene. Each vial cor	ticHRNAisiRNA to inhibit UBA1 expression using RNA interferenceUBA1 siRNA (Human) is a target-specific 19-23 nt siRNA oligo knock down gene expression.Lyophilized powderUBA1A1S9T; UBE1; Ubiquitin-like modifier-activating enzyme 1; Pro Ubiquitin-activating enzyme E17317 (Human)P22314 (Human)> 97%Oligonucleotide synthesis is monitored base by base through appropriate coupling efficiency. The oligo is subsequently pur phase extraction. The annealed RNA duplex is further analyze spectrometry to verify the exact composition of the duplex. E the previous lot by mass spectrometry to ensure maximum lo We offers pre-designed sets of 3 different target-specific siRN human UBA1 gene. Each vial contains 5 nmol of lyophilized si be transfected individually or pooled together to achieve kno gene, which is most commonly assessed by qPCR or westernComponent15 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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UBA1 siRNA (Human) - B	5 nmol x 1	5 nmol x 2
UBA1 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 μΙ
		10 nM	0.25 μl	1 μΙ
		100 nM	5 μl	2 μl
12-well	1 ml	50 nM	2.5 μl	2 μΙ
		10 nM	0.5 μl	2 μΙ
		100 nM	10 µl	5 µl
6-well	2 ml	50 nM	5 µl	5 μΙ
		10 nM	1 μΙ	5 μΙ

Storage/Stability

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

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