

Product Data Sheet

CGA siRNA (Human)

Source	Reactivity	Applications		
Synthetic	н	RNAi		
siRNA	A to inhibit CGA expre	ssion using RNA interference		
CGA	CGA siRNA (Human) is a target-specific 19-23 nt siRNA oligo duplexes designed to			
knoc	k down gene expressi	on.		
Lyopl	hilized powder			
CGA	CGA			
ames Glyco	protein hormones al	bha chain; Anterior pituitary glycoprotein hormones		
comr	mon subunit alpha; Ch	oriogonadotropin alpha chain; Chorionic gonadotrophin		
subu	nit alpha; CG-alpha; F	ollicle-stimulating hormone alpha chain; FSH-alpha;		
Follit	ropin alpha chain			
1081	1081 (Human)			
P012	P01215 (Human)			
> 97%	6			
ol Oligo	nucleotide synthesis	s monitored base by base through trityl analysis to ensure		
appro	opriate coupling effici	ency. The oligo is subsequently purified by affinity-solid		
phase	e extraction. The anne	ealed RNA duplex is further analyzed by mass		
spect	rometry to verify the	exact composition of the duplex. Each lot is compared to		
the p	revious lot by mass s	pectrometry to ensure maximum lot-to-lot consistency.		
We o	ffers pre-designed se	s of 3 different target-specific siRNA oligo duplexes of		
huma	an CGA gene. Each via	l contains 5 nmol of lyophilized siRNA. The duplexes can		
be tra	ansfected individually	or pooled together to achieve knockdown of the target		
gene	, which is most comm	only assessed by qPCR or western blot.		
	Synthetic siRN/ CGA knoc Lyop CGA CGA com subu Follit 1081 P012 > 979 Oligo appro phas spect the p We o huma	SyntheticHsiRNA to inhibit CGA expressionCGA siRNA (Human) is a takeknock down gene expressionLyophilized powderCGACGAGlycoprotein hormones algocommon subunit alpha; CG-alpha; Feisubunit alpha; CG-alpha; FeiFollitropin alpha chain1081 (Human)P01215 (Human)> 97%Oligonucleotide synthesis in appropriate coupling efficient phase extraction. The anneesspectrometry to verify the the previous lot by mass spectrometry to verify the the previous lot by mass spectrometry to verify the the previous lot by mass spectrometry in the previous lot by mass spectrometry to verify the the previous lot by mass spectrome		

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
CGA siRNA (Human) - A	5 nmol x 1	5 nmol x 2
CGA siRNA (Human) - B	5 nmol x 1	5 nmol x 2
CGA 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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