

Product Data Sheet

TXLNB siRNA (Human)

Reactivity	Applications	
Н	RNAi	
NA to inhibit TXLNB expression using	RNA interference	
TXLNB siRNA (Human) is a target-specific 19-23 nt siRNA oligo duplexes designed to		
ock down gene expression.		
ophilized powder		
bol TXLNB		
Alternative Names C6orf198; MDP77; Beta-taxilin; Muscle-derived protein 77; hMDP77		
e 167838 (Human)		
Q8N3L3 (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.		
Components We offers pre-designed sets of 3 different target-specific siRNA oligo duplexes of		
human TXLNB gene. Each vial contains 5 nmol of lyophilized siRNA. The duplexes can		
transfected individually or pooled tog	ether to achieve knoc	kdown of the target
gene, which is most commonly assessed by qPCR or western blot.		
Component 15 nmol 30 nmol		
	5 nmol x 1	
	NA to inhibit TXLNB expression using LNB siRNA (Human) is a target-specific ock down gene expression. ophilized powder LNB orf198; MDP77; Beta-taxilin; Muscle-c 7838 (Human) 3N3L3 (Human) 37% gonucleotide synthesis is monitored b propriate coupling efficiency. The oligo ase extraction. The annealed RNA dup ectrometry to verify the exact compos e previous lot by mass spectrometry to e offers pre-designed sets of 3 different man TXLNB gene. Each vial contains 5 transfected individually or pooled tog ne, which is most commonly assessed	H RNAi RNA to inhibit TXLNB expression using RNA interference LNB siRNA (Human) is a target-specific 19-23 nt siRNA oligo ock down gene expression. ophilized powder LNB orf198; MDP77; Beta-taxilin; Muscle-derived protein 77; hN 7838 (Human) RN3L3 (Human) 207% gonucleotide synthesis is monitored base by base through t propriate coupling efficiency. The oligo is subsequently puri- ase extraction. The annealed RNA duplex is further analyzed ectrometry to verify the exact composition of the duplex. Ea e previous lot by mass spectrometry to ensure maximum lot e offers pre-designed sets of 3 different target-specific siRNA man TXLNB gene. Each vial contains 5 nmol of lyophilized si transfected individually or pooled together to achieve knoc ne, which is most commonly assessed by qPCR or western b

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

5 nmol x 1

5 nmol x 2

TXLNB siRNA (Human) - B

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Product Data Sheet

TXLNB 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 μΙ	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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