

Supplementary Table S1 Primers used in this study.

Primer name	Primer equence (5' – 3')
Construction of cloning site in pCHL3-SB	
Y3-5-NheI	<u>gctagc</u> GTGTGTAGTAATTGGCA
Y3-StuI-BglII-CP3	<u>agatctCaggcctGACTCGACTCAATTCTACG</u>
Y3-StuI-BglII-CP5	<u>GAGTCaggcctGagatct</u> ATGGACAAATCTGAATCAAC
Y3-3-EcoRI	CGgaattcTGGTCTCCTTTTGGAGGC
Construction of cloning site in pCHL3-EB	
Y3-5-NheI	<u>gctagc</u> GTGTGTAGTAATTGGCA
Y3-EcoRV-BglII-CP3	<u>agatctCgatateGACTCGACTCAATTCTACG</u>
Y3-EcoRV-BglII-CP5	<u>GAGTCgatateGagatct</u> ATGGACAAATCTGAATCAAC
Y-CP-3-100	GAAAGCTGCTGCGACAAGACTC
Isolation of <i>LIPDS</i>	
LIPDSdf	GCAGAATTTGTTTGGGGAAC
LIPDSer	CAATGGGGTGACATGCTCTA
Creation of 33-base <i>LIPDS</i> insert	
PDS33f	CATagatctGCCTTCTACCTGCTATG
LIPDSbr	TTGGAGCCATGCTTTTCCTG
Creation of 52-base <i>LIPDS</i> insert	
LIPDShf	GCGgatateCGAGAGGTTGTGCATACCAAT
LIPDSdr	GCGagatctTTGACCACCAATGATTGAA
Amplification of RNA1 genome	
CY2T7	CCGGATCCGATTAATACGACTCACTATAGTTTATTTACAAGAGCC
HL-1-3	TGGTCTCCTTGTGGAGCC
Gene specific primers for qRT-PCR	
LIPDScf	CAGGAAAAGCATGGCTCCAAG
LIPDSfr	TCTCTCCATTCTTGAGGCAAA
Lh18SrRNAaf	TGCAACAAACCCCGACTTTC
Lh18SrRNAbr	CCGTCACCCGTCAATACCAT
Detection of CMV-HL vector and the deletion of the <i>PDS</i> fragments	
HL3ff	CTCCGCGAGATTGCGTTAT
HL3er	ACGACCAGCTGCTAACGTCT
Detection of U6 small nuclear RNA	
U6af	CGGGGACATCCGATAAAATTGGAACG
U6br	CGATTTGTGCGTGTCAATCCTTGC

Overlapped regions were underlined. Lower-case letters indicate restriction sites.

Supplementary Table S2 Sequences of siRNAs and primers used for stem-loop pulsed RT and end-point PCR (5' – 3')

siRNA derived from *LIPDS 33*

Expected siRNA sequence *AAUCAUAGCAGGUAG* AAGGCA

PDS33_Stem-loop RT primer GTTGGCTCTGGTGCAGGGTCCGAGGTATTCGCACCAGAGCCAACTGCCTT

PDS33_Forward primer *TCGCGAATCATAGCAGGTAG*

siRNA derived from *LIPDS 52*

Expected siRNA sequence *UUGGUAUGCACAACC* UCUCGG

PDS52_Stem-loop RT primer GTTGGCTCTGGTGCAGGGTCCGAGGTATTCGCACCAGAGCCAACCCGAGA

PDS52_Forward primer *TCGCGTTGGTATGCACAACC*

Reverse complimentary sequences are underlined, and same forward sequences are in italic.

A universal reverse PCR primer in all cases is: GTGCAGGGTCCGAGGT.