

Supplementary Information

Construction of plasmid pHH21-5-3U-S1

5'-3' UTR-S1:

agtgttagtctacgtggaccgacaagacagattttgagggagctaagctcaacgttagttcaacagtttttaattagagaggcagatctgtatgaa
taaccaacgaaaaaggcgagaaaatacgccttcaatatgctgaaacgcgagagaaaccgcgttag**GAATT**Caggcaaaactaacatga
aacaaggctagaagtccggattaagccatagtacgaaaaactatgctacctgtgagccccgtccaaggacgtaaaagaagtccggat
tacaaatgccatagcttgactaaactgtgcagccctgtagetccacctgagaaggtgtaaaaatctggaggccacaaaccatggaagctgtacgca
tggcttagtggacttagcggttagaggagaccccccacaaacaaaaacagcatattgacgctggaaagaccagagatcctgcttcctcagcatattccag
gcacagaacgccagaaaatggaatggctgtgaatcaacagg**tct****GGATCCATGGGACCGACCAGAATCATGCAA**
GTGCGTAAGATAGTCGCGGCCGGCCATG

GAATT highlighted with red color is the link of 5' and 3' UTR.

GGATCCATGGGACCGACCAGAATCATGCAAGTGC**GTAAGATAGTCGCGGCCGGCCAT**

G highlighted with yellow color is the sequence of Bio Aptamer S1.

5' UTR-F-T7

TAATACGACTCACTATAAGGAGtgttagtctacgtggac

3' UTR-R-S1:

CATGGCCCAGCCCCGCGACTATCTTACGCACCTGCATGATTCTGGTCGGTCCCATGGATCCa
gaacctgttattcaacagc

The restriction sites of vector pHH21:

CGTCTCNTATTAGTAGAA....TTTGCTCCNGAGACG

GCAGAGNATAA**TCATCTT**....AAA**ACGA**GGGN**CTCTGC**

pHH21-5'-3' U-S1

CGTCTCCTATTAGtgttagtctacgtggaccgacaagacagattttgagggagctaagctcaacgttagttcaacagtttttaattagag
agcagatctgtatgataaccacgaaaaaggcgagaaaatacgccttcaatatgctgaaacgcgagagaaaccgcgttag**GAATT**Ca
ggcaaaactaacatgaaacaaggctagaagtccggattaagccatagtacgaaaaactatgctacctgtgagccccgtccaaggacgtta
aaagaagtccggcattacaaatgccatagcttgactaaactgtgcagccctgtagctccacctgagaaggtgtaaaaatctggaggccacaaac
catggaaagtctgtacgcattggctgtggacttagcggttagaggagaccccccacaaacaaaaacagcatattgacgctggaaagaccagagatcctgctgt
ctcctcagcatcattccaggcacagaacgccagaaaatggatggctgtgaatcaacagg**tct****GGATCCATGGGACCGACCA**
GAATCATGCAAGTGC**GTAAGATAGTCGCGGCCGGCCATG****CCCCGGAGACG**

The primers used to clone the 5'-3' UTR-S1 into pHH21.

5' UTR-F-BsmB:

5'-**CGTCTC**CTATTAGTTGTTAGTCTACGTGGAC-3'

3' UTR-Bio-R-BsmB:

5'-CGTCTCCGGGCATGGCCCGCCCGACTATC-3'

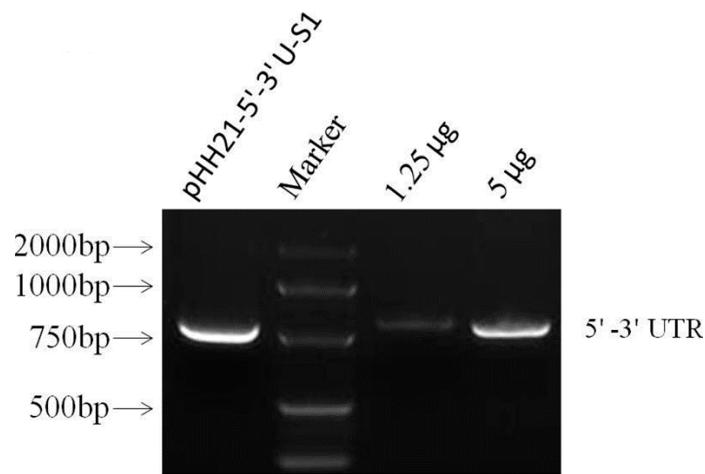


Figure S1. The RNA used in Figure 1D was reverse transcript to cDNA, and then was subjected to PCR using primers of 5' UTR-F and 3' UTR-R. The bands at around 750 bp proved that the DENV-UTR construct is transcribed as one intact RNA.

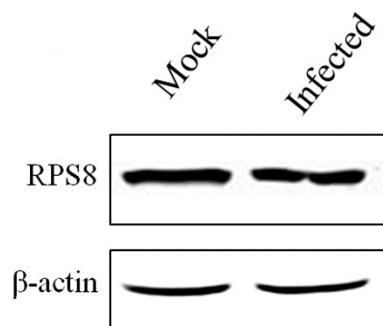


Figure S2. The protein level of RPS8 in DENV mock-infected and infected cells was assayed with western blotting using antibody against RPS8, and β -actin act as internal control. The result showed that there was no statistical difference between the two groups.

Tabel S1. The C_t value of RT-qPCR in the Figure 1D.

Group	β -Actin	5'-3' UTR
1.25 μ g	17.78	18.36
2.5 μ g	17.95	17.96
5 μ g	17.85	17.75