

Electronic Supplementary Material

Development and Evaluation of a Universal and Supersensitive NS1-Based Luciferase Immunosorbent Assay to Detect Zika Virus-Specific IgG

Tianyu Wang^{1,2#}, Ying Zhan^{2.#}, De Wu^{3#}, Zihai Chen^{4#}, Wei Wu², Yao Deng², Wenling Wang², Wenjie Tan^{2✉}, Shixing Tang^{1✉}

1.Guangdong Provincial Key Laboratory of Tropical Disease Research, Department of Epidemiology, School of Public Health, Southern Medical University, Guangzhou 510515, China.

2.NHC Key Laboratory of Biosafety, National Institute for Viral Disease Control and Prevention, China CDC, Beijing 102206, China.

3.Key Laboratory for Repository and Application of Pathogenic Microbiology, Research Center for Pathogens Detection Technology of Emerging Infectious Diseases, Guangdong Provincial Center for Disease Control and Prevention, Guangzhou 511430, China.

4.The National Clinical Key Department of Infectious Disease, Beijing Ditan Hospital, Capital Medical University, Beijing 100015, China.

Supporting information to DOI: 10.1007/s12250-019-00160-x

Table S1. Primers used for NS1 amplification in this study

Target gene	Primer name	Primer sequence (5'-3')	Size	Restriction site
NS1	Full-F	CGGAATTCGACGTGGGCTGCTCCGTGGA	1056bp	<i>EcoR</i> I
	Full-R	GCTCTAGAGGCGGTCACCATGCTCCGCA		<i>Xba</i> I
NS1-N	N-F	CGGAATTCGACGTGGGCTGCTCCGTGGA	513bp	<i>EcoR</i> I
	N-R	GCTCTAGACACCTTCAGCCACACGCTGG		<i>Xba</i> I
NS1-C	C-F	CGGAATTCGAGGACTACAGCCTGGA	543bp	<i>EcoR</i> I
	C-R	GCTCTAGAGGCGGTCACCATGCTCCGCA		<i>Xba</i> I
NS1-C1	C1-F	CGGAATTCGAGGACTACAGCCTGGAG	210bp	<i>EcoR</i> I
	C1-R	GCTCTAGATTACAGATCGGACTCCTCCACGCCGT		<i>Xba</i> I
NS1-C2	C2-F	CGGAATTCCTGTGGACAGACGGCGTGGAG	210bp	<i>EcoR</i> I
	C2-R	GCTCTAGATTAGCTTCTCAGAGAGGGGCCCGTG		<i>Xba</i> I
NS1-C3	C3-F	CGGAATTCACCAAGGTGTATGTGGAGGAG	210bp	<i>EcoR</i> I
	C3-R	GCTCTAGATTAGGCGGTCACCATGCTCCGCACCA		<i>Xba</i> I