

## Electronic Supplementary Information

Table 1. Comparison of RT-PCR and POC NAT System in Testing Clinical Samples

	H1pdm 2009 Specific Primer *		Influenza A Consensus Primer **		Housekeeping Gene Primer ***		
	Positive	Negative	Positive	Negative	Positive	Negative	
Multiple-Gene Detection test (45 samples)	23	22	23	22	44	1	RT-PCR
	21 <sup>a</sup>	24	21 <sup>a</sup>	24	43 <sup>a</sup>	2	POC NAT System
Sensitivity test (42 samples)	22	20	22	20	42	0	RT-PCR
	20 <sup>b</sup>	22	N.D.	N.D.	16 <sup>c</sup>	26	POC NAT System

\* The H1pdm-specific primer detects the HA gene in RT-PCR.

\*\* The influenza A consensus primer detects the matrix protein gene in RT-PCR.

\*\*\* The housekeeping gene primer detects the human RNase P gene in RT-PCR and the human  $\beta$ -actin in the POC NAT System.

a “Positive” was defined as the detection of a nucleic acid amplification signal in at least one of three wells within 30 min.

b “Positive” was defined as the detection of a nucleic acid amplification signal in at least one of eight wells within 30 min.

c Because we divided and diluted one swab sample for comparison between two methods in this experiment, the amount of sample was insufficient to detect human  $\beta$ -actin gene which was tested in only one well.