

# **A lab-on-chip device for sample-in-result-out detection of viable *Salmonella* using loop-mediated isothermal amplification and real-time turbidity monitoring**

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**Table S1.** Nucleotide sequences of LAMP primers

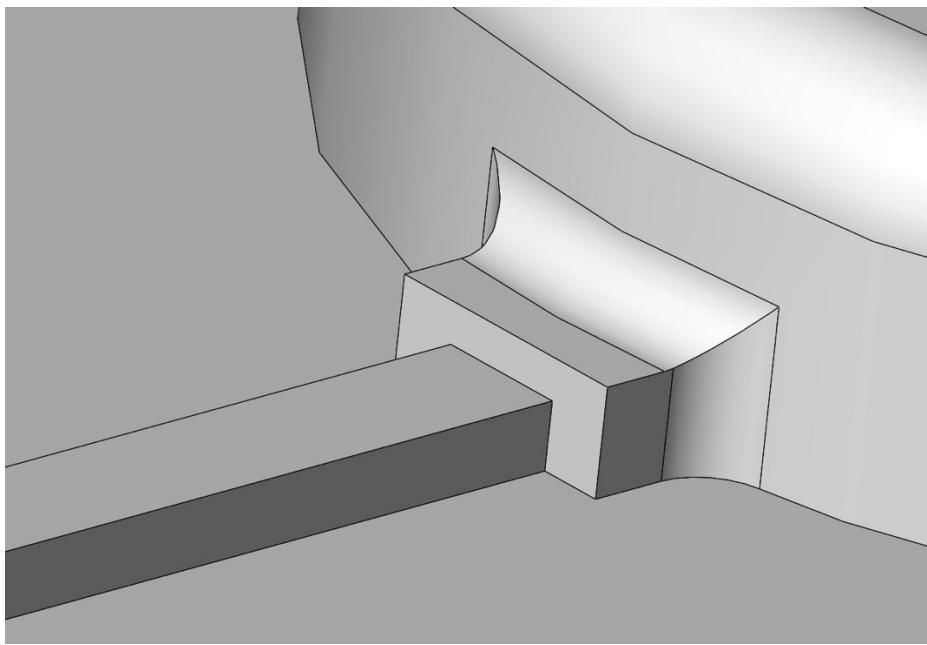
Primer Type	Primer Name	Nucleotide Sequence
Inner primer 1	F3	5'-CGGCCCGATTTCTCTGG-3'
Inner primer 2	B3	5'-CGGCAATAGCGTCACCTT-3'
Outer primer 1	FIP	5'-GCGCGGCATCCGCATCAATA-TGCCCGTAAACAGATGAGT-3'
Outer primer 2	BIP	5'-GCGAACGGCGAAGCGTACTG-TCGCACCGTCAAAGGAAC-3'
Loop primer 1	LF	5'- GGCCTTCAAATCGGCATCAAT-3'
Loop primer 2	LB	5'-GAAAGGGAAAGCCAGCTTACG-3'

**Table S2.** Comparison of this proposed device with previously reported detection methods

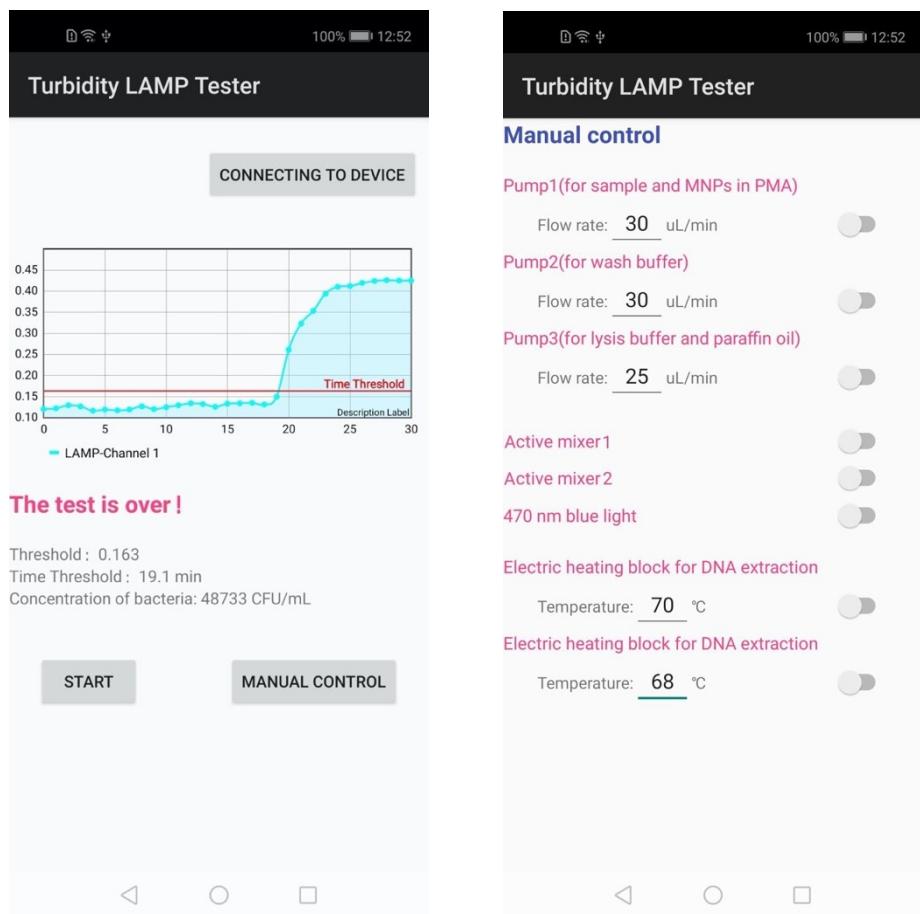
Method	Target Bacteria	Detection Limit	References
PCR	<i>Bacillus cereus</i>	$9.2 \times 10^1$ CFU/mL	1
ELISA	<i>Salmonella/E.coli/Vibrio choera</i>	$1.0 \times 10^4$ CFU/mL	2
Plate count	<i>Enterobacter sakazakii</i>	$1.0 \times 10^2$ CFU/mL	3
Biosensor	<i>Staphylococcus aureus</i>	$1.0 \times 10^1$ CFU/mL	4
LAMP	<i>Salmonella spp./E. coli O157:H7</i>	$3.0 \times 10^1$ CFU/sample	5
LAMP	<i>Salmonella typhimurium</i>	$1.4 \times 10^1$ CFU/mL	This study

## References

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**Fig S1.** The connection between channel and chamber



**Fig S2.** The interfaces of the smartphone App