

## Electronic Supplementary Information

# Fully automated and colorimetric foodborne pathogen detection on an integrated centrifugal microfluidic device

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**Table S1.** Primer sequence information for the multiplexed LAMP reaction

Target gene / Accession #	Primer	Sequence (5' to 3')	Length [bp]
<i>E. coli</i> O157:H7 fliC gene (CP002967.1 : 2147323-2149017)	FIP	CTACCAACCTGTCTGAAGCGCATCGACATGTTGGACACTTCG	42
	BIP	GGTGACTGCGGAATCCAGACGTGAGGCAATTGCATCCATCG	41
	F3	GCCTGCTGGATGATCTGC	18
	B3	CGGCTCTGCAACCAAAGA	18
<i>S. Typhimurium</i> invA gene (M90846.1)	FIP	TCCGCGACACGTTTCTGAACCCTCTATTGTCACCGTGGTCC	40
	BIP	TGCCGATTTGAAGGCCGGTACAGTACGCTTCGCCGTTTC	38
	F3	GCGGTGGGTTTTGTTGTCT	19
	B3	CGTAAAGCTGGCTTTCCCTT	20
<i>V. parahaemolyticus</i> toxR gene (L11929.1)	FIP	TCGACTCCACATTCACTCGATTACTGATAACTTGCCAGACG	41
	BIP	CCTGCCGAATGGCGATTACAATTAATTGTCGATTGAGCCG	40
	F3	CAACCATGGTGACTGTGA	18
	B3	CCAGCGACCTTTCTCTGA	18
<i>L. monocytogenes</i> iap gene (JQ015300.1)	FIP	ATCAAATGTAGTTGGTCCGTTACCAGCTGAAGCTCAAAAACACC	44
	BIP	ATGTATTTGCTAAAGCGGGAATCTGTAGTGCTAGCGTATTGTGC	44
	F3	AATTCAAGTGCAAGTGCTAT	20
	B3	GTTTTGCTTGAGATTCAGAGA	21



**Fig. S1.** Singleplex *Listeria monocytogenes* detection on the lab-on-a-disc. A real sample containing  $10^4$  cells of *L. monocytogenes* was used. Colour change was only observed in the reaction chamber #4, where the LAMP primer set targeting *L. monocytogenes* had been pre-stored, which confirmed specific detection of that bacteria species.



**Fig. S2.** Negative control experiments using a real sample spiked with non-pathogenic *Escherichia coli* K-12. No colour change was shown in the reaction chambers. This result proves the ability of this assay to distinguish bacteria sub-types within *E. coli* species.