## **Supporting Information for**

## Tryptamine-functionalized magnetic nanoparticles for highly

## sensitive detection of Salmonella Typhimurium

Target	Sequences (5'-3')	T <sub>m</sub> (°C)	Product size (bp)	
S. Typhimurium-Forward	5'-TATCGCCACGTTCGGGCAA-3'	59.7	275	
S. Typhimurium-Reverse	5'-TCGCACCGTCAAAGGAACC-3'	58.1	275	

**Table S1.** Specific sequence and primers we used to identify DNA of S. Typhimurium.



Figure S1. A TEM image of Indole@MNPs



**Figure S2**. Evaluation of interference of milk during sample preparation using commercial kit. No Cq values were obtained after amplification of no-template controls (NTCs) in any experiments. Error bars indicate standard deviation from mean based on three independent experiments.



**Figure S3.** Capture efficiencies of Indole@MNPs with concentrations of *S*. Typhimurium (1000, 100, 10 CFU in 10 mL PBS buffer and milk). The original samples without binding with Indole@MNPs were counted as 100 % and served as controls. Error bars indicate standard deviation from mean based on three independent experiments.



**Figure S4**. Evaluation of different Indole@MNP amounts in 10 mL PBS sample using 10<sup>3</sup> CFU of *S*. Typhimurium. Error bars indicate standard deviation from mean based on three independent experiments.



**Figue S5.** Gel electrophoresis image of PCR product. 25/100 bp mixed DNA ladder (Bioneer Inc.); NC: negative control with no DNA template; Lane 1: 1000 CFU/10 mL sample; Lane 2: 100 CFU/10 mL sample; Lane 3: 10 CFU/10 mL sample; Lane 4: 5 CFU/10 mL sample.

## **DLVO model**

The potential interaction energy profile between *S*. Typhimurium and Indole@MNP was calculated based on the DLVO theory. [1-4] Following equations were used to make the computation:

$$V_{Tot} = V_{LW} + V_{EL}$$

$$V_{LW} = -\frac{A(a_1a_2)}{6d(a_1 + a_2)}$$

$$V_{EL} = \frac{\pi \epsilon a_1 a_2(\zeta_1^2 + \zeta_2^2)}{(a_1 + a_2)} \left[ \frac{2\zeta_1\zeta_2}{\zeta_1^2 + \zeta_2^2} ln \frac{1 + exp[in](-kd)}{1 - exp[in](-kd)} + ln[in]\{1 - exp(-2kd)\} \right]$$

 $V_{LW}$  = Electrical double layer repulsive energy

 $V_{EL} =$  Van-der-Waals attractive energy

 $V_{Tot} = DLVO$  energy barrier

 $A = \text{Hamaker constant where } A_{132} = (\sqrt{A_1} - \sqrt{A_3})(\sqrt{A_2} - \sqrt{A_3}) = 1.40 \times 10^{-21} J \text{ for the interaction between bacterial pathogen } (A_1 = 5.2 \times 10^{-20} J) \text{ and iron oxide nanoparticle } (A_2 = 10 \times 10^{-21} J) \text{ separated by a water medium } (A_3 = 3.7 \times 10^{-20} J).$ 

 $a_1 =$  Radii of microbial cells; 770 nm for *Salmonella* Typhimurium

 $a_2$  = Radii of Indole@MNP (iron oxide nanoparticle) = 425 nm

d = Separation distance: 0–3000 nm (dependent variable)

 $\varepsilon = \varepsilon_0 \times \varepsilon_r = 7.083 \times 10^{-10} F/m$  where the relative permittivity of the medium,  $\varepsilon_r = 80$  (for water at 20°C) and ther permittivity of a vacuum,  $\varepsilon_0 = 8.854 \times 10^{-12} F/m$ 

 $\zeta_1 =$  Zeta potential of bacteria where -11.7 mV for *Salmonella* Typhimurium

$$\zeta_2$$
 = Zeta potential of Indole@MNPs (pH 7)= -32 mV

k = Inverse Debye-Huckel length = 729 nm for pH 6.5

References:

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**Table S2.** An overview on recently reported MNP-based methods for the detection of *Salmonella* species.

Materials used	Detection method	Detection limit	References	
Antibody functionalized MNPs	Multiplex PCR	10 CFU/ml (Pure culture)	culture) [1] at)	
		100 CFU/ml (Meat)		
Antibody functionalized	Fluorescence (DNA-	13.6 CFU/ml (PBS)	[2]	
MNPs	QD-AuNP)			
Antibody functionalized	Electrochemistry	10 CFU/ml (Milk)	[3]	
MINP chains	-			
Aptamer functionalized MNPs	Chemiluminescence			
	(Rolling circle	10 CFU/mL (PBS)	[4]	
	amplification)			
Aptamer functionalized	SERS	15 CFU/mL (PBS)	[5]	
MNPS				
Oligonucleotide	RT-PCR	10 CFU/mL (Milk)	[6]	
iunctionalized MINPs				
Indole functionalized	qPCR	10 CFU/10 mL (Milk)	This study	
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