# Rapid screening and quantitative detection of Salmonella using a quantum dot nanobead based biosensor

Jiao Hu,\*a,c Feng Tang, d Yong-Zhong Jiang e and Cui Liu\*b

<sup>a</sup>·Hubei Key Laboratory of Environmental and Health Effects of Persistent Toxic Substances, Institute of Environment and Health, Jianghan University, Wuhan, 430056, China. E-mail: hujiao@whu.edu.cn.

<sup>b.</sup>Institute of Medical Engineering, Department of Biophysics, School of Basic Medical Sciences, Xi'an Jiaotong University Health Science Center, Xi'an, 710061, China. E-mail: liucui.tree@163.com.

<sup>c</sup>Wuhan Academy of Agricultural Sciences, Wuhan, 430072, China.

<sup>d.</sup>Department of Laboratory Medicine, Wuhan Children's Hospital (Wuhan Maternal and Child Healthcare Hospital), Tongji Medical College, Huazhong University of Science & Technology, Wuhan, 430016, China.

<sup>e.</sup>Hubei Provincial Center for Disease Control and Prevention, Wuhan, 430072, China.

## S1. Evaluating the fluorescence of QDNS

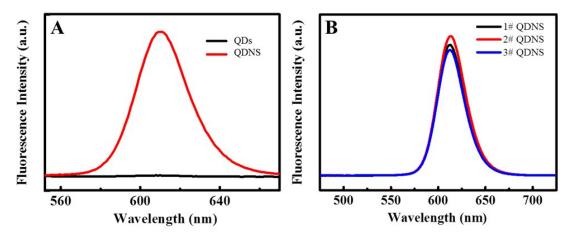


Fig. S1 (A) Fluorescence intensity of QDNS (read line, fabricated by layer-by-layer assembly method), quantum dots (black line) at the same particle concentrations. (B) Fluorescence intensity of the three batches of QDNS.

## S2. Verification of the fabrication of antibody conjugation

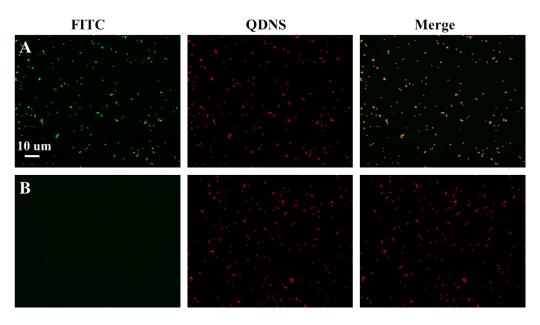


Fig. S2 The fluorescence response of IQDNS (A) and QDS (B) after reacted with FITC-goat antimouse IgG, respectively.

# S3. Optimization of the detection conditions

In order to realize highly sensitive detection of *Salmonella typhimurium*, the amount of QDNS was optimized. As we know, the signal of positive sample was increased with the amounts of QDNS within a certain range. Then, the signal was decreased due to the hook effect. As shown in Figure S2, the fluorescence intensity of

test line was the highest at 30  $\mu g$  QDNS. Hence, 30  $\mu g$  QDNS was ascertained as the optimal dose of one test.

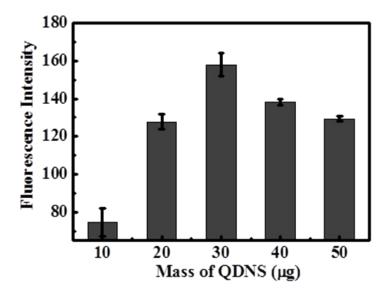


Fig. S3 Histogram of fluorescence intensity of test line with different amount of QDNS for the detection of *Salmonella typhimurium* (The *Salmonella typhimurium* concentrations were 10<sup>7</sup> CFU/mL).

## S4. Reproducibility analysis of the QDNS based biosensor

For practical application, good precision and reproducibility are very important. Different concentrations of *S. typhi* broth samples were used to evaluate the reproducibility. As shown in Table S1, the intra-assay and inter-assay variability was calculated to be 4.99% and 7.15%, respectively. The results suggested that the QDNS based biosensor exhibited high reproducibility.

Table S1. Reproducibility analysis of the QDNS based biosensor

	Intra-assay			Inter-assay		
S. typhi concentration (CFU/mL)	mean <sup>a</sup>	$SD^b$	CV <sup>c</sup> (%)	mean <sup>a</sup>	$SD^b$	CV <sup>c</sup> (%)
5×10 <sup>4</sup>	45.8	2.59	5.66	41.2	3.95	9.59
1×10 <sup>5</sup>	60.8	4.10	6.74	69	5.23	7.58
5×10 <sup>5</sup>	135.2	4.86	3.59	138.6	8.02	5.79
1×10 <sup>6</sup>	164.2	6.53	3.98	163.4	9.18	5.62
	Intra	a-assay variability		Inter-assay variability		
	4.99			7.15		

<sup>&</sup>lt;sup>a</sup>Values represent the average of detected fluorescence intensity of parallel samples (n=5). <sup>b</sup>Values represent the standard deviation of parallel results (n=5). <sup>c</sup>CV=SD/mean.