## **Electronic Supporting Information**

## A competitive, bead-based assay combined with microfluidics for multiplexed toxin detection

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**Figure S1** Fluorescence images of retained beads in the collection chambers. The images show the decrease in the fluorescence signal of beads from left to right by increasing the target toxin concentration.



**Figure S2** Monitoring the signal stability of the assay. The images were taken from thre beads retained in the chamber in a three-week period.



**Figure S3.** A) The chamber of Toxin-Chip where magnetic beads were collected. B) The chamber after washing. C) the herringbone structure after washing.



Figure S4. Fluorescence images of detecting injected toxin from lake water samples.

## Table S1 Toxin-Chip dimensions

Geometry	Dimensions	
Mixing module length	30 cm	
Mixing module width	300 µm	
Mixing module height	45 μm	
Herringbone structures height	45 μm	
Chamber diameter	1.5 mm	
Chamber height	45 μm	

## Table S2 Microscope setting for collecting the fluorescence images

Setting	Value
Magnification	30X
Exposure time	10ms
Lookup Tables (LUT)	100-2500

**Table S3** Spiked and calculated concentrations of toxin related to Figure 6.

Spiked conc.	Calculated MC-LR conc.	Calculated OA conc.
Sample 1- 0 µg/ml MC-LR, 0 µg/ml OA	0.0008 µg/ml	0.00075 μg/ml
Sample 2- 1 µg/ml MC-LR, 0 µg/ml OA	1 μg/ml	0.00001 μg/ml
Sample 3- 0 µg/ml MC-LR, 1 µg/ml OA	0 μg/ml	0.75 μg/ml
Sample 4- 1 µg/ml MC-LR, 1 µg/ml OA	0.85 µg/ml	1 μg/ml

	Table S4	Spiked and recovered	concentration from	lake water experiment.
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Spiked conc.	0 μg/ml	1 μg/ml
MC-LR	0.000028 µg/ml	0.76 µg/ml
error	12%	10%
OA	0 μg/ml	0.87 µg/ml
error	8%	17%