

## Supplementary Information

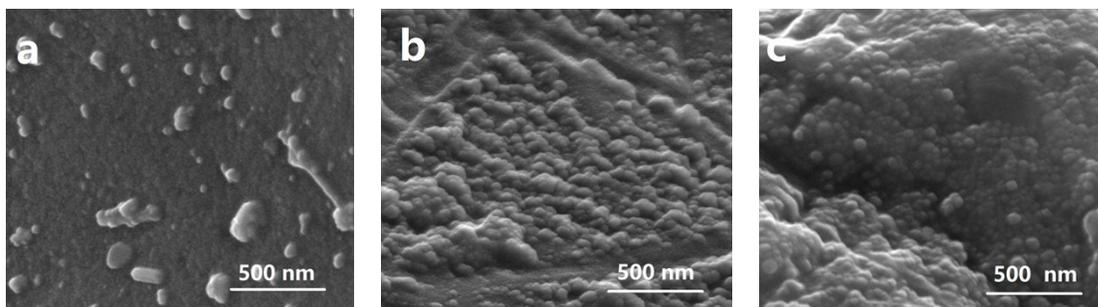
### Dual functional PDMS sponge SERS substrate for the on-site detection of pesticides both on fruit surfaces and in juice

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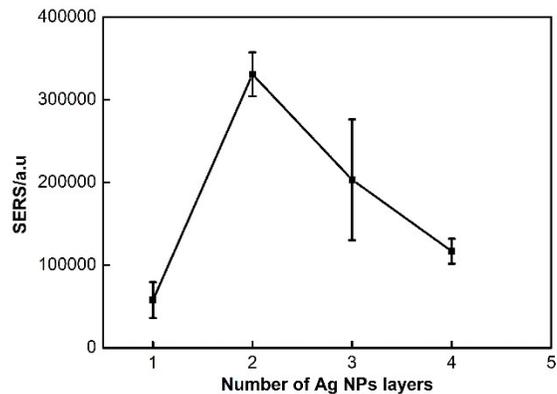
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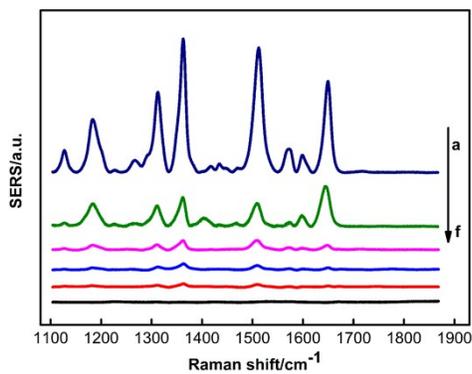
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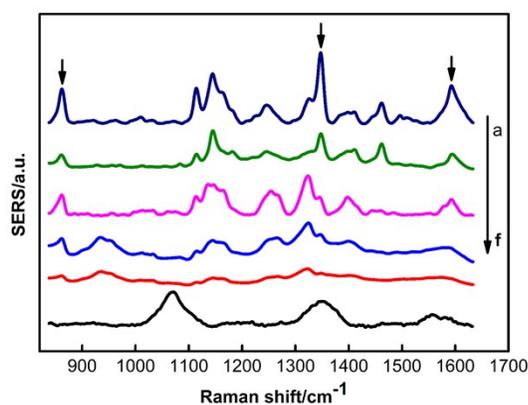
**Fig. S1** SEM images of different rounds of Ag NPs deposited on PDMS sponge. a-c represented one, two and three rounds of Ag NPs deposition, respectively.



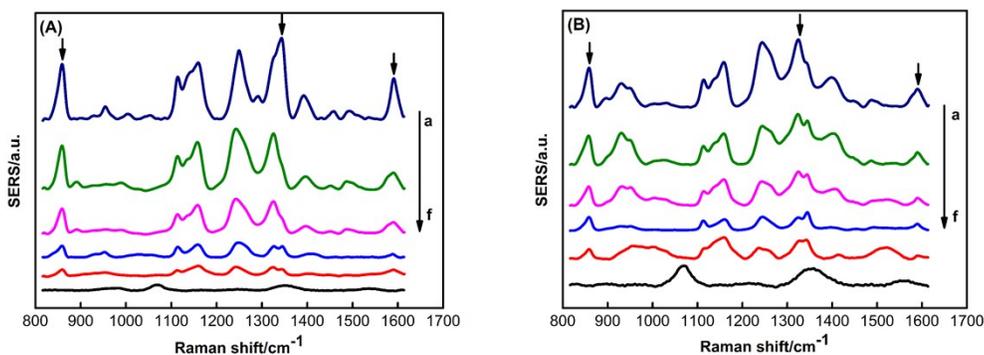
**Fig. S2** SERS performance of different Ag NPs rounds-modified PDMS sponge. 20  $\mu$ L triazophos (50 ppm) was used as the Raman probe. Laser power: 14 mW. Integration time: 50 s. Objective: 50 $\times$ .



**Fig. S3** SERS spectra of R6G on the PDMS sponge-SERS substrate. The concentrations of R6G were: (a-f) 100 nM, 10 nM, 5 nM, 1 nM, 0.1 nM, 0 nM, respectively. Laser power: 10 mW. Integration time: 5 s. Objective: 20 $\times$ .



**Fig. S4** SERS spectra of methyl parathion from the surface of glass. The concentration of methyl parathion were: (a-f) 50 ppm, 10 ppm, 5 ppm 1 ppm, 0.1 ppm, 0 ppm, respectively. Laser power: 10 mW. Integration time: 50 s. Objective: 20 $\times$ .



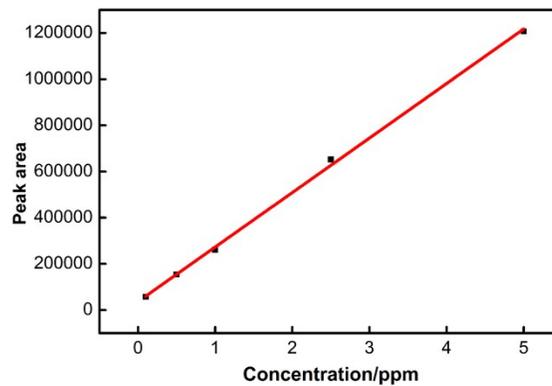
**Fig. S5** SERS spectra of methyl parathion from the surface of Cherry tomatoes (A) and Plum (B). The concentrations of methyl parathion were: (a-f) 50 ppm, 10 ppm, 5 ppm, 1 ppm, 0.1 ppm, 0 ppm, respectively. Laser power: 10 mW. Integration time: 50 s. Objective: 20 $\times$ .

Samples were analyzed by HPLC on LiChrospher®100 RP 18 column and UV-detector. Methanol and water were used as mobile phase, at a flow rate of 0.8 mL/min with 15 min and detection at 254 nm. The gradient elution profile of methanol and water was programmed as methanol/water = 70:30 (v/v). The

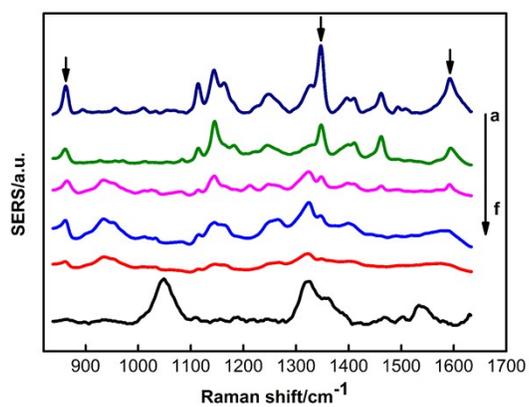
column oven was set at 28 °C, and the injection volume was 20 μL. The found value of the pesticides in the sponge SERS substrate was then plugged into equation 3 to determine the sampling efficiency.

100 ppm triazophos was prepared by using methanol as solvent, then this solution was diluted to 10 ppm, 5 ppm, 2.5 ppm, 1ppm, 0.5ppm, 0.1 ppm. These samples were used for the construction of calibration curve.

The standard curve for HPLC in Fig. S6 showed that, the fitting equation is  $y = ax + b$ , ( $a = 236214$ ,  $b = 36373$ ,  $R^2 = 999$ ). The data of the three parallel tests were averaged, and the absolute amount of triazophos from Cherry Tomatoes and Plum obtained by sticking to the HPLC standard curve.



**Fig. S6** HPLC operating curves for different concentrations of triazophos used for recovery detection.



**Fig. S7** SERS spectra of methyl parathion in carrot juice. The concentrations of methyl parathion were: (a-f) 100 ppm, 50 ppm, 10 ppm, 5 ppm, 1 ppm, 0 ppm, respectively. Laser power: 10 mW. Integration time: 50 s. Objective: 20 $\times$ .