

Supporting Information

Horseradish peroxidase-repeat assay based on tyramine signal amplification for highly sensitive H₂O₂ detection by Surface-Enhanced Raman Scattering

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Optimization of Experimental Conditions

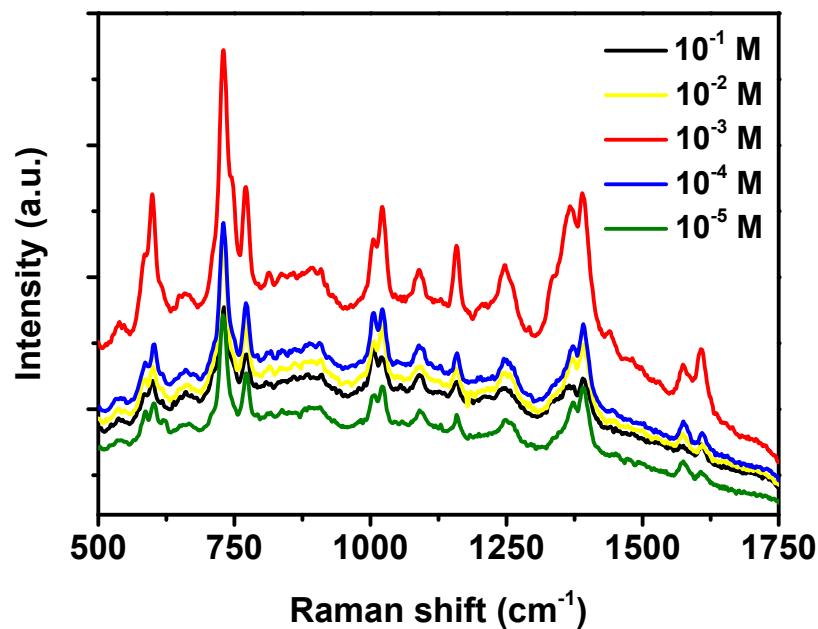


Fig. S1. SERS spectra of the sensing systems with different cysteamine concentrations.

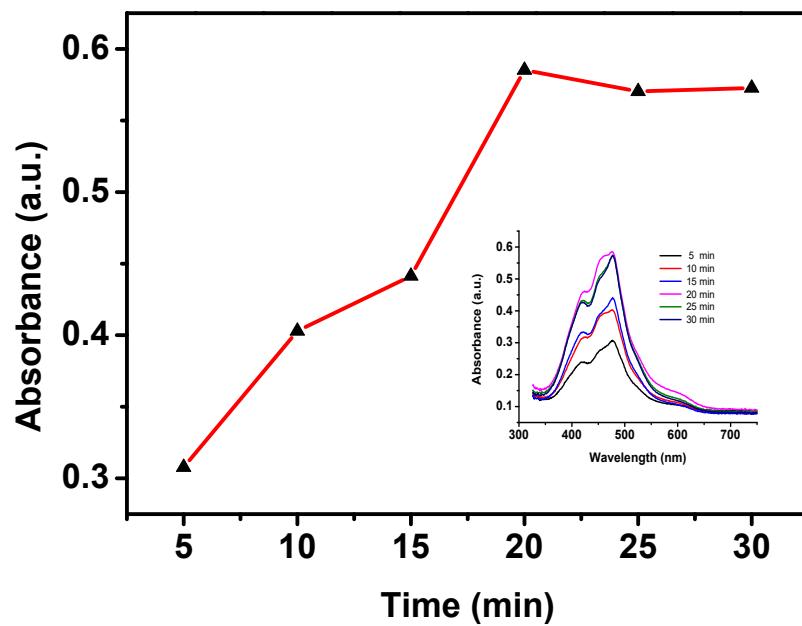


Fig. S2. UV-Vis spectra of the catalytic reaction with different incubation times for Au-HRP conjugation.

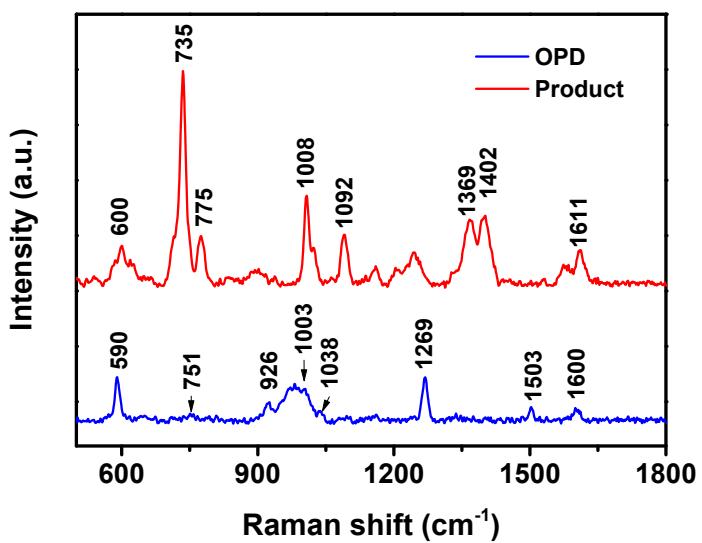


Fig. S3. SERS spectra of OPD and the product DAP.

Table S1. Main SERS Band Assignments of OPD and DAP

Vibrational bands/cm ⁻¹		Assignment
OPD	DAP	
590	600	ring deformation
751	735	C-H wagging
	775	ring deformation, ring stretching
1003	1008	ring stretching and in-plane bending
		ring deformation, ring breathing
1038		
1269	1369, 1402	C-N stretching in the benzenoid C-N ⁺ -stretching, C=N stretching
1503		ring deformation, C-H in-plane bending
1600	1611	NH ₂ scissoring

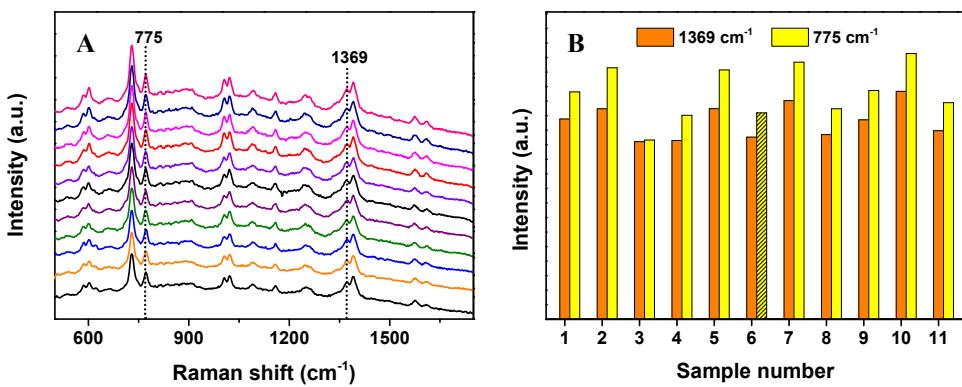


Fig. S4. (A) SERS spectra with Au-re-HRP assay in the substrate solution of 0.1 mM H_2O_2 from 11 different sensors. (B) The intensities of two different bands from 11 trials.

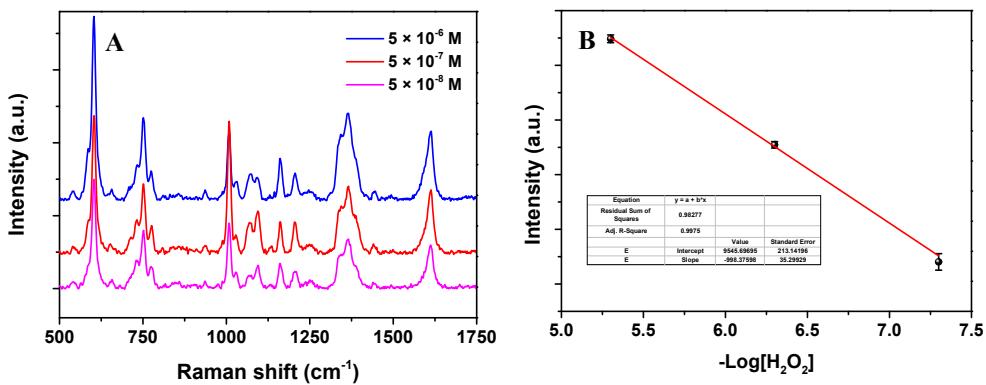


Fig. S5. (A) SERS spectra of samples with different H_2O_2 concentrations in serum samples. (B) Quantitative analysis plot showing the relationship between band intensity at 1366 cm^{-1} and the logarithm of H_2O_2 concentration. The error bar indicates three independent measurements.