

Electronic Supplementary Information for

High-index {hk0} faceted platinum concave nanocubes with enhanced peroxidase-like activity for ultrasensitive immunoassay

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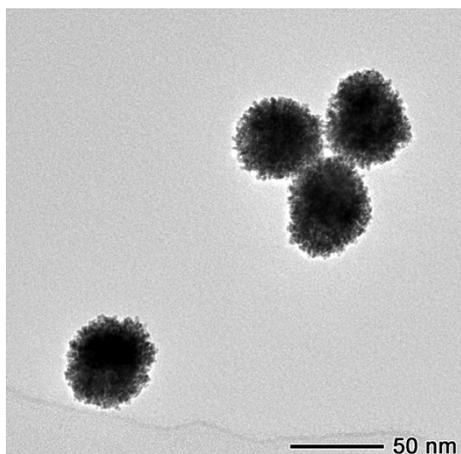


Fig. S1 TEM image of ~ 48.0 nm Pt nanospheres.

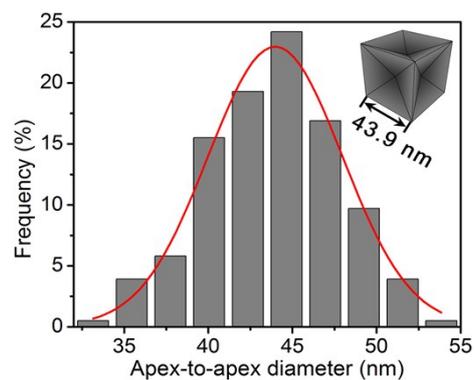


Fig. S2 Distribution of apex-to-apex diameter of the as-synthesized HIF-Pt-CNCs based on counting more than 200 particles by using “NANO MEASURER software”.

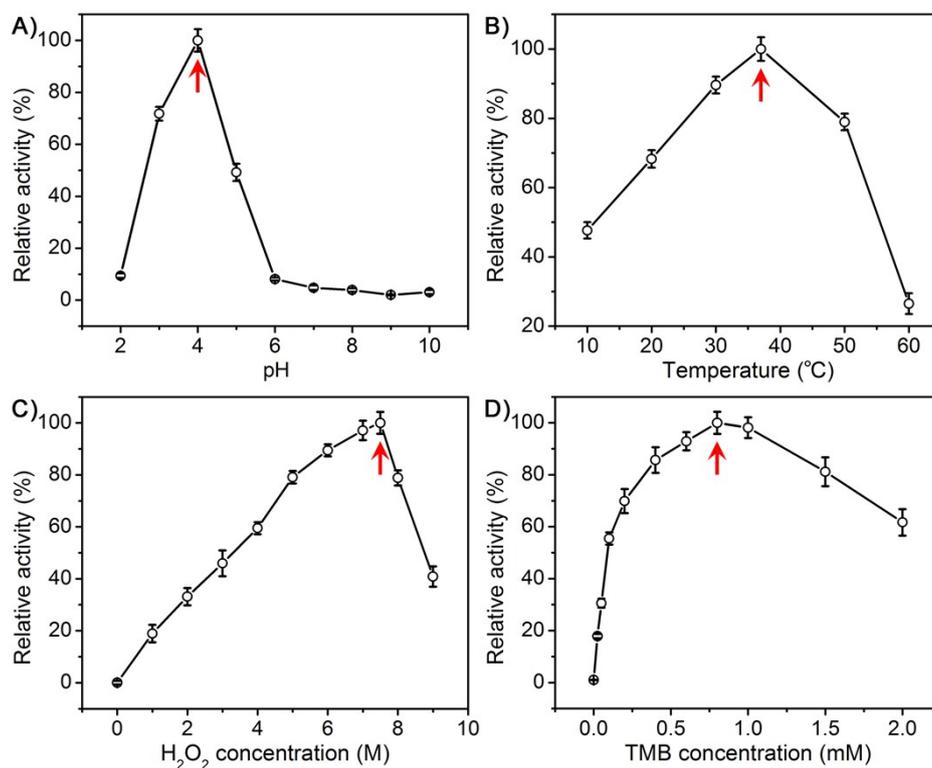


Fig. S3 The effect of pH (A), temperature (B), H₂O₂ concentration (C), and TMB concentration (D) on the peroxidase-like catalytic activity of the HIF-Pt-CNCs. The maximum point in each curve is set as 100%. Error bars indicate standard deviations ($n = 3$).

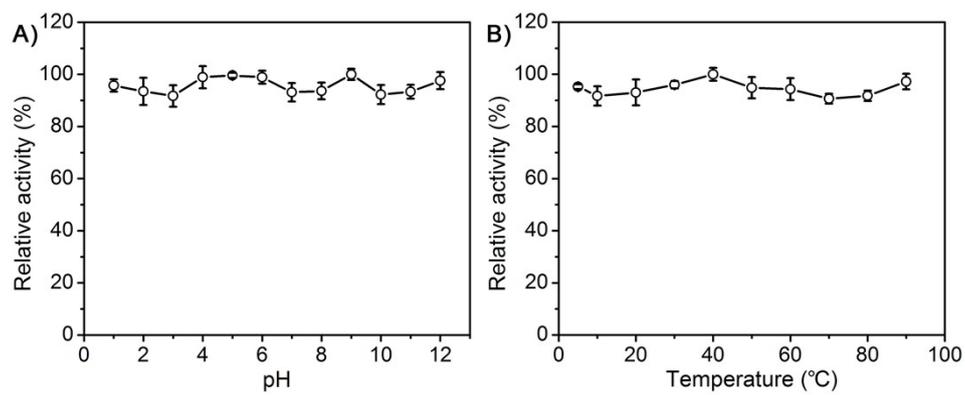


Fig. S4 Relative catalytic activities of HIF-Pt-CNCs after incubation for 2 h at various pH values (A) and temperatures (B), respectively. The maximum point in each curve is set as 100%. Error bars indicate standard deviations ($n = 3$).

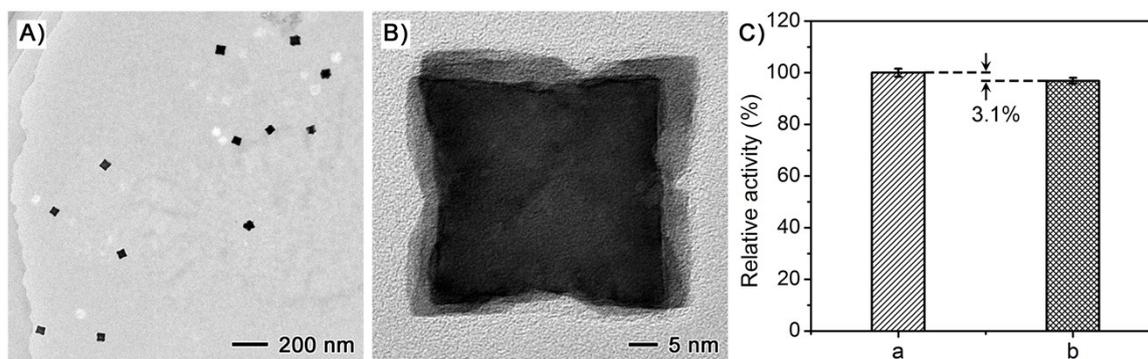


Fig. S5 (A, B) Regular TEM image (A) and HRTEM image (B) of the DAb-conjugated HIF-Pt-CNCs after the colorimetric immunoassay. (C) Relative catalytic activities of the DAb-conjugated HIF-Pt-CNCs before (a) and after (b) the colorimetric immunoassay. The activity before the colorimetric immunoassay is set as 100%. Error bars indicate standard deviations ($n = 3$).

The experiments were carried out as follows: i) after the colorimetric immunoassay of 300 PSA standards (20 ng mL^{-1} as an example), the detection solutions in the wells of the plates were discarded, and the wells were washed three times with water; ii) $100 \mu\text{L}$ of 0.1 M Glycine-HCl buffer ($\text{pH } 2.3$) was added into the wells to dissociate the antibody-antigen complexes and release the DAb-conjugated HIF-Pt-CNCs;^{S1} iii) the released DAb-conjugated HIF-Pt-CNCs in the buffer were collected and washed with water twice *via* centrifugation, and the collected DAb-conjugated HIF-Pt-CNCs were dispersed in 1 mL of water; iv) the structure and morphology of these collected HIF-Pt-CNCs were characterized by TEM; and v) the catalytic activities of these collected HIF-Pt-CNCs and the unused DAb-conjugated HIF-Pt-CNCs toward H_2O_2 -TMB system were determined and compared under the same conditions including the same Pt concentration (Please note that the concentrations of Pt element in the collected HIF-Pt-CNCs suspension and the unused DAb-conjugated HIF-Pt-CNCs suspension were determined to be 1.7 mg L^{-1} and 36.2 mg L^{-1} , respectively, by ICP-OES.).

Fig. S5A and B show the TEM images of the collected HIF-Pt-CNCs. It can be seen that the structure and morphology of these collected HIF-Pt-CNCs were almost the same as those of the original HIF-Pt-CNCs (Fig. 1C and D), indicating a good structural stability of HIF-Pt-CNCs during the detection process. Also the catalytic activity of the collected HIF-Pt-CNCs only decreased 3.1% compared to the activity obtained by the unused DAb-conjugated HIF-Pt-CNCs, suggesting a good catalytic stability of HIF-Pt-CNCs during the detection process (Fig. S5C).

Table S1. Projection angles and geometrical parameters of concave nanocubes bounded by different types of high-index facets.^{S2}

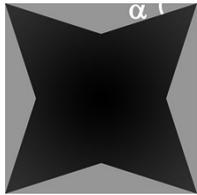
projection direction	geometrical model of the polyhedron	equation for the projection angle	calculated projection angle	
			(hkl)	α
[001]		$\alpha = \arctan(k/h)$	(410)	14.0°
			(720)	15.9°
			(310)	18.4°
			(830)	20.6°
			(520)	21.8°
			(730)	23.2°

Table S2. Comparison of the kinetic parameters of various Catalysts toward the oxidation of TMB by H₂O₂.^a

catalyst	[E] ₀ (M)	substance	V _{max} (M s ⁻¹)	K _{cat} (s ⁻¹)	references
HRP	2.5 × 10 ⁻¹¹	TMB	1.0 × 10 ⁻⁷	4.00 × 10 ³	S3
	2.5 × 10 ⁻¹¹	H ₂ O ₂	8.71 × 10 ⁻⁸	3.48 × 10 ³	
Fe ₃ O ₄ nanoparticles	1.14 × 10 ⁻¹²	TMB	3.44 × 10 ⁻⁸	8.58 × 10 ⁴	S3
	1.14 × 10 ⁻¹²	H ₂ O ₂	9.78 × 10 ⁻⁸	3.02 × 10 ⁴	
Co ₃ O ₄ nanoparticles	3.43 × 10 ⁻¹⁰	TMB	6.27 × 10 ⁻⁸	1.83 × 10 ²	S4
	3.43 × 10 ⁻¹⁰	H ₂ O ₂	1.21 × 10 ⁻⁷	3.53 × 10 ²	
Fe ₂ O ₃ nanoparticles	3.09 × 10 ⁻¹⁰	TMB	1.06 × 10 ⁻⁶	3.43 × 10 ³	S5
	3.09 × 10 ⁻¹⁰	H ₂ O ₂	1.17 × 10 ⁻⁶	3.79 × 10 ³	
MnO ₂ nanoparticels	3.01 × 10 ⁻⁸	OPD	8.21 × 10 ⁻⁸	2.73	S6
	3.01 × 10 ⁻⁸	H ₂ O ₂	5.71 × 10 ⁻⁸	1.90	
Au@Pt nanorods	1.25 × 10 ⁻¹¹	TMB	1.81 × 10 ⁻⁷	1.40 × 10 ⁴	S7
	N/A	N/A	N/A	N/A	
Au@Pd nanoparticles	9.60 × 10 ⁻¹¹	TMB	2.00 × 10 ⁻⁶	2.10 × 10 ⁴	S8
	9.60 × 10 ⁻¹¹	H ₂ O ₂	4.40 × 10 ⁻⁶	4.60 × 10 ⁴	
Pt nanoparticles	8.12 × 10 ⁻¹¹	TMB	1.26 × 10 ⁻⁶	2.27 × 10 ⁴	S9
	8.12 × 10 ⁻¹¹	H ₂ O ₂	1.85 × 10 ⁻⁶	1.55 × 10 ⁴	
Pd nanocubes	1.40 × 10 ⁻¹²	TMB	9.70 × 10 ⁻⁸	6.90 × 10 ⁴	S10
	1.40 × 10 ⁻¹²	H ₂ O ₂	6.50 × 10 ⁻⁸	4.60 × 10 ⁴	
Pd-Ir nanocubes	3.40 × 10 ⁻¹⁴	TMB	6.50 × 10 ⁻⁸	1.90 × 10 ⁶	S10
	3.40 × 10 ⁻¹⁴	H ₂ O ₂	5.10 × 10 ⁻⁸	1.50 × 10 ⁶	
HIF-Pt-CNCs	2.54 × 10 ⁻¹⁴	TMB	1.52 × 10 ⁻⁷	5.98 × 10 ⁶	Present work
	2.54 × 10 ⁻¹⁴	H ₂ O ₂	1.29 × 10 ⁻⁷	5.08 × 10 ⁶	

^a [E]₀ is the catalyst concentration, K_m is the Michaelis constant, V_{max} is the maximal reaction rate, and K_{cat} is the catalytic constant, where K_{cat}=V_{max}/[E]₀.

REFERENCE

