

Supporting information

An ultrasensitive electrochemical aptasensor for thrombin based on the triplex-amplification of Hemin/G-quadruplex horseradish peroxidase-mimicking DNAzyme and horseradish peroxidase decorated FeTe nanorods

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S1. Optimization of the concentration of H₂O₂ for thrombin detection

As we know, both signal amplification and noise reduction are critical for sandwich-type electrochemical aptasensors. To obtain the best performance of the aptasensor, concentration of H₂O₂ was investigated as it may also influence the sensitivity of the aptasensor. After incubated with a fixed concentration of thrombin through sandwich assay, the aptasensor was tested in 0.1 M PBS (pH 7.0) containing H₂O₂ with different concentrations of 0.5, 1.0, 1.5, 2.0, 2.5 mM. It could be evidently observed that the reduction peak current of proposed aptasensor increased rapidly with increasing the concentration of H₂O₂ and the current tended to reach at a saturation value when the concentration of H₂O₂ was higher than 2.0 mM (Fig. S1). Therefore, 2.0 mM H₂O₂ was adopted in the subsequent work for signal amplification.

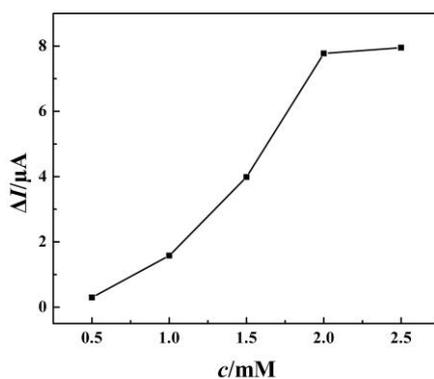


Fig. S1. Influence of the concentration of H_2O_2 on the current response of the aptasensor.

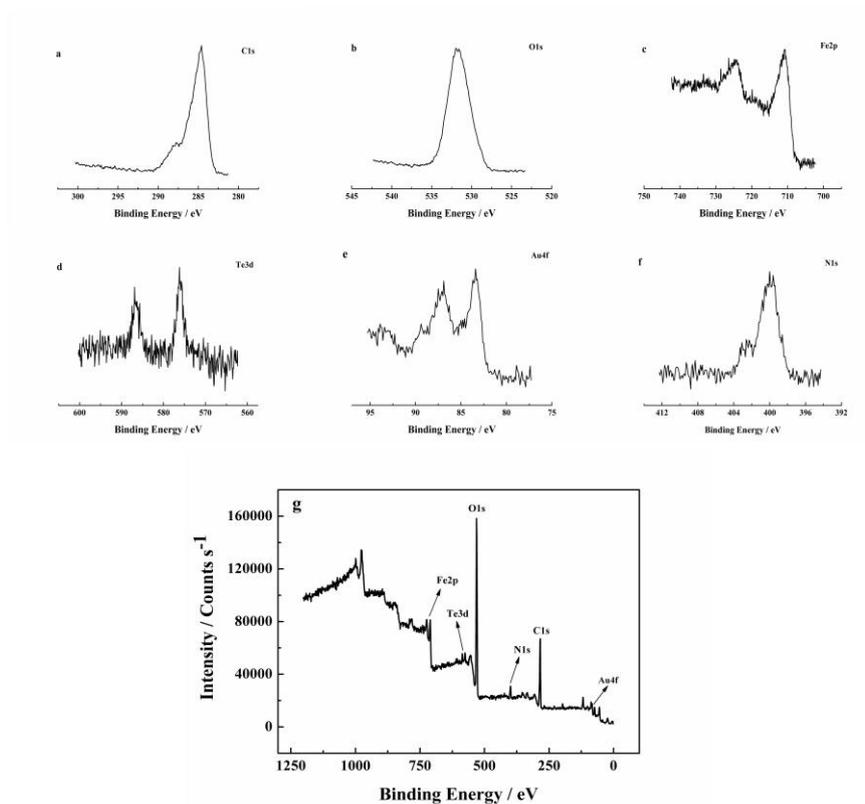


Fig. S2 XPS analysis of different nanomaterials. (a) and (b) were the C1s and O1s core level spectrum; (c), (d), (e), (f), represented the Fe2p, Te3d, Au4f, N1s core level spectrum of as-prepared Au@FeTe NRs nanocomposite respectively; (g) was the coverage of Au@FeTe NRs nanocomposites.

Table 1
Analytical application of the aptasensor

Serum samples	Added thrombin /M	Found Thrombin/M	Relative standard deviation ^a / %	Recovery / %
1	1.0×10^{-11}	1.1×10^{-11}	4.4	107.0
2	1.0×10^{-10}	1.1×10^{-10}	5.7	106.5
3	5.0×10^{-10}	4.8×10^{-10}	4.7	95.3
4	5.0×10^{-9}	4.6×10^{-9}	7.3	91.1

a. Determination of thrombin added in human blood serum ($n=3$) with the proposed aptasensor.

Table 2

Comparisons of proposed aptasensor with other sandwich-type electrochemical aptasensor for thrombin detection.

Analytical method	Detection limit	Linear range	Ref.
DPV	2 pM	0.01~50 nM	1
DPV	0.34 nM	0.001~50 nM	2
DPV	4.6 nM	0.007~70 nM	3
DPV	0.2 nM	0.8~15 nM	4
DPV	0.06 nM	0.1~5 nM	5
DPV	0.5 pM	0.001~20 nM	Our work

From the Table 2 we can see that our proposed aptasensor exhibits a much higher sensitivity and wider linear range, which provides a vigorous evidence of our strategy for highly sensitive detection of thrombin.

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Electronic supplementary information (ESI): Optimization of the concentration of H₂O₂ for thrombin detection, analytical application of the aptasensor and XPS analysis of Au@FeTe NRs nanocomposite. This material is available free of charge via the Internet at <http://pubs.acs.org>.