Au Nanoparticles Fluorescence Switch-mediated Target Recycling Amplification Strategy for Sensitive Nucleic Acid Detection

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Fig. S1. The change in fluorescence during the detection procedure. Condition: pH: 8.0, Temperature: 37 °C, F-DNA: 5.0×10^{-8} mol L⁻¹, T-DNA: 5.0×10^{-9} mol L⁻¹, Exo III: 1.3×10^{6} U L⁻¹.



Fig. S2. Effects of the concentration of AuNPs. Condition: pH: 8.0, Temperature: 37 °C, F-DNA: 5.0×10^{-8} mol L⁻¹, T-DNA: 5.0×10^{-9} mol L⁻¹, Exo III: 1.3×10^{6} U L⁻¹. 10 min incubation time of F-DNA and AuNPs, 30 min hybridization time of F-DNA on AuNPs surface and T-DNA, 60 min incubation time of Exo III.



Fig. S3. Effects of the pH. Condition: Temperature: 37 °C, F-DNA: 5.0×10^{-8} mol L⁻¹, T-DNA: 5.0×10^{-9} mol L⁻¹, Exo III: 1.3×10^{6} U L⁻¹. 10 min incubation time of F-DNA and AuNPs, 30 min hybridization time of F-DNA on AuNPs surface and T-DNA, 60 min incubation time of Exo III.



Fig. S4. Effects of the reaction temperature. Condition: pH: 8.0, F-DNA: 5.0×10^{-8} mol L⁻¹, T-DNA: 5.0×10^{-9} mol L⁻¹, Exo III: 1.3×10^{6} U L⁻¹. 10 min incubation time of F-DNA and AuNPs, 30 min hybridization time of F-DNA on AuNPs surface and T-DNA, 60 min incubation time of Exo III.



Fig. S5. Effects of concentrations of Exo III. Condition: pH: 8.0, Temperature: 37 °C, F-DNA: 5.0 \times 10⁻⁸ mol L⁻¹, AuNPs: 7.8 \times 10⁻⁹ mol L⁻¹, T-DNA: 5.0 \times 10⁻⁹ mol L⁻¹.10 min incubation time of F-DNA and AuNPs, 30 min hybridization time of F-DNA on AuNPs surface and T-DNA, 60 min incubation time of Exo III.



Fig. S6. Effects of incubation time of Exo III. Condition: pH: 8.0, Temperature: 37 °C, F-DNA: 5.0×10^{-8} mol L⁻¹, AuNPs: 7.8×10^{-9} mol L⁻¹, T-DNA: 5.0×10^{-9} mol L⁻¹, Exo III: 1.3×10^{6} U L⁻¹. 10 min incubation time of F-DNA and AuNPs, 30 min hybridization time of F-DNA on AuNPs surface and T-DNA.

Strategy	Nanomaterial	Synthesis procedure of the nanomaterial	Detection limit (mol·L ⁻¹)	Ref.
The fluorescent oligonucleotide with carbon nanotubes and target DNA	single walled carbon nanotubes	The purchased single walled carbon nanotubes were sonicatied in DMF for 5 h.	$4.0 imes 10^{-9}$	1
The carbon nanotube-based nanobeacon for detection of DNA	single walled carbon nanotubes	The purchased nanotubes were refluxed in HNO ₃ for 2 days, and the precipitates were oxidized by HNO ₃ /H ₂ SO ₄ for 4 h, and then the products were kept in an oven at 50 °C for 5 h.	4.2×10^{-11}	2
A graphene platform for sensing biomolecules	graphene oxide	The graphene oxide were synthesized from natural graphite powder by a modified Hummers method.	$1.0 imes 10^{-9}$	3
Graphene and HpaII-based fluorescent strategy for DNA detection	graphene oxide	The graphene oxide were synthesized by the Hummers method.	4.3 × 10 ⁻¹¹	4
A conjugation polymer nanobelts platform for nucleic acid detection	conjugation polymer nanobelts	The conjugation polymer nanobelts were synthesized with chemical oxidation polymerization of p-phenediamine monomers.	$3.0 imes 10^{-8}$	5
Fluorescence-enhanced nucleic acid detection with Pd nanowires	Pd nanowires	Te nanowires were synthesized with hydrothermal method, and Te nanowires mixed with H ₂ PdCl ₄ solution was stirred, and then centrifuged and washed for Pd nanowires.	3.0 × 10 ⁻¹⁰	6
The different interaction between ss- DNA or ds-DNA and MoS ₂ nanosheets	MoS ₂ nanosheets	MoS_2 nanosheets were synthesized by exfoliating bulk MoS_2 using electrochemical lithium-intercalation method.	5.0 × 10 ⁻¹⁰	7
The affinity change of carbon nitride nanosheets to DNA probes upon their recognition to targets	carbon nitride nanosheets	Carbon nitride nanosheets were prepared by sonicating bulk graphitic-phase carbon nitride, which was first synthesized by polymerization of melamine.	8.1 × 10 ⁻¹¹	8
Au nanoparticles fluorescence switch- mediated target recycling strategy	Au nanoparticles	The mixture of chloroauric acid and sodium citrate were maintained at 100 °C for 20 min with stirring.	1.0×10^{-11}	this method

Table S1 Comparison of the nanomaterial-based fluorescent methods for DNA detection

References

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