

A facile and sensitive peptide-modulating graphene oxide nanoribbon catalytic nanoplasmon analytical platform for human chorionic gonadotropin

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Abstract: The nanogold reaction between HAuCl_4 and citrate is very slow, and the catalyst graphene oxide nanoribbon (GONR) enhanced the nanoreaction greatly to produce gold nanoparticles (AuNPs) that exhibited strong surface plasmon resonance (SPR) absorption (Abs) at 550 nm and resonance Rayleigh scattering (RRS) at 550 nm. Upon addition of the peptide of human chorionic gonadotropin (hCG), the peptide could adsorb on the GONR surface, which inhibited the catalysis. When hCG was added, peptides were separated from the GONR surface due to the formation of stable peptide-hCG complex, which led to the activation of GONR catalytic effect. With the increase in hCG concentration, the RRS and Abs signal enhanced linearly. The enhanced RRS value showed a good linear relationship with hCG concentration in the range of 0.2–20 ng/mL, with a detection limit of 70 pg/mL. Accordingly, two new GONR catalytic RRS/Abs methods were established for detecting hCG in serum samples.

Keywords: nanocatalysis, graphene oxide nanoribbon, peptide regulation, hCG, RRS

Introduction

Nanoenzyme is a kind of mimetic enzyme that has unique properties and catalytic function. It has the characteristics of high efficiency, stability, economy and large-scale preparation and has been applied in different fields of materials science, physics, chemistry, biology, medicine, and environment.¹⁻³ The nanomaterials with nano analog enzyme activity have important significance in analytical chemistry. At present, the analytical application is mainly involved in the detection of heavy metal ions and biological molecules.³⁻⁶ Seok and Jongsoo³ detected mercury ions using ssDNA magnetic nanoparticles to inhibit the oxidation of H_2O_2 o-phenylenediamine to make no color change; the detection range was 5–75 $\mu\text{mol/L}$. Li et al⁴ used the principle that mercury ions inhibited the color reaction of oxidation of tetramethyl benzidine catalyzed by nano lead mimetic enzyme to analyze Hg^{2+} in water samples, with a detection limit (DL) of 7.2 nmoL/L. Peng et al⁵ detected mercury and copper using the principle of fluorescence quenching based on the specific binding of the fluorescent silver nanoclusters and the heavy metal ions; the detection ranges were 6–160 nmoL/L and 6–240 nmoL/L, respectively. Lien et al⁶ used fluorescence enhancement to detect thrombin due to amplex reagent (AR) for hydrogen peroxide oxidation catalyzed by thrombin transforming protein-mediated Bi-gold nanoparticles (AuNPs). Wang et al⁷ reduced the DL of H_2O_2 to 30 nmoL/L successfully by resonance Raman scattering with the oxidation of tetramethyl benzidine catalyzed by Au@Ag nanoparticles. Jiang et al⁸ achieved a low concentration using the glucose detection method, which was based

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on the catalytic effect of nanogold by spectrophotometry, with a detection range of 2–10 nmol/L. Wang et al⁹ used Cu²⁺-ion-modified graphene oxide (GO) nanoparticles as a heterogeneous catalyst mimicking functions of horseradish peroxidase (HRP), and a simple colorimetric method was proposed to detect 30–140 μ M ascorbic acid.¹⁰ Our research group has established some sensitive resonance Rayleigh scattering (RRS) and surface enhanced Raman scattering methods for rapid detection of melamine,¹¹ PDGF¹² and Cu,¹³ based on the nanoparticle probe aggregation and catalysis. Graphene oxide nanoribbons (GONRs) were prepared by using the method of facile unzipping of nitrogen-doped multiwalled carbon nanotubes (N-MWCNTs) with the help of microwave energy,^{14,15} which exhibited good water solubility and has been used in nanoanalysis. Zhang et al¹⁶ reported a GONR-modified electrochemical biosensor for the detection of 0.25–1.25 mmol/L amino acid. Dong et al¹⁷ developed a GONR biosensor for determination of 5–100 μ mol/L ATP. Zhu et al¹⁸ fabricated an MWCNT@GONR electrochemical sensor for 8–500 nmol/L polycyclic aromatic amines. In recent years, nanoplasmonic sensors have become widely used in medical, biotechnology and environmental science. One of the most promising directions has been surface-based nanoplasmonic sensors, and the potential of such technologies is still emerging.^{19–21} To date, there have been no reports about GONR catalytic gold nanoplasmon RRS/absorption (Abs) analytical platform for human chorionic gonadotropin (hCG), based on peptide reaction-modulating nanoenzyme activity, without preparation of nanoprobe and nanoparticle aggregation.

hCG is a glycoprotein secreted by the placenta trophoblast cells, and it is an important medical diagnostic marker of pregnancy. hCG also is one of the important markers of clinical disease. Its content is closely related to some diseases, such as gestational trophoblastic disease, germ cell tumors and Down syndrome.^{22–24} In addition, the quantitative detection of hCG is of great significance to the analysis of clinical medicine and the abuse of stimulants. At present, the detection methods of hCG are mainly electrochemical immunoassay, radioimmunoassay, chemiluminescent immunoassay, fluorescence resonance energy transfer method and optical immunoassay.^{25–30} Among these methods, radioimmunoassay is widely used, but there are radiation hazards. The electrochemical immunoassay method has a high sensitivity, but the operation is complex. The cost of fluorescence immunoassay is low, but there is a fluorescence quenching effect. Immunogold method is the most mature and widely used method for rapid detection of hCG, but nanogold aggregation caused instability. Most of these hCG immune

analysis methods are based on the antigen–antibody reaction; the biggest disadvantage is that the antibody biological protein is easily affected by the environment deterioration and then affects the detection effect. Recently, some new assays were developed for the detection of hCG.^{31–36} Zhao et al³¹ reported a microplate magnetic chemiluminescence enzyme immunoassay for the determination of hCG using the alkaline phosphatase (ALP) as a labeled reagent and 3-(2'-spiroadamantane)-4-methoxy-4-(3''-phosphoryloxy) phenyl-1,2-dioxetane (AMPPD) as a chemiluminescence reagent. Chang et al³² combined catalytic gold nanoparticles with an hCG-specific peptide aptamer to create a simple, sensitive, label-free colorimetric assay for hCG. Ding and Yang³³ reported an antibody-free and label-free mechanism for detecting hCG using a short oligopeptide as an hCG receptor to bind hCG, with a DL of 2 nmol/L hCG. Lei et al³⁴ reported a strategy of chemiluminescence resonance energy transfer for detection of 0.12 nmol/mL hCG using graphene as an efficient long-range energy acceptor and the magnetic nanoparticles for magnetic separation and immobilization of HRP-labeled anti-hCG antibody. Chen et al³⁵ reported an immunomagnetic assay for quantitative analysis of total β -subunit of hCG concentration in urine. The peptide of hCG was more stable than hCG antibody. Thus, Xia et al³⁶ found a new and stable method to detect hCG by using the polypeptide nano silver electrochemical sensor with voltammetry, with a DL of 0.4 mIU/mL. In this article, a new, simple, sensitive and low-cost nanoplasmon RRS/Abs analytical platform was developed, which was based on the polypeptide–hCG reaction regulation of GONR catalytic nanogold particle reaction.

Materials and methods

Materials

A DXR SmartRaman Spectrometer (Thermo Fisher Scientific, Waltham, MA, USA) with a laser wavelength of 633 nm and a power of 3.0 mW, Cary Eclipse Fluorescence Spectrophotometer (Varian Company, Salt Lake City, UT, USA), TU-1901 double beam ultraviolet (UV)–visible spectrophotometer (Beijing Purkinje General Instrument Co., Ltd., Beijing, China) and a thermostatic bath were used.

Reagents

A 0.50 mg hCG (Beijing Boosen Biotechnology Co., Ltd., Beijing, China) freeze-dried powder was mixed with 1.0 mL water and diluted to 10 mL; hCG polypeptide (HP; Biological Engineering Shanghai Limited by Share Ltd., Shanghai, China) was prepared by using Pro–Pro–Leu–Arg–Ile–Asn–Arg–His–Ile–Leu–Thr–Arg (PPLRINRHILTR),

1.0% $\text{HAuCl}_4 \cdot 4\text{H}_2\text{O}$ (Sinopharm Chemical Reagent Co., Ltd., Shanghai, China), 1.0% trisodium citrate (TCA; Guangdong Shantou Timong Chemical Factory, Guangzhou, China) and 0.1 mol/L HCl. All reagents were analytically pure, and water was double distilled.

GONR was prepared by the modified KMnO_4 procedure. Briefly, 50 mg of MWCNT powder was added into a 50 mL flask containing 10 mL concentrated H_2SO_4 solution for 1 hour. Then, 500 mg of KMnO_4 was added and heated at 60°C for 2 h. The product was poured into a 5 mL 30% H_2O_2 solution that was cooled by ice water. The mixture was dispersed by ultrasonic treatment for 10 min and centrifuged at 7,000 rpm for 10 min to obtain the supernatant containing 230 $\mu\text{g}/\text{mL}$ GONR that was calculated by adding MWCNT. Before use, it was neutralized with 50 mmol/L NaOH and then was diluted to the desired concentration.

Methods

A moderate amount of hCG and 100 μL of 1.4 $\mu\text{mol}/\text{L}$ HP were added to a 5 mL test tube for ~ 15 min and then 480 μL of 1 $\mu\text{g}/\text{mL}$ GONR was added and mixed well. After ~ 10 min, 75 μL of 3.4 mmol/L TCA, 150 μL of 0.01 mol/L HCl and 100 μL of 0.1% (84 $\mu\text{mol}/\text{L}$) HAuCl_4 were put into the test tube, diluted to 1.5 mL, mixed well, incubated at 60°C in a water bath for ~ 12 min and cooled with tap water. Finally, the mixture was diluted to 2 mL with water. The mixture was transferred into a quartz cell, and its RRS spectra were recorded. The RRS peak intensity at 550 nm (I) and the I_0 blank without hCG were recorded, and the value of $\Delta I = I - I_0$ was calculated.

Results and discussion

Analysis principle

Nanocatalytic reaction is an important way for signal amplification and enhancing sensitivity. The new nanocatalytic particle reaction of citrate- HAuCl_4 nanoparticles was investigated and used in nanoplasmon molecular spectral analysis. The AuNP reaction was very slow in the absence of nanocatalyst. Hence, in the non-catalytic reaction system, there was a low concentration of the produced AuNPs with weak surface enhanced Raman scattering signal due to small amount of VB4r molecular probes being adsorbed on the AuNPs. The GONRs with abundant surface hydroxyl groups had a strong catalytic effect on the AuNP reaction between citrate and HAuCl_4 . The AuCl_4^- and citrate could adsorb on the GONR catalyst surface, and redox reaction took place to form Au. The surface electrons of GONR enhanced the electron transfer in the redox reaction and speeded up the redox reaction. The produced AuNPs exhibited surface plasmon resonance (SPR)-RRS and Abs, and both increased with GONR concentration. HP had a high affinity with GONR and a high specificity for targeting hCG. It could easily adsorb to the GONR surface through electrostatic attraction that weakened GONR catalysis. When the hCG was present in the solution, the HP selectively recognized and tightly bound to hCG to form a stable HP-hCG complex that escaped the nanosurface to recover the nanocatalysis. With the increase in hCG concentration, the catalysis enhanced due to the increase in the free GONR concentration. Thus, we could establish a new nanogold plasmon RRS/Abs platform for the determination of hCG (Figure 1) with high sensitivity, high selectivity and simplicity.

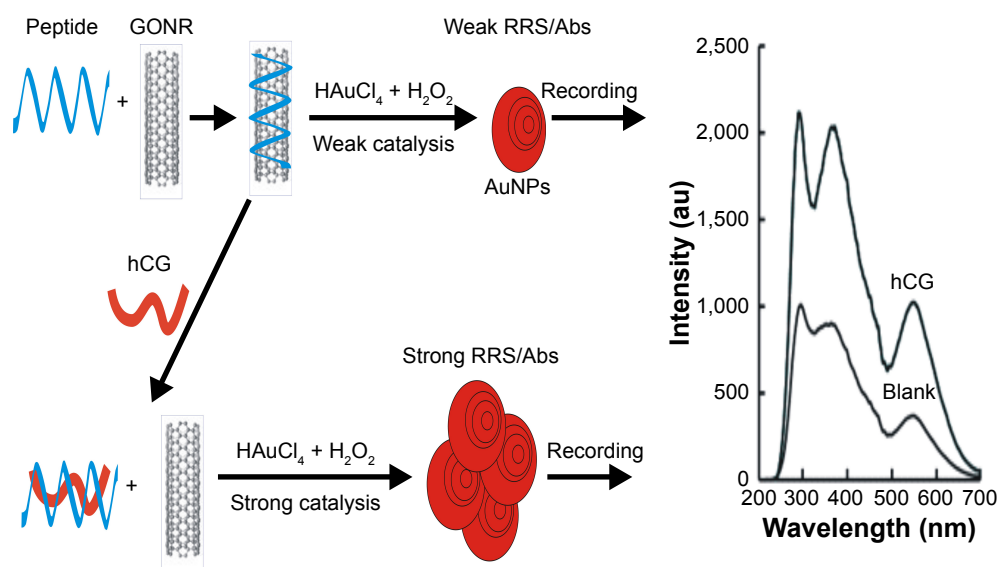


Figure 1 Scheme of peptide-controlling GONR catalytic reaction-RRS detection of hCG.

Abbreviations: Abs, absorption; AuNP, gold nanoparticle; au, atomic unit; GONR, graphene oxide nanoribbon; hCG, human chorionic gonadotropin; RRS, resonance Rayleigh scattering.

RRS spectra

RRS is a simple and sensitive molecular spectral technique. It is suitable for nanoparticle reaction and is selected for the study of the nanocatalytic system. To date, there have been no reports about RRS study of the GONR catalytic nanogold reaction and its coupling with immunoreactions without nanoparticle aggregation that could cause instability of the analytical system. Thus, RRS was selected for the study of the analytical system. The catalysis of GONR and GO in the redox reaction was considered; as the GONR is stronger

than the GO, GONR was selected as the nanoenzyme. For the HP-hCG-GONR-HAuCl₄-TCA analytical system, the generated AuNPs exhibited three RRS peaks at 290, 370 and 550. The RRS peak at 550 nm had a good linear relationship between the signal and hCG concentration and was selected for assay of hCG (Figure 2A). The nanocatalytic system of GONR-HAuCl₄-TCA exhibited three RRS peaks at 290 and 545 nm (Figure 2B), and the RRS peak intensity increased with GONR concentration. Upon addition of HP, HP could inhibit the catalytic activity of GONR nanoenzyme; the

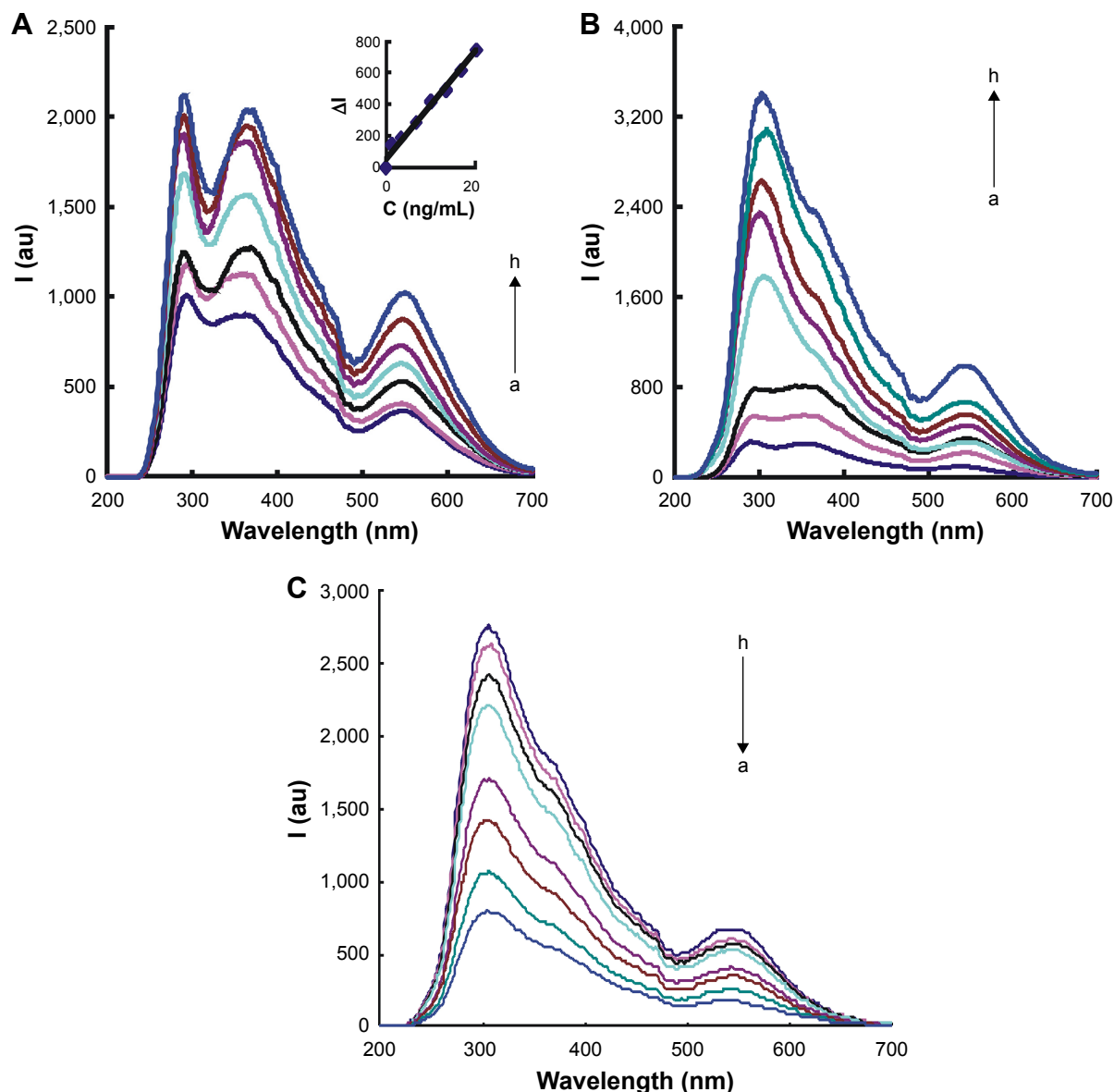


Figure 2 RRS spectra of the GONR systems.

Notes: (A) HP-hCG-GONR-HAuCl₄-TCA system – a: 35 nmol/L HP +240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄; b: a+0.33 ng/mL hCG; c: a+1 ng/mL hCG; d: a+2 ng/mL hCG; e: a+4 ng/mL hCG; f: a+6 ng/mL hCG; g: a+10 ng/mL hCG; and h: a+13.33 ng/mL hCG. The inset graph was the working curve. (B) GONR-HAuCl₄-TCA system – a: 0.167 mmol/L HCl +0.34 mmol/L TCA+ 5.6 μmol/L HAuCl₄; b: a+16 ng/mL GONR; c: a+40 ng/mL GONR; d: a+80 ng/mL GONR; e: a+120 ng/mL GONR; f: a+160 ng/mL GONR; g: a+240 ng/mL GONR; and h: a+320 ng/mL GONR. (C) HP-GONR-HAuCl₄-TCA system – a: 240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄; b: a+1.4 nmol/L HP; c: a+3.5 nmol/L HP; d: a+7 nmol/L HP; e: a+10.5 nmol/L HP; f: a+14 nmol/L HP; g: a+21 nmol/L HP; and h: a+28 nmol/L HP.

Abbreviations: C, concentration; I, intensity; RRS, resonance Rayleigh scattering; GONR, graphene oxide nanoribbon; HP, hCG polypeptide; hCG, human chorionic gonadotropin; TCA, trisodium citrate.

catalytic activity decreased and the two RRS peaks at 300 and 550 nm decreased with the increasing HP concentration (Figure 2C).

Ultraviolet Abs spectra

A spectrophotometer is a very simple and low-cost spectral instrument that was selected for studying the SPR Abs of nanogold nanoparticles. For the HP-hCG-GONR-HAuCl₄-TCA system, the SPR Abs peak at 540 nm increased with hCG concentration and was selected for assay of hCG (Figure 3A). The GONR catalyzed the TCA reduced HAuCl₄ to form AuNPs with a SPR Abs peak at 550 nm (Figure 3B)

that corresponds to the RRS peak at 545 nm, in which the peak increased with the GONR increasing. HP can adsorb on the GONR surface and inhibit the catalysis; the SPR Abs peak decreased with the increasing HP concentration (Figure 3C).

Catalysis and inhibition

In the experimental conditions, the reduction of HAuCl₄-TCA could be catalyzed by GONR or GO to form AuNPs. The RRS and Abs values increased with the increasing catalyst concentration. The catalytic action of GONR was stronger than that of the GO, because the slope of GONR is

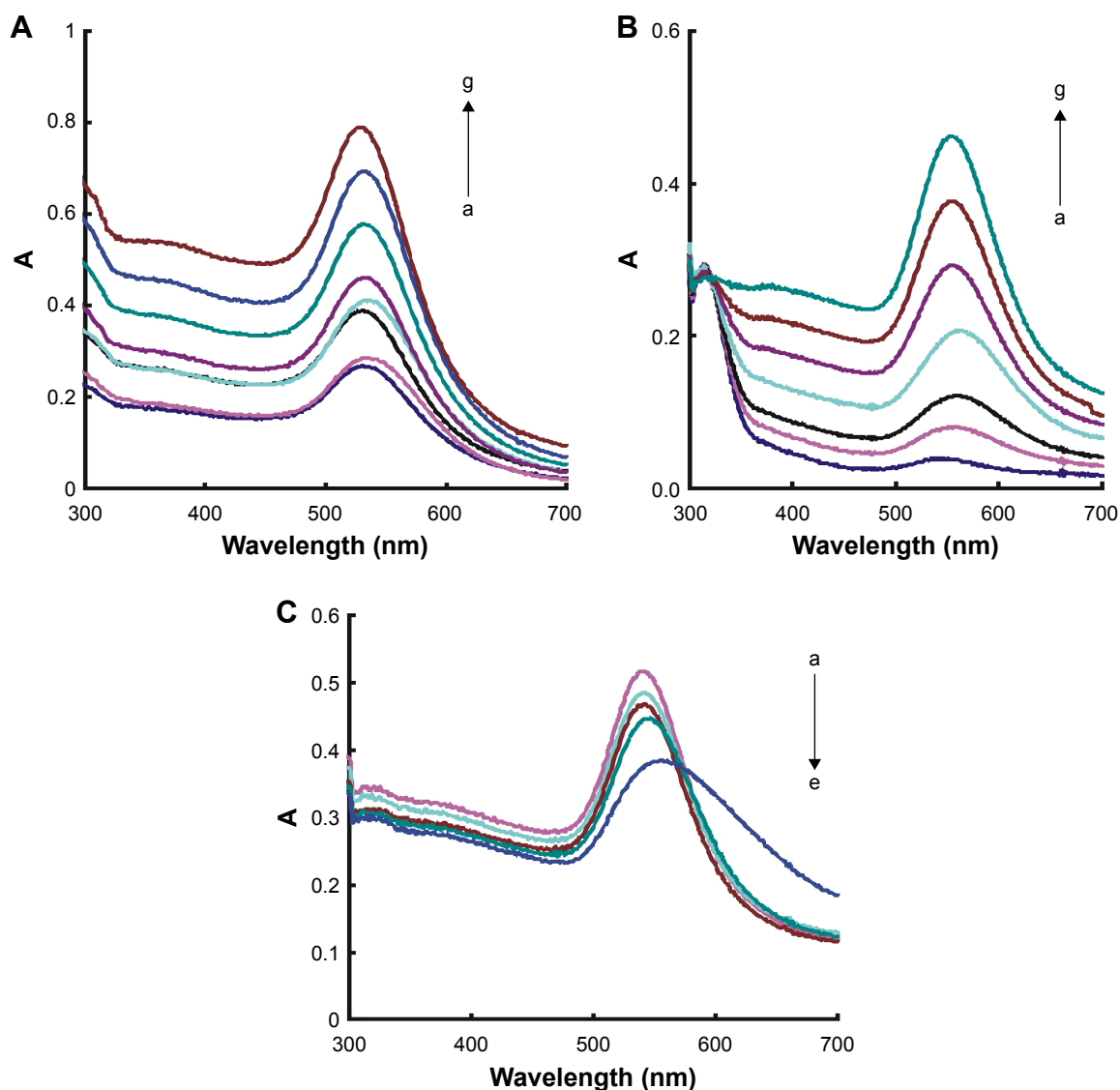


Figure 3 UV spectra of the GONR systems.

Notes: (A) HP-hCG-GONR-HAuCl₄-TCA system – a: 35 nmol/L HP +240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄; b: a+0.67 ng/mL hCG; c: a+1.67 ng/mL hCG; d: a+3.33 ng/mL hCG; e: a+6.67 ng/mL hCG; f: a+10 ng/mL hCG; and g: a+13.33 ng/mL hCG. (B) GONR-HAuCl₄-TCA system – a: 0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄; b: a+16 ng/mL GONR; c: a+40 ng/mL GONR; d: a+80 ng/mL GONR; e: a+120 ng/mL GONR; f: a+160 ng/mL GONR; and g: a+240 ng/mL GONR. (C) HP-GONR-HAuCl₄-TCA system – a: 240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄; b: a+5 nmol/L HP; c: a+10 nmol/L HP; d: a+20 nmol/L HP; and e: a+30 nmol/L HP.

Abbreviations: A, absorbance; UV, ultraviolet; GONR, graphene oxide nanoribbon; HP, hCG polypeptide; hCG, human chorionic gonadotropin; TCA, trisodium citrate.

Table 1 Comparison of the nanocatalysis and the inhibition of HP on the reaction of TCA–HAuCl₄

System	Linear range	Linear equation	Correlation coefficient
GONR	1.0–30 ng/mL GONR	$\Delta I_{545\text{ nm}} = 61.5C + 203$	0.9918
AuNP	1.0–25 ng/mL AuNP	$\Delta I_{545\text{ nm}} = 58.3C + 150$	0.9834
(d = 10 nm)			
HP–GONR	2.0–30 nmoL/L HP	$\Delta I_{545\text{ nm}} = 47.8C + 69.5$	0.9804
GO	2.0–30 ng/mL GO	$\Delta I_{545\text{ nm}} = 36.9C - 89.8$	0.9621

Note: I is measured in au.

Abbreviations: HP, hCG polypeptide; TCA, trisodium citrate; GONR, graphene oxide nanoribbon; AuNP, gold nanoparticle; GO, graphene oxide; hCG, human chorionic gonadotropin; I, intensity; C, concentration; d, diameter.

larger than the GO due to the size of GONR being smaller than the GO. When HP was added into the system, the RRS peak and Abs values decreased with increasing HP concentration (Table 1). The reason was that HP could be attached to the surface of the nanocatalyst by physical electrostatic adsorption, so as to block the exchange between the redox electrons and catalyst surface electrons.

Transmission electron microscopy

Transmission electron microscopy (TEM) is used to observe the particulate particle size and surface morphology. The blank reaction solution of HP–hCG–GONR–HAuCl₄ system was obtained by the procedure, and the TEM (Figure 4A) was recorded with particle average size of 80 nm. When hCG was added into the solution, more AuNPs formed (Figure 4B and C); the particle size was 60 and 50 nm.

Optimization of analysis conditions

The reaction conditions of HP–hCG–GONR–TCA–HAuCl₄ were optimized according to the experimental method. The effect of HP concentration on ΔI was investigated when it reached 35 nmol/L. The value of ΔI was the largest (Figure 5A), so the selection of the concentration of HP was 35 nmol/L. The effects of GONR concentration on the catalytic system at ΔI were investigated; according to the result, 240 ng/mL GONR was selected for use (Figure 5B). The concentration of TCA was researched; when the TCA

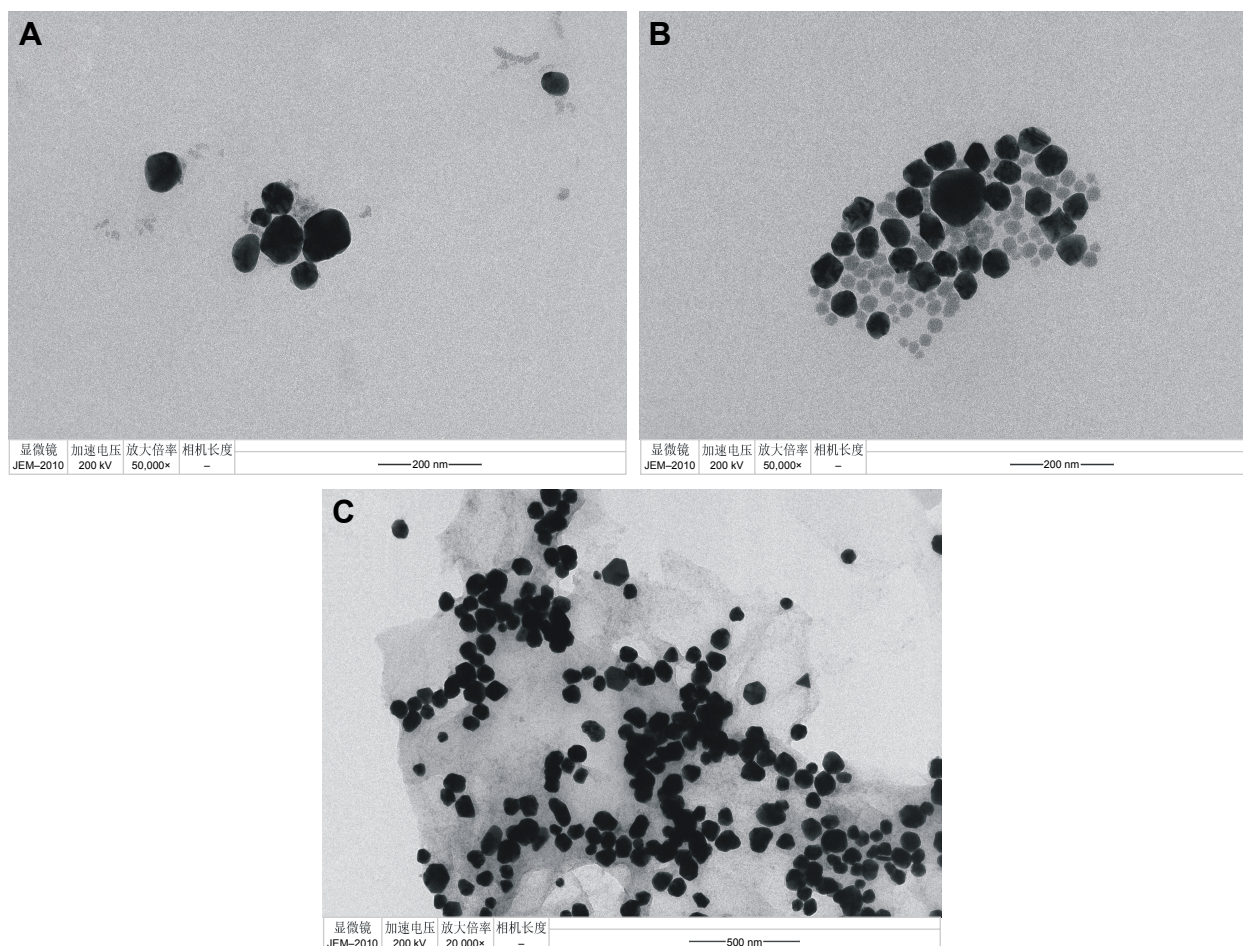


Figure 4 TEMs of analysis system.

Notes: (A) 35 nmol/L HP + 240 ng/mL GONR + 0.167 mmol/L HCl + 0.34 mmol/L TCA + 5.6 μ mol/L HAuCl₄; (B) A + 4 ng/mL hCG. (C) A + 10 ng/mL hCG.

Abbreviations: TEM, transmission electron microscopy; HP, hCG polypeptide; GONR, graphene oxide nanoribbon; TCA, trisodium citrate; hCG, human chorionic gonadotropin.

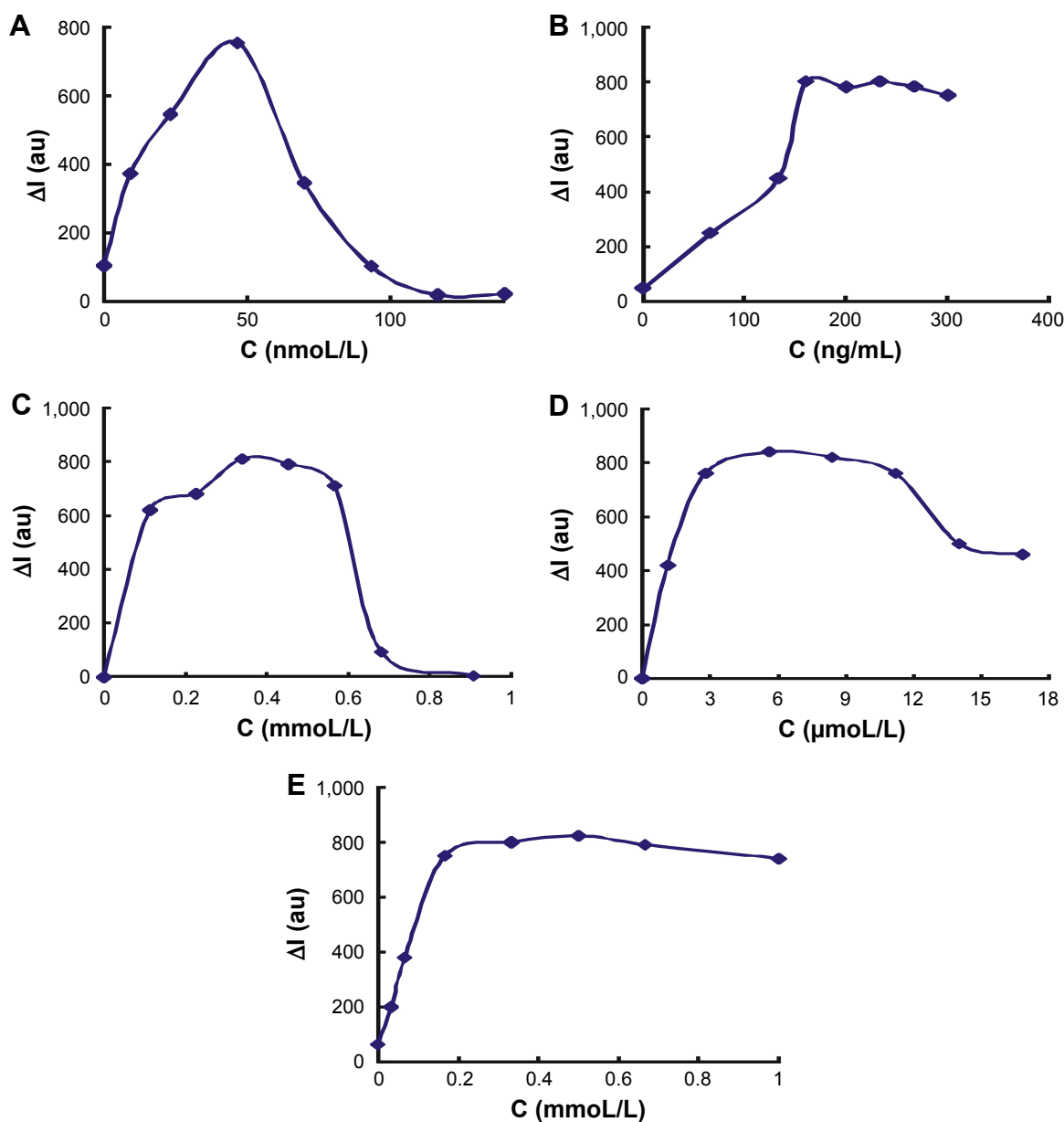


Figure 5 Effect of reagent concentration.

Notes: (A) HP -10 ng/mL hCG +240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μ mol/L HAuCl₄; (B) GONR -35 nmol/L HP +10 ng/mL hCG +GO +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μ mol/L HAuCl₄; (C) TCA -35 nmol/L HP +10 ng/mL hCG +240 ng/mL GONR +0.167 mmol/L HCl +5.6 μ mol/L HAuCl₄; (D) HAuCl₄ -35 nmol/L HP +10 ng/mL hCG +240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA+HAuCl₄; (E) HCl -35 nmol/L HP +10 ng/mL hCG +240 ng/mL GONR+HCl +0.34 mmol/L TCA +5.6 μ mol/L HAuCl₄.

Abbreviations: HP, hCG polypeptide; hCG, human chorionic gonadotropin; GONR, graphene oxide nanoribbon; TCA, trisodium citrate; GO, graphene oxide; I, intensity; C, concentration.

concentration was 0.34 mmol/L, the value of ΔI was the largest (Figure 5C); the selection of the concentration of TCA was 0.34 mmol/L; when the HAuCl₄ concentration was 5.6 μ mol/L, the ΔI value was the largest (Figure 5D); 5.6 μ mol/L HAuCl₄ was selected. The dosage of HCl was optimized, and when the concentration of HCl was 0.167 mmol/L, ΔI reached the maximum value (Figure 5E); 0.167 mmol/L HCl was chosen; The optimization of the aforementioned reaction was carried out at 60°C water bath condition; the reaction time was 10 min. The reaction

temperature and time were examined. A temperature of 60°C for 12 min was chosen for use (Figure 6).

Working curve

The working curves for hCG were obtained according to the experimental method. For the GONR analytical system, the enhanced values of the RRS peak at 550 nm and the Abs peak at 550 nm were linear with hCG concentration in the range of 0.2–20 ng/mL and 10–15 ng/mL, respectively, with a linear equation of $\Delta I = 91.5C + 34.0$

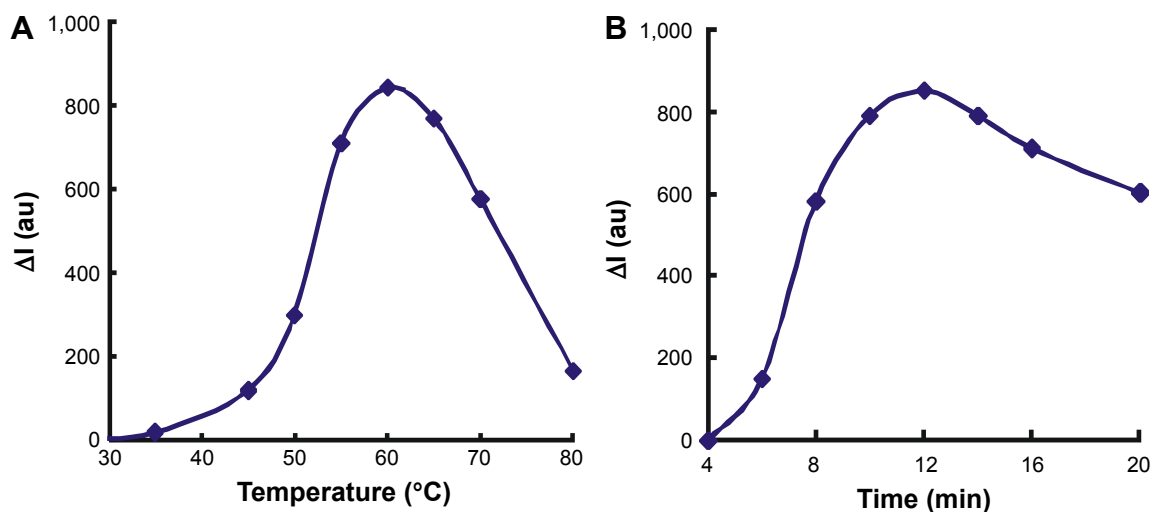


Figure 6 The effect of reaction temperature and time.

Notes: (A) Reaction temperature: 35 nmol/L HP +10 ng/mL hCG +240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄. (B) time: 35 nmol/L HP +10 ng/mL hCG +240 ng/mL GONR +0.167 mmol/L HCl +0.34 mmol/L TCA +5.6 μmol/L HAuCl₄.

Abbreviations: HP, hCG polypeptide; hCG, human chorionic gonadotropin; GONR, graphene oxide nanoribbon; TCA, trisodium citrate; I, intensity.

and $\Delta A_{550\text{ nm}} = 0.0334C + 0.0223$, correlation coefficient of 0.9863 and 0.9661, and DL of 0.07 and 3 ng/mL, respectively. We can see that the RRS analytical system is more sensitive than the Abs and was thus chosen for sample detection. Compared to the reported hCG methods,^{25–36} this RRS method is not only sensitive and selective but also facile, of low-cost and without nanoparticle aggregation. The clinical method of chemiluminescent immunoassay had a DL of 0.143 ng/mL, limit of quantitation (LQ) of 5 ng/mL and linear range of 5–540 ng/mL hCG;³⁷ the DL and LQ of this RRS method are better than the clinical method, and the linear range of the two methods are close to each other, as the upper LQ divided by the lower LQ is 108 and 100, respectively.

Interference

The effect of the coexisting substances on the determination of 10 ng/mL hCG was investigated. The tested common interfering ions and amino acids, IgG and IgM, did not interfere with the determination when the relative error was within 10% (Table 2). This indicated that this method had a good selectivity.

Analysis of samples

Seven serum samples of women were provided by the No 5 People's Hospital of Guilin, Guangxi, China. A 100 μL sample was diluted to 10 mL with water before determination, the next detections were according to the procedure, and the content was equal to detection value multiplying dilution times of 100. In addition, recovery tests were performed, and the recovery was equal to ((Found value–Detection

value]/Added value) multiplied by 100%. The results are listed in Table 3, and the recoveries were in the range of 90.1%–106%; the relative standard deviation (RSD) was in the range of 0.90%–3.97% according to five detection values. The obtained results were not obviously different from clinical diagnosis values from the No 5 People's Hospital of Guilin,³⁷ so the results demonstrate that the method was reliable and practical in clinical detection.

Conclusion

In this work, GONR exhibited strong catalysis of the AuNP nanoreaction between HAuCl₄ and citrate that produced an SPR Abs peak at 550 nm and an RRS peak at 550 nm. Upon addition of HP, HP adsorbed on the surface of GONR nanoparticles, which blocked the electron transfer in the redox reaction and inhibited GONR catalytic action. When the target hCG was added, it reacted with HP to form stable

Table 2 Effect of coexisting substances

Coexisting substance	Relative multiple	Relative error (%)	Coexisting substance	Relative multiple	Relative error (%)
K ⁺	100	–1.0	SO ₃ ^{2–}	100	–5.7
Ca ²⁺	100	–7.4	NO ₂ [–]	100	8.0
Mg ²⁺	100	–6.3	S ₂ O ₃ ^{2–}	100	–6.0
Fe ³⁺	100	–5.3	CO ₃ ^{2–}	100	–6
IgM	10	–0.8	Glycine	100	–4
IgG	10	–3.5	Lysine	100	–7
Tryptophan	10	5.0	Aspartic acid	100	8.0
Glutamate	50	1.2	Valine	100	–4.0
Phenylalanine	50	–1.0	Tyrosine	100	–6.0

Table 3 Analysis of hCG samples

Sample	Detection value (ng/mL)	Content (ng/mL)	Added value (ng/mL)	Found value (ng/mL)	Recovery (%)	RSD (%)	Ref value (ng/mL)
1	0.548	55.6	0.25	0.7842	94.5	1.05	56.6
2	0.432	43.7	0.5	0.9258	98.8	3.97	45.7
3	0.404	40.9	1.0	1.364	90.1	1.43	39.6
4	0.480	48.6	0.5	1.010	106	0.90	46.9
5 ^a	0.310	3.10	0.25	0.550	96.0	3.76	32.1
6	5.24	524	5.0	10.52	105.6	2.85	518
7	16.7	1,670	1.0	17.63	93.0	2.45	1,692

Note: ^aThe sample dilution time was 10 min.

Abbreviations: hCG, human chorionic gonadotropin; Ref, reference; RSD, relative standard deviation.

HP-hCG complex and free GONR nanocatalyst that caused the SPR Abs value and the RRS intensity to increase due to the recovery of nanocatalysis. According to this principle, RRS and SPR Abs methods for detection of hCG were established based on the control of the catalytic reaction of GONR nano enzyme by the peptide reaction. With the increase in the amount of the analyte hCG, the RRS signal of the system increased gradually. The increased RRS intensity, ΔI , at 550 nm showed a good linear relationship with the concentration of hCG in the range of 0.2–20 ng/mL, with a DL of 70 pg/mL. The RRS method was used in analyzing serum samples, with a recovery of 90.1%–106% and RSD of 0.90%–3.97%.

Acknowledgments

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Disclosure

The authors report no conflicts of interest in this work.

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