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A frogspawn-like Ag@C core–shell structure for an ultrasensitive label-free electrochemical immunosensing of carcinoembryonic antigen in blood plasma

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A three-dimensional (3D) frogspawn-like structure was achieved by simply coating nano-carbon outside silver nanospheres (Ag@C NFS) and used as a probe to capture the anti-carcinoembryonic antigen for the electrochemical immunosensing of carcinoembryonic antigen (CEA), a typical biomarker of several diseases such as gastric cancer, intestinal cancer and colon cancer. Moreover, Ag@C nanocables (Ag@C NCs) were also synthesized. By comparison, the globular 3D frogspawn-like structure endowed Ag@C NFS with a larger surface area, which is preferred to improve the capability of loading antibodies, higher water solubility, better biocompatibility and improved electrical conductivity, which was likely attributed to the synergistic effects of Ag and crystalline graphite carbon and the different structure with more hydroxyl groups exposed. Therefore, the resultant Ag@C NFS was used as an electrochemical immunosensing platform to fabricate a label-free immunosensor for the analysis of CEA, which showed an excellent immunosensing performance with a wide linear CEA detection range from 0.0001 ng mL⁻¹ to 100 ng mL⁻¹ and a low detection limit of 5.12 pg mL⁻¹. In particular, the good reproducibility, high stability and specificity of the proposed immunosensor ensured the successful application in the quantitative determination of CEA in cancerous human serum samples, providing a promising alternative to detect other biomarkers.

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1. Introduction

In recent years, great strides have been made to reduce death rates from cancer in virtue of various diagnostic technologies, such as magnetic resonance (MR) imaging,¹ X-ray computed tomography (CT) imaging² and photoacoustic (PA) imaging.³ However, these techniques can only detect established tumors. In some cases, it's too late. Therefore, it's urgent to find one method that could detect tumor markers generated by the body or by the tumor tissue itself, rapidly and quantitatively for cancer preliminary screening and risk evaluation.^{4,5} As one of the most reliable and preferred tumor markers, CEA has most typically been used to predict the potential for patients having mammary gland, lung, or colorectal cancer, with a CEA level above 5 ng mL⁻¹ for tumor existence in the abovementioned organs.^{6,7} So far, although several methods have been used for detecting CEA, such as electrochemiluminescence

immunoassay,⁸ enzyme-linked immunosorbent assay (ELISA),⁹ photoelectrochemical immunoassay¹⁰ and radioimmunoassay,¹¹ the fairly high required time or costs limit their further application to some extent.¹² As a preferred alternative, electrochemical immunosensors have been widely used for testing CEA and some other tumor markers due to their high efficiency, low cost and miniaturized operating equipment.^{13–16} As an important type of electrochemical immunosensor, label-free electrochemical immunosensors have a big advantage over labeled (sandwich-type) electrochemical immunosensors because of their simpler architecture.^{17,18} Compared with labeled immunosensors, label-free electrochemical immunosensors don't require a complicated labeling process. In other words, label-free electrochemical immunosensors could directly monitor the binding process of antibody–antigen interaction by amperometric measurements or electrochemical impedance spectroscopy without the involvement of secondary antibodies.^{14,18,19} Therefore, more and more attention has been paid to the application of label-free electrochemical immunosensors in the field of tumor marker detection.^{20–22} In addition, the detection principle of the electrochemical immunosensor is specific binding of antibodies (Ab) and antigens (Ag), ensuring a fairly high specificity.²³

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To make the designed electrochemical immunosensor with a satisfactory detection limit and high sensitivity, it is critical to explore optimal electrode materials that can not only accelerate charge transfer on the surface of the electrode but also can provide a high specific surface area to anchor more antibodies. To date, noble metal nanomaterials, such as gold nanorods,²⁴ palladium nanocubes⁵ and silver nanoparticles,²⁵ were commonly used for electrode materials because of their favourable electrical conductivity and biocompatibility, among which silver nanomaterials were the most commonly used, owing to their relatively low price for raw materials and comparable conductivity. Unfortunately, silver nanomaterials are apt to agglomerate and very easily denature when exposed to moist air.²⁶ To address this dilemma, functional shells were usually constructed to protect the metal cores from agglomeration, as well as to improve their sensitivity for the detection of substances.^{27,28} As a promising candidate, carbon coating is of great research significance. On the one hand, the as-formed carbon coating could enhance the electrical conductivity through the cooperative effects with inner metal cores. On the other hand, the carbon shell is to a great extent the security umbrella of its inner silver NPs.²⁸ Moreover, the hydrophilic hydroxyl groups on the carbon shell could significantly promote the composite's water solubility and biocompatibility.²⁹

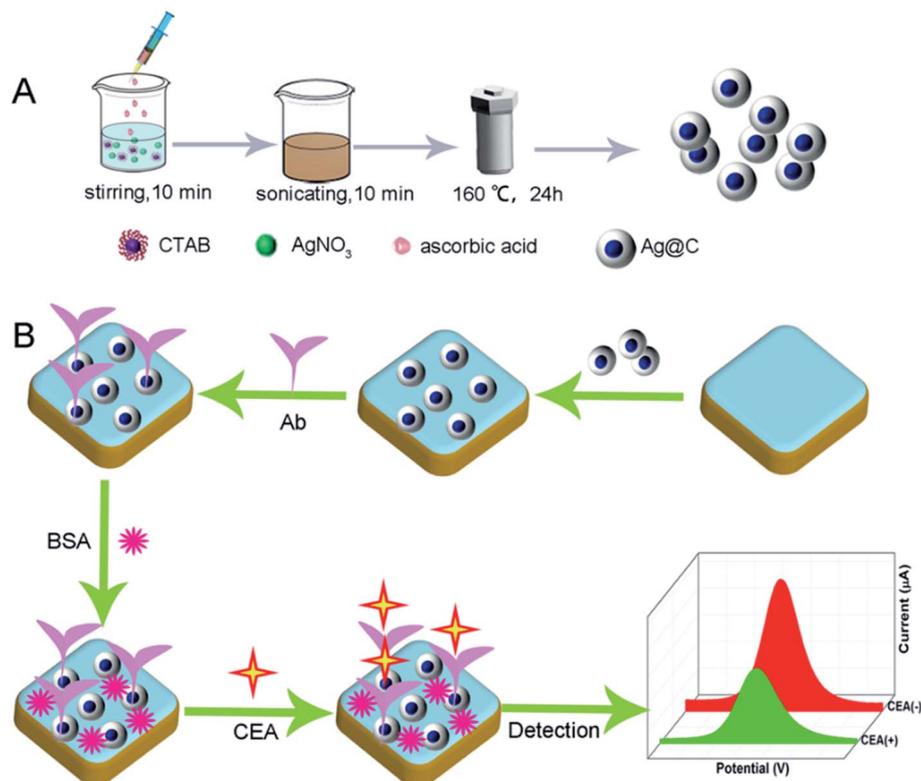
Herein, a three dimensional (3D) frogspawn-like structure core-shell nanocarbon coated silver nanosphere (Ag@C NFS)

was directly constructed by a one-pot solvothermal method with the assistance of hexadecyl trimethyl ammonium bromide (CTAB) and utilized to capture anti-CEA (Ab) for the construction of a label-free electrochemical immunosensor for CEA detection, as shown in Scheme 1. For comparison, nanocarbon coated silver nanocables (Ag@C NCs) were synthesized and used to fabricate a label-free electrochemical immunosensor for CEA detection as well. It was found that the Ag@C NFSs demonstrated obviously better electrochemical properties and higher Ab loading capacity than Ag@C NCs, which was likely ascribed to the larger surface area, better biocompatibility, faster electron transfer as a result of different structures between Ag@C NFSs and Ag@C NCs and the synergistic effect of silver and nanocarbon. As expected, the obtained label-free immunosensor performed well in the detection of CEA in terms of sensitivity, selectivity, reproducibility and stability, indicating great potential for the quantitative detection of other tumor markers.

2. Materials and methods

2.1. Reagents and materials

Ascorbic acid (AA), CTAB, glycerol, sulfuric acid (H_2SO_4), sodium phosphate monobasic dihydrate ($\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$), phosphate buffered solution (PBS) and sodium phosphate dibasic dodecahydrate ($\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$) were purchased from Aladdin Reagents Company (Shanghai, China).



Scheme 1 Schematic illustration of (A) the preparation of Ag@C NFSs and (B) the fabrication of the Ag@C NFS-based electrochemical immunosensor for CEA detection.



Polyvinylpyrrolidone K30 (PVP K30), silver nitrate (AgNO_3), gamma alumina powder, potassium ferrocyanide trihydrate ($\text{K}_4\text{Fe}(\text{CN})_6 \cdot 3\text{H}_2\text{O}$), potassium ferrocyanide ($\text{K}_3\text{Fe}(\text{CN})_6$), and potassium chloride (KCl) were gained from Sinopharm Chemical Reagent Company (Shanghai, China). CEA antibody, CEA, PSA, human IgG, AFP and bovine serum albumin (BSA) were provided by Shanghai Linc-Bio Science Co. Ltd. All the reagents used in the experiments were of analytical grade and used as received. Double distilled water was provided by automatic double pure water distillation equipment (Shanghai Yarong Biochemical Instrument Factory) for the preparation of all aqueous solutions.

2.2. Apparatus

All electrochemical performance parameters are tested on an Autolab PGSTAT 100 potentiostat/galvanostat (Shanghai Nano Industrial Co., Ltd, China). A three electrode system with a platinum wire electrode as the counter electrode, a saturated calomel electrode ($\text{Hg}/\text{Hg}_2\text{Cl}_2$) as the reference electrode and a glassy carbon electrode (GCE) as the working electrode was used. The water contact angle was measured using the contact angle system OCA (Beijing Eastern-Dataphy Instruments Co., Ltd, China). Scanning electron microscopy (SEM) images and energy dispersive spectroscopy (EDS) spectra were obtained using a field emission SEM (Zeiss, Germany). Transmission electron microscopy (TEM) images were recorded using a JEOL JEM 2100F transmission electron microscope (TEM). X-ray powder diffraction (XRD) was tested using a Rigaku Ultima IV

X-ray diffractometer from 5° to 80° . Fourier transform infrared (FT-IR) spectra were measured on a Nicolet 380 FT-IR spectrophotometer. Ultraviolet-visible absorption spectra were recorded on a UV-1900 spectrophotometer (Shimadzu, Japan). The hydrodynamic diameter was measured using a NanoBrook 90Plus Particle Size Analyzer.

2.3. Preparation of frogspawn-like Ag@C NFSs

The Ag@C NFSs were synthesized by a one-pot hydrothermal method.²⁴ Briefly, 0.0849 g AgNO_3 and 0.0546 g CTAB were completely dissolved in 30 mL double-distilled water with the assistance of magnetic stirring and ultrasonication. Then, the solution mixture was mixed with 10 mL of $0.03522 \text{ g mL}^{-1}$ AA solution under sonication to obtain a homogeneous solution. Afterward, the final solution mixture was transferred to a 50 mL Teflon-lined autoclave and then maintained at 160°C for 24 h after being well sealed. When the autoclave cooled to room temperature naturally, the resulting brown products were collected through centrifugation, washed with distilled water three times, and finally placed in a freeze-dryer for 6 h to remove all residual moisture.

2.4. Preparation of Ag@C NCs

The Ag@C NCs were prepared with the mediation of H_2SO_4 according to the previously reported procedure.³⁰ Typically, 0.0015 mol of glycerol and 0.015 mol of AgNO_3 were dissolved in 30 mL of 2 mol L^{-1} H_2SO_4 solution in sequence. After being stirred evenly, the mixed solution was redirected to a 50 mL

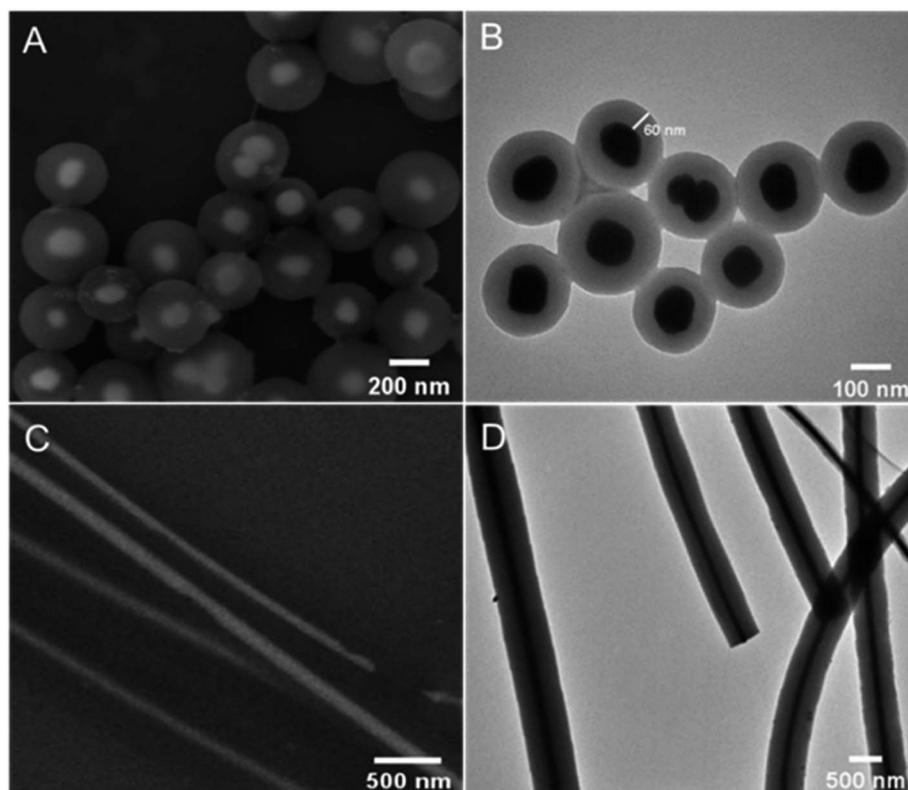


Fig. 1 SEM images of (A) Ag@C NFSs and (C) Ag@C NCs and TEM images of (B) Ag@C NFSs and (D) Ag@C NCs.



autoclave and reacted at 180 °C for 12 h in an oven to produce the preferred Ag@C NCs.

2.5. Construction of the Ag@C NFSs-based label-free electrochemical immunosensor

The whole construction process of the electrochemical immunosensor using Ag@C NFSs as an Ab capturer is clearly demonstrated in Scheme 1. First, a bare GCE was polished with alumina slurry with different diameters, and then ultrasonically rinsed with deionized water to achieve a mirrored surface. Then, 6 μL of 3.5 mg mL^{-1} Ag@C NFS containing aqueous solution was dropped on the pregrinded GCE and spontaneously dried in an incubator at 37 °C, followed by incubation with 5 μL of 5 $\mu\text{g mL}^{-1}$ Ab. Afterwards, Ab was immobilized onto the Ag@C NFS modified GCE by the Ag–N bond. Subsequently, 3 μL of 1% BSA solution was pipetted onto the Ab captured electrode so as to seal the potential nonspecific active sites. Finally, the obtained electrode was covered with 6 μL CEA at different concentrations and then stored in a refrigerator at 4 °C for the following detection.

3. Results and discussion

3.1. Characterization of the synthesized Ag@C NFSs and Ag@C NCs

The morphology and structure of the obtained Ag@C NFSs and Ag@C NCs were characterized by both SEM and TEM (Fig. 1). As shown in Fig. 1A and B, the obtained Ag@C nanomaterials were of frogspawn-like structure in which the silver nanosphere

acted as a core and nanocarbon with a nearly uniform thickness of about 60 nm as the shell. From Fig. 1C and D, it could be seen that the silver NCs were well wrapped with the carbon nano-coating with a similar diameter, quite smooth surface and straight morphologic structure. The elemental mapping images clearly show the elemental distribution of Ag and C in the structures of both Ag@C NFSs (Fig. 2A–C) and Ag@C NCs (Fig. 2D–F), verifying their successful synthesis.

In order to determine the actual size of the obtained Ag@C NFSs in the aqueous solution, dynamic light scattering (DLS) was selected to measure the hydrodynamic diameter of Ag@C NFSs. As shown in Fig. 3A, the hydrodynamic diameter was ~ 225 nm, in good accordance with that recorded by SEM and TEM. In addition, XRD was used to know more about the crystalline structure of Ag@C NFSs. The main Bragg peaks shown in Fig. 3B could completely match the standard face-centered cubic crystal of Ag (PDF card JCPDS no. 87-0597). The atomic ratio of Ag to C within Ag@C NFSs was measured to be 26.89 and 69.41, respectively, by EDS (Fig. 3C). According to the FT-IR spectrum of Ag@C NFSs (Fig. 3D), the wavelengths at 1647, 1773 and 2900 cm^{-1} belonged to the vibrations of C=C, C=O and C–H groups, respectively.^{28,31} The absorption bands in the range of 3300 cm^{-1} to 3500 cm^{-1} were attributed to the O–H stretching vibration.³² The form of carbon in Ag@C NFSs was further investigated using Raman spectroscopy. As shown in Fig. 3E, two characteristic peaks located at 1356 cm^{-1} and 1584 cm^{-1} could be easily seen that corresponded to the D band and G band, respectively, indicative of two forms of carbon existing within Ag@C NFSs: the disordered amorphous carbon and the crystalline graphite carbon.³³ Here, the prominent G

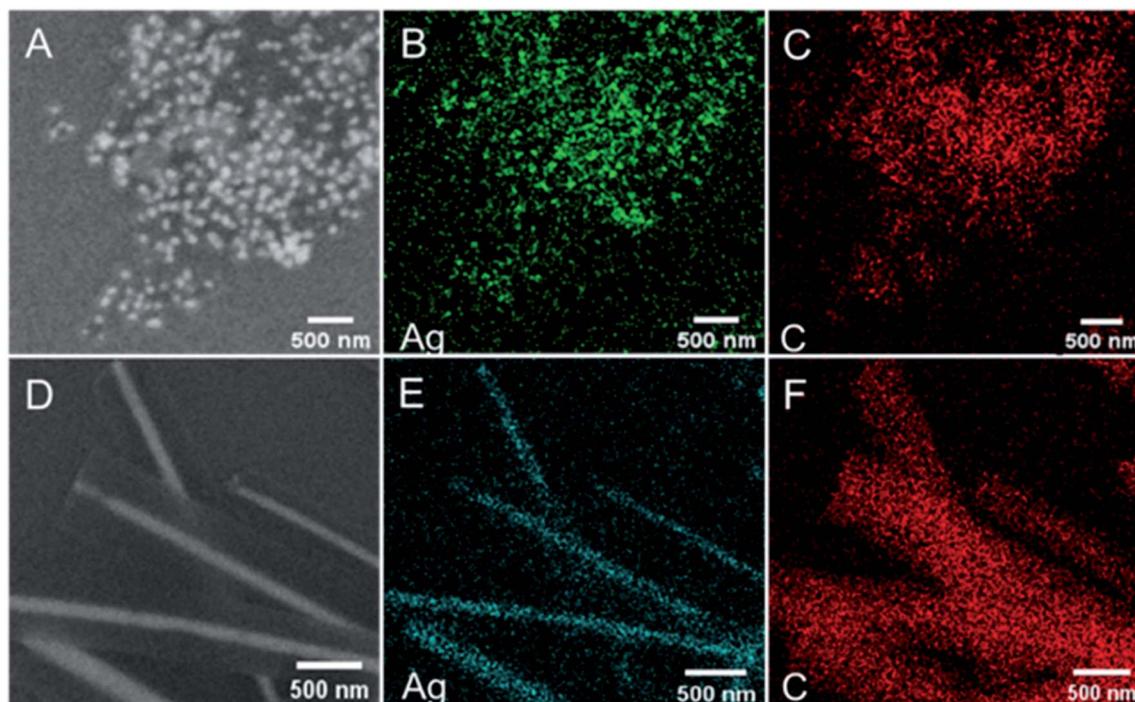


Fig. 2 SEM images of (A) Ag@C NFSs and (D) Ag@C NCs and energy-dispersive X-ray mapping images of Ag and C in (B and C) Ag@C NFSs and (E and F) Ag@C NCs.



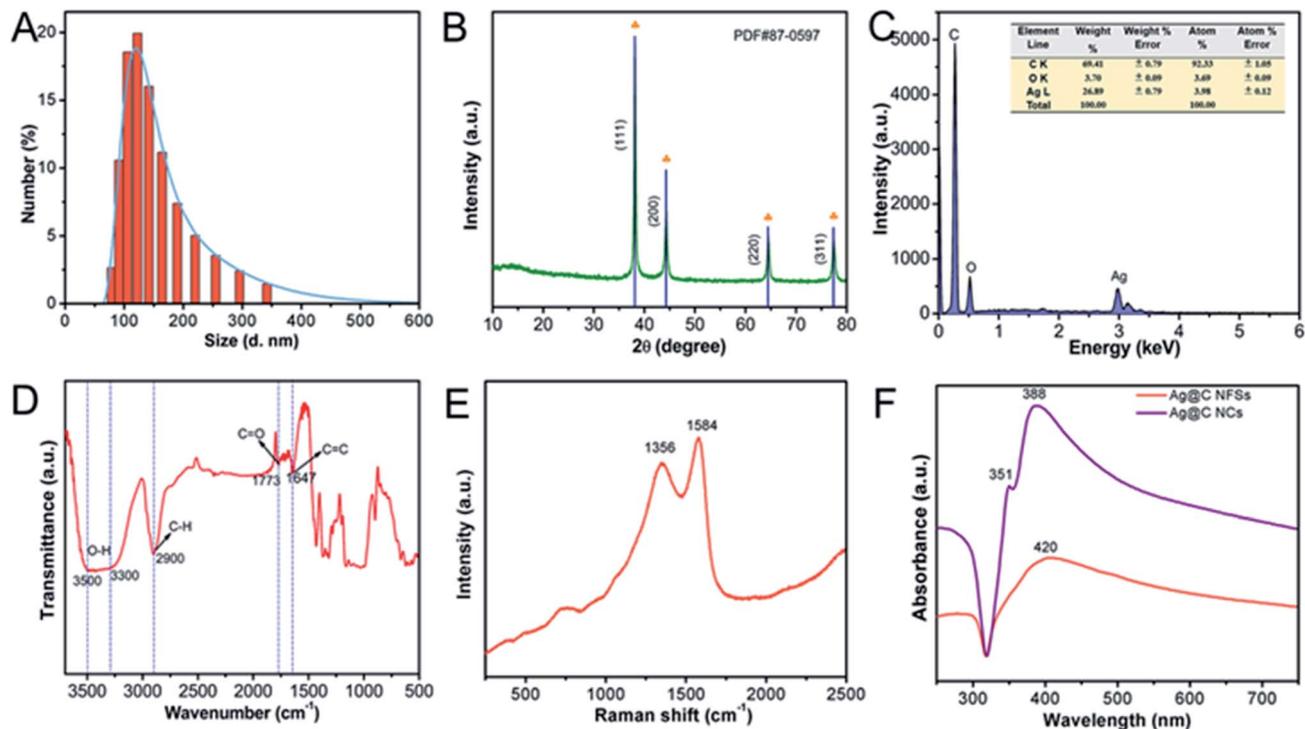


Fig. 3 (A) Hydrodynamic diameter of Ag@C NFSS. (B) XRD pattern of Ag@C NFSS. (C) The EDS of Ag@C NFSS. (D) FT-IR spectra of Ag@C NFSS. (E) Raman spectrum of Ag@C NFSS. (F) UV-vis absorbance spectra of Ag@C NFSS and Ag@C NCs.

band means that the as-formed carbon shell was mainly composed of highly graphitized ordered carbon, which contributes to the fast electron transfer over the electrode. Through the ultraviolet absorption spectrum, it could be clearly seen that the absorption of Ag@C NCs happened at 420 nm, while the absorption of Ag@C NFSS happened at 351 nm and 388 nm. The observations were consistent with the respective characteristic UV-vis absorption peaks previously reported.^{31,34}

3.2. The electrochemical behaviors of the Ag@C NFS modified electrode

Both the electrochemical impedance spectroscopy (EIS) and cyclic voltammetry (CV) behaviours of the Ag@C NFS modified electrode (Ag@C NFSS/GCE) were examined by comparison with that of the Ag@C NC modified GCE (Ag@C NCs/GCE). As depicted in Fig. 4A and B, Ag@C NFSS as a substrate material exhibited obviously better electrochemical performances through showing a smaller circle diameter of EIS and higher CV peak current than Ag@C NCs, which might be because of the significant difference in their morphology and structure. The frogspawn-like structure endowed Ag@C NFSS with a larger surface area and different crystal structure from Ag@C NCs, which likely contributed to the improved electrochemical behaviours of Ag@C NFSS/GCE. Additionally, a slight peak potential shift was observed after electrode modifications. The possible reason might be ascribed to the formation of different double electric layers on the electrode surfaces modified with different materials which might cause different over-potential of the electrochemical reaction.

In addition, the relationship between peak currents and different scan rates was systematically studied to determine the electroactive surface area of Ag@C NFSS/GCE, as shown in Fig. 4C. Obviously, the redox peak currents (I) were proportional to the square root of scan rates ($\nu^{1/2}$) in the range of scan rate from 50 mV s^{-1} to 290 mV s^{-1} . The regression equations of both anodic (I_{pa}) and cathodic (I_{pc}) peaks were $I_{\text{pa}} = 199.09\nu^{1/2} + 25.13$ and $I_{\text{pc}} = -174.46\nu^{1/2} - 30.35$, respectively, indicating that the redox reaction was controlled by diffusion.³⁵ Generally, the electro-active surface area of the modified electrode can be calculated according to the Randles-Sevcik equation $I = 2.69 \times 105A \times D^{1/2}n^{3/2}\nu^{1/2}C$,³⁶ where n is the number of electrons transferred in the redox reaction ($n = 1$), A is the electrode electrochemical active area, D is the diffusion coefficient ($6.70 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$, 25°C), C is the concentration of the reactant ($5 \times 10^{-3} \text{ mol L}^{-1} \text{ Fe}(\text{CN})_6^{3-/4-}$), ν is the scan rate (V s^{-1}) and I is the peak current (μA) of the CV measurement. Here, the electro-active surface area of Ag@C NFSS/GCE was calculated to be 0.0827 cm^2 , 1.17 times higher than the initial surface area of the bare GCE ($d = 3 \text{ mm}$, 0.0707 cm^2), suggesting that Ag@C NFSS would greatly enlarge the surface area of the modified electrode, leading to remarkably accelerated electron transfer on the electrode surface. Moreover, the electron transfer rate constant (k_s) was also used to investigate the electrical conductivity of the Ag@C NFS modified electrode. Fig. 4D shows the relationship between peak potentials (E_{pa} and E_{pc}) and denary logarithm of scan rate ($\lg \nu$), where both E_{pa} and E_{pc} were linearly related to $\lg \nu$ with two linear regression equations for the redox peaks achieved as follows: $E_{\text{pa}} = 0.9972 \lg \nu + 0.4493$ and $E_{\text{pc}} =$



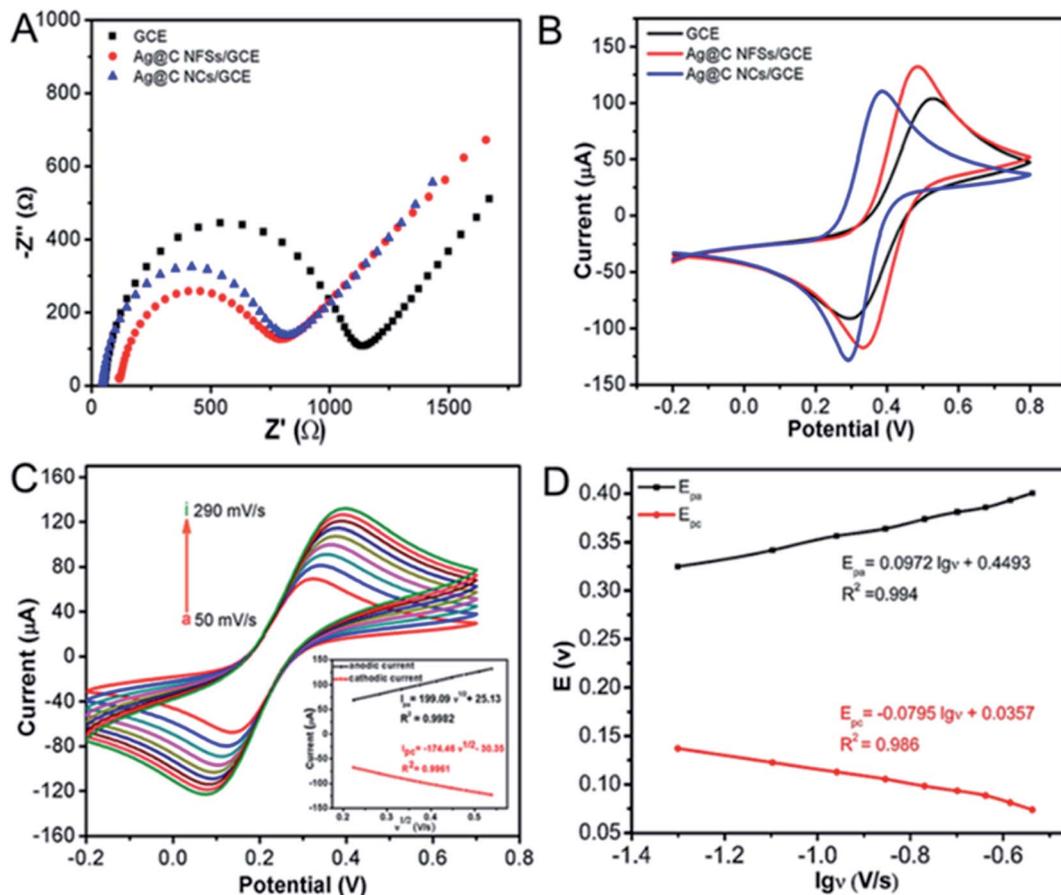


Fig. 4 (A) EIS and (B) CV curves of bare GCE, Ag@C NFSSs/GCE and Ag@C NCs/GCE recorded under the same conditions. (C) Influence of scan rate on the electrochemical responses of Ag@C NFSSs/GCE in 5 mM $[\text{Fe}(\text{CN})_6]^{3-/4-}$ in the scanning potential range of -0.2 to 0.6 V. Scan rates were set at 50, 80, 110, 140, 170, 200, 230, 260 and 290 mV s^{-1} (from a to i), respectively (inset: linear relationship of I_{pa} (black line) and I_{pc} (red line) versus $\nu^{1/2}$). (D) The relationship between peak potentials (E_{pc} and E_{pa}) and $\lg \nu$.

$-0.0795 \lg \nu + 0.0357$, respectively. The k_s was calculated based on the following Laviron equations:³⁷

$$\lg \frac{k_a}{k_c} = \lg \left(\frac{\alpha}{1 - \alpha} \right) \quad (1)$$

$$\lg k_s = \alpha(1 - \alpha) + (1 - \alpha) \lg \alpha - \lg \frac{RT}{nF\nu} - \alpha(1 - \alpha) \frac{nF\Delta E_p}{2.3RT} \quad (2)$$

where k_a and k_c are the slopes of E_{pa} versus $\lg \nu$ and E_{pc} versus $\lg \nu$, respectively, α is the charge-transfer coefficient, R is Faraday constant, n is the apparent number of electrons transferred, ν is the scan rate, T is room temperature and ΔE_p is the potential difference of the redox peaks. The final value of k_s was estimated to be 1.07 s^{-1} , much higher than that (0.584 s^{-1}) of the previously reported silver nanowire-polyaniline nanocomposite based electrochemical sensor, further verifying the greatly improved electron transfer behavior of Ag@C NFSSs/GCE.³⁸

3.3. Biocompatibility of Ag@C NFSSs

The biocompatibility of Ag@C NFSSs plays a crucial role in the construction of the electrochemical immunosensor, so the hydrophilicity, cytotoxicity and stability of the obtained

nanocomposites were investigated to assess the biocompatibility. Fig. 5A and B show the contact angles of Ag@C NFSSs and Ag@C NCs before and after the immobilization of Ab. Clearly, the contact angle of Ag@C NFSSs was about 32° lower than that of Ag@C NCs, revealing the obviously better hydrophilicity of Ag@C NFSSs. After the incubation with Ab, the contact angle of Ag@C NFSSs greatly decreased to $\sim 15^\circ$, while the Ag@C NCs still showed a contact angle as high as 54° , further verifying the excellent hydrophilicity of Ag@C NFSSs favorable for immobilizing biomolecules. Moreover, the cytotoxicities of both Ag@C NFSSs and Ag@C NCs were carefully examined using cancer cells, 4T1 cells (Fig. 5D). After the incubation of 4T1 cells with the Ag@C NFS and Ag@C NC suspensions at various concentrations (12.5 , 25 , 50 , 100 , and $200 \mu\text{g mL}^{-1}$) for 24 h, the cell viability was measured to be above 90% for Ag@C NFSSs by the Cell Counting Kit-8 (CCK-8) method within the set concentration range of Ag@C NFSSs, suggesting no obvious cytotoxicity of the Ag@C NFSSs toward 4T1 cells, while the Ag@C NCs demonstrated relatively poor biocompatibility (Fig. 5E). Moreover, the remarkable stability of Ag@C NFSSs against the tested media was observed. As shown in Fig. 5C and F, Ag@C NFSSs exhibited excellent dispersity in aqueous solution as no sedimentation was seen after standing for 30 min, while obvious



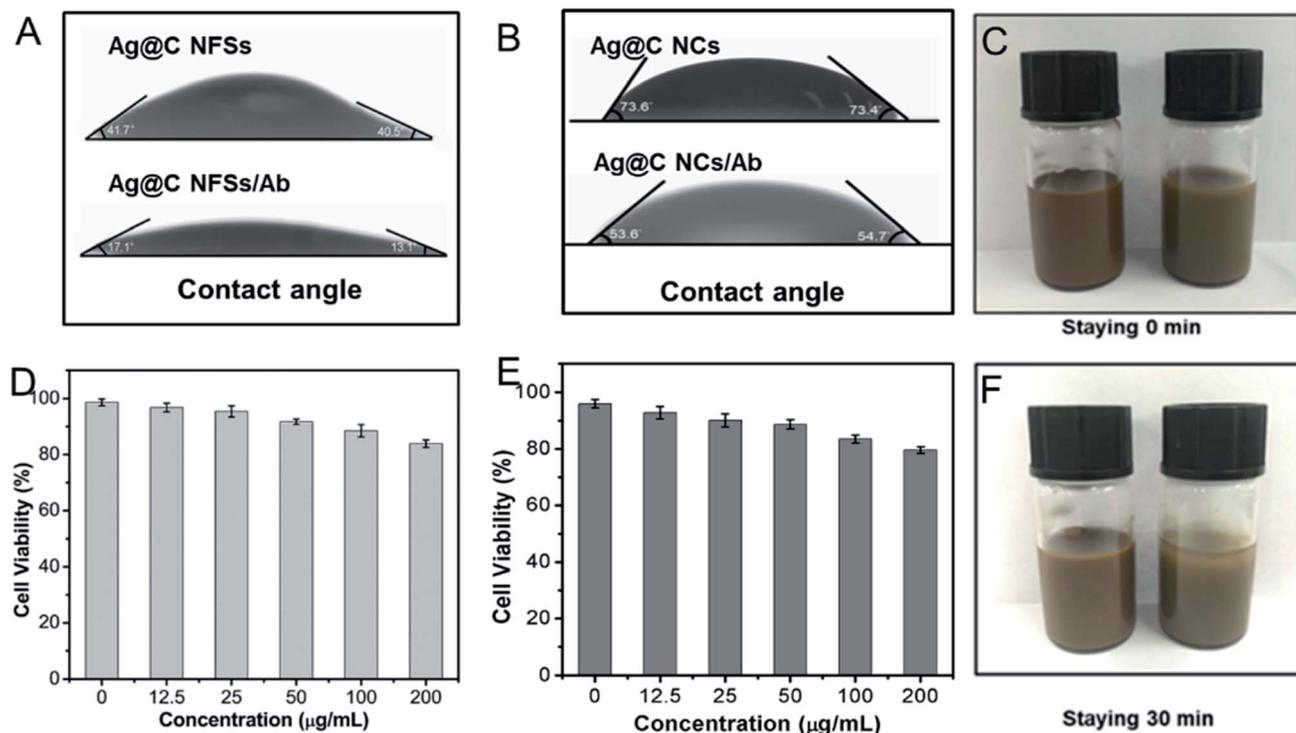


Fig. 5 Contact angles of (A) Ag@C NFSs and (B) Ag@C NCs before and after being modified with anti-CEA. Relative viabilities of 4T1 cells incubated with Ag@C NFSs and Ag@C NCs (D) at various concentrations (12.5, 25, 50, 100, and 200 $\mu\text{g mL}^{-1}$) for 24 h. (D and E) Photographic images of Ag@C NFSs (left) and Ag@C NCs (right) dissolved in water at different standing times (C and F).

stratification was observed in the Ag@C NC aqueous solution. All the results clearly confirmed the favorable biocompatibility of the obtained Ag@C NFSs preferred for the fabrication of the immunosensor. Thereby, the Ag@C NFSs were selected to fabricate the subsequent electrochemical immunosensing platform for CEA detection.

3.4. Fabrication of the Ag@C NFS-based immunosensor

CV was commonly adopted to investigate the electrode interface properties of the immunosensor during the preparation process. Fig. 6A illustrates the CV plots during the preparation

of the Ag@C NFS-based immunosensors. Compared with the bare GCE the modification of Ag@C NFSs greatly improved the current response signal of the electrode; this might benefit from the unique 3D frogspawn-like structure of the Ag@C nanocomposite with a large conductive specific area and the synergistic effect of silver cores and graphite carbon. Although the successive immobilization of Ab, BSA and CEA weakened the current of the modified electrode due to the introduction of non-conductive protein molecules blocking the electron-transfer process of the probe, the current was still high enough for the following CEA detection. On the other hand, the

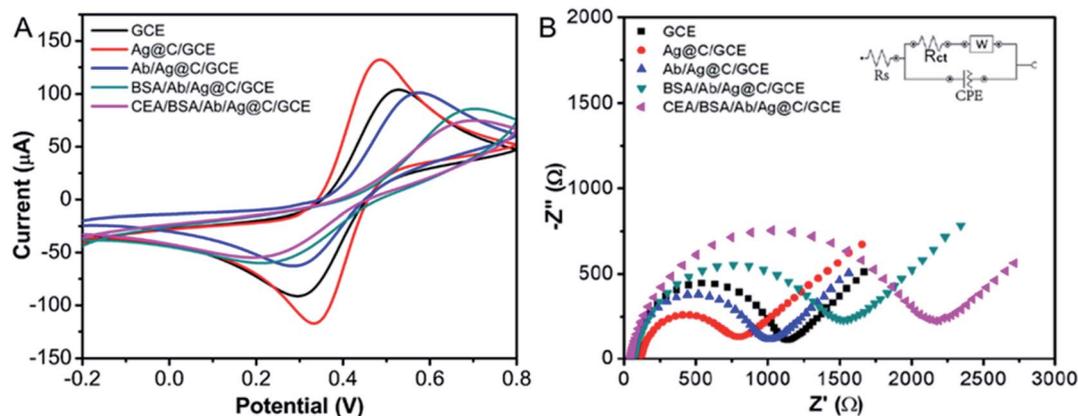


Fig. 6 (A) CV curves and (B) Nyquist diagram of AC impedance of bare GCE, Ag@C NFSs/GCE, Ab/Ag@C NFSs/GCE, BSA/Ab/Ag@C NFSs/GCE, and CEA/BSA/Ab/Ag@C NFSs/GCE by using 5.0 mM $[\text{Fe}(\text{CN})_6]^{3-/4-}$ as an electrochemical indicator, respectively.



Table 1 Simulation parameters of the equivalent circuit components

Electrode	R_s (Ω)	R_{ct} (Ω)	C_{dl}	Z_w
GCE	54.8	1030	1.2676×10^{-6}	0.001416
Ag@C NFSs/GCE	11.3	669	2.6506×10^{-6}	0.001083
Ab1/Ag@C NFSs/GCE	51.3	903	1.4113×10^{-6}	0.001562
BSA/Ab1/Ag@C NFSs/GCE	70.3	1360	2.6766×10^{-6}	0.001038
CEA/BSA/Ab1/Ag@C NFSs/GCE	45.7	1790	1.9889×10^{-6}	0.001349

results confirm the successful construction of the Ag@C NFS-based immunosensors to detect CEA.

Meanwhile, the Nyquist diagram of AC impedance spectroscopy was also used to confirm the stepwise modification process of the Ag@C NFS-based immunosensor (Fig. 6B). In general, the Nyquist diagram is made of a linear portion and semicircular portion which correspond to the diffusion process and the electron transfer resistance (R_{ct}), respectively. As seen in Fig. 6B, the bare GCE showed a very small semicircle, while the semicircle diameter clearly decreased when the Ag@C NFSs were dropped onto the spotless GCE surface by virtue of the good electron transfer behavior of Ag@C NFSs. Subsequently, the semicircle domain was increased in sequence with the

successive modification of Ab and BSA, indicating the successful immobilization of non-conductive biomolecules onto the electrode. The largest semicircle diameter appeared after the incubation of CEA with the above modified electrode, revealing the successful specific recognition between Ab and CEA. The model of the Randles equivalent circuit, which consists of the resistance of solution (R_s), the charge-transfer resistance (R_{ct}), the Warburg impedance (W), and the double layer capacitance (CPE), was fitted as shown in Fig. 6B (inset). Moreover, the corresponding values of stepwise modification were simulated with the aid of ZSimpwin software and are listed in Table 1. Obviously, the variation rule of R_{ct} was consistent with the changes of the semicircle domain, indicative of the effective

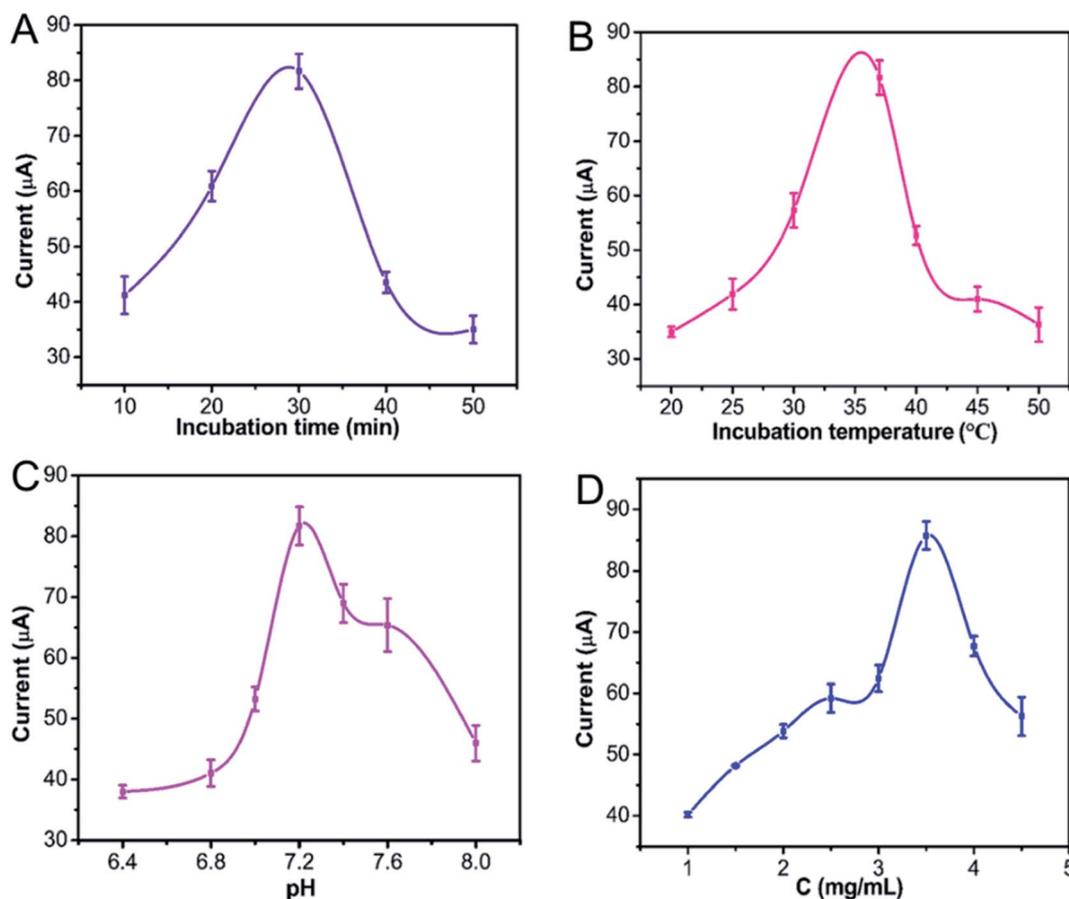


Fig. 7 Effects of (A) the incubation time, (B) incubation temperature, (C) pH of PBS and (D) Ag@C NFS concentration on the current response of the Ag@C NFS-based immunosensor. Error bar refers to the relative standard deviation (RSD, $n = 3$).



fabrication of the proposed Ag@C NFS-based electrochemical immunosensor for CEA detection.

3.5. Optimization of the experimental conditions

The optimum working condition of the immunosensor is crucial to sensitive CEA detection. Thus, the experimental conditions of the Ag@C NFS-based immunosensor were optimized by differential pulse voltammetry (DPV), including the incubation time and temperature of CEA with Ab, the pH of PBS and the concentrations of Ag@C NFSs. The effect of incubation time of Ab with CEA on the electrochemical performance of the Ag@C NFS-based immunosensor was first studied. Fig. 7A shows that the peak current of the prepared immunosensor increased with the time increasing from 10 to 30 min and went down after 30 min. Thus, 30 min was selected as the optimal incubation time between Ab and Ag. In the optimum incubation period, the optimum incubation temperature was investigated in the range of 20–50 °C (Fig. 7B). Clearly the highest peak current appeared at 37 °C, preferred for the specific interaction between Ab and Ag. A temperature lower or higher than 37 °C was not good to maintain the biological activity of protein molecules. Therefore, the optimal incubation temperature was set at 37 °C for the interaction between Ab and Ag. In addition, the acidity of aqueous buffer influences the biological activity of protein molecules as well,³⁹ so the effect of pH on the current response of the Ag@C NFS-based immunosensor toward CEA was also optimized. As demonstrated in Fig. 7C, within the pH range of 6.4–8.0 in PBS solution, the current signal increased with pH increasing until 7.2 and irreversibly dropped afterwards as the alkalinity increased constantly. The maximum signal response showed up at pH 7.2 which was very close to the physiological pH of the human body, favorable for the binding interaction. Therefore, the PBS solution at pH 7.2 was chosen as the detection medium of CEA. Ag@C NFSs are another un-neglectable important factor and majorly contributed to the immunosensing performance of the prepared immunosensor, so the concentration of Ag@C NFSs ranging from 1 mg mL⁻¹ to 3.5 mg mL⁻¹ was optimized to receive the best response signal

to CEA detection. As shown in Fig. 7D, when the Ag@C NFS concentration increased, the response current went up dramatically and reached the highest at 3.5 mg mL⁻¹, but went down with the concentration exceeding 3.5 mg mL⁻¹. The increase of Ag@C NFS concentration not only improved the electrode conductivity but also boosted the affinity binding between Ab and Ag;^{27,40} however, when too many Ag@C NFSs were dropped on the electrode surface, the composite layer was too thick to facilitate electron transfer and efficient specific interaction between Ab and CEA. Hence, 3.5 mg mL⁻¹ was used as the most appropriate concentration of Ag@C NFS to fabricate the immunosensor.

3.6. Analytical performance of the Ag@C NFS-based immunosensor

In order to assess the detection performance of the as-prepared immunosensor, a series of different CEA concentrations were detected using the Ag@C NFS-based immunosensor by DPV under the optimal experimental conditions. Fig. 8A shows the DPV curves for the detection of CEA at concentrations ranging from 0.1 pg mL⁻¹ to 100 ng mL⁻¹, where the current signal was found to decrease gradually with the increase of the CEA concentration. The linear equation of the Ag@C NFS-based immunosensor toward CEA detection was $y = -6.6743 \lg c + 66.9988$ ($R^2 = 0.996$ with the limit of detection (LOD) calculated to be 5.12 pg mL⁻¹ (signal to noise ratio = 3), much lower than many existing ones). Given its wide linear range and low LOD, it is of great potential to detect CEA in the early clinical diagnosis of tumors. Upon comparison with some previously reported noble metal-based CEA immunoassays (Table 2), the proposed Ag@C NFS-based electrochemical immunosensor exhibited a significantly lower LOD and superior sensitivity for the detection of CEA.

3.7. Selectivity, reproducibility and stability

In order to assess the reproducibility of the immunosensor, five immunosensors were constructed to detect 10 ng mL⁻¹ CEA. As shown in Fig. 9A, the current responses of the five replicative

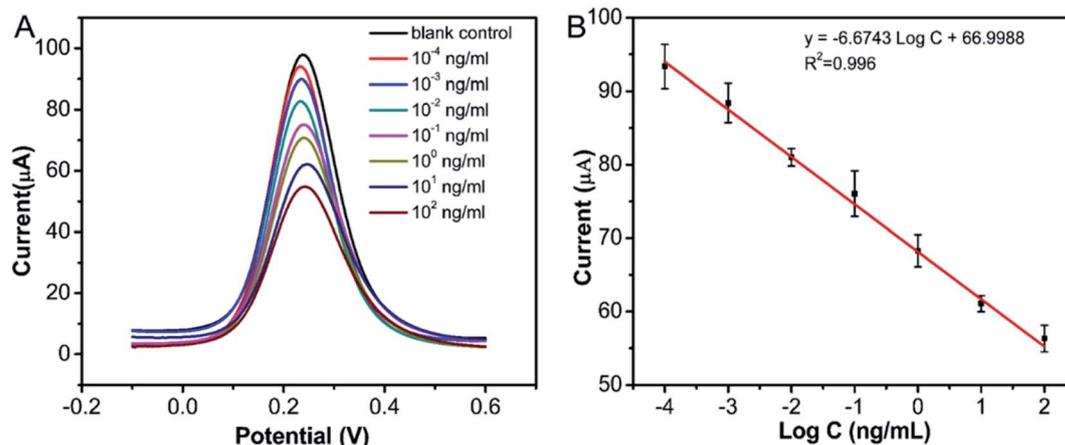


Fig. 8 (A) DPVs and (B) calibration curve obtained by detecting CEA at different concentrations with the proposed Ag@C NFS-based electrochemical immunosensor. Error bar = RSD ($n = 3$).



Table 2 Comparison of the analytical performance of the proposed Ag@C NFS-based electrochemical immunosensor with other existing immunosensors for the detection of CEA^a

Immunosensor	Line range (ng mL ⁻¹)	LOD (ng mL ⁻¹)	Reference
Au-TiO ₂ film	0.2–160	0.06	41
g-C ₃ N ₄ NS	0.1–150	0.06	42
rGO	0.1–5	0.05	43
Au NPs–MWCNTs	0.4–125	0.09	44
GO/MWCNTs–COOH/Au@CeO ₂	0.05–100	0.02	45
NCMTs@Fe ₃ O ₄ @Cu silicate	0.03–6	0.00538	46
Fe ₃ O ₄ @Au NPs	0.001–100	0.01	19
Au@CeBi _{0.4} O _{3.7}	0.01–100	0.00012	47
Au-PB	0.005–50	0.0033	48
Pt NPs@rGO@PS NSs	0.05–70	0.01	49
AuNPs/PB-PEDOT	0.05–40	0.01	50
Silicon nanoribbon	0.01–100	0.01	51
BSA-NC-rGO	0.01–200	0.067	52
Ag@C NFSs	0.0001–100	0.00512	This work

^a NS: nanosheet; MWCNTs: multi walled carbon nanotubes; NCMTs: nitrogen-doped magnetic carbon microtubes; PB: Prussian blue; PS NSs: polystyrene nanospheres; NC: nitidine chloride.

immunosensors were basically the same and the RSD was less than 5%, exhibiting the good reproducibility of the proposed immunosensor. In addition, the stability also affects the detection performance of the immunosensor. As shown in Fig. 9B and C, after the immunosensor was incubated with 0.01

ng mL⁻¹ CEA, the current fluctuation was less than 5% within the successive CV scans of 30 cycles, proving that the proposed Ag@C NFS-based immunosensor had an acceptable short-term stability. Additionally, the long-term storage stability of the Ag@C NFS-based immunosensor was further investigated after

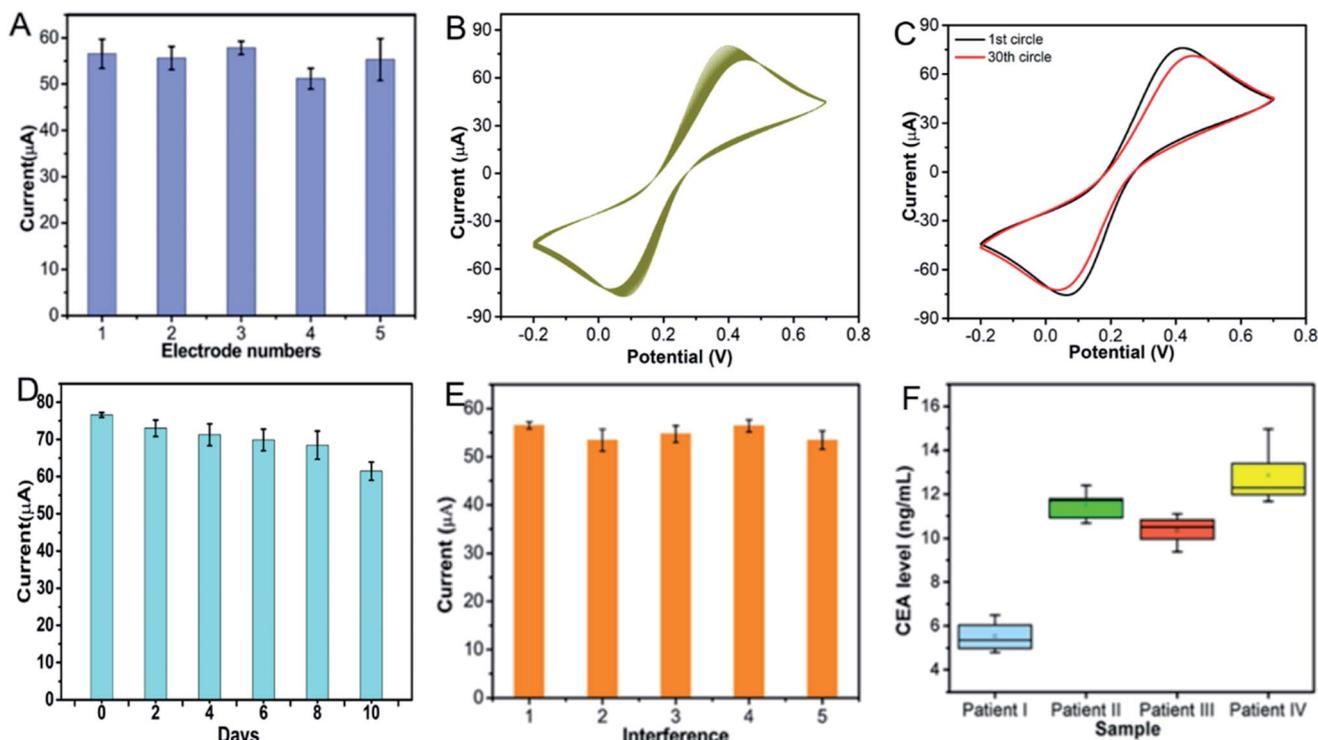


Fig. 9 (A) Reproducibility and (B–C) short-term and (D) long-term stabilities of the proposed Ag@C NFS-based immunosensors. (E) Influence of various interfering proteins on the current responses of the proposed immunosensor toward CEA. (Groups 1–5 refer to 10 ng mL⁻¹ CEA, 10 ng mL⁻¹ CEA + 10 ng mL⁻¹ BSA, 10 ng mL⁻¹ CEA + 10 ng mL⁻¹ AFP, 10 ng mL⁻¹ CEA + 10 ng mL⁻¹ IgG and 10 ng mL⁻¹ CEA + 10 ng mL⁻¹ PSA, respectively.) Error bar = RSD ($n = 3$). (F) The original CEA levels in blood samples from four cancerous patients detected with the proposed Ag@C NFS-based immunosensors.



Table 3 Human serum sample analysis using the proposed method and the radionuclide method

Content of CEA in the serum (ng mL ⁻¹)	The addition content (ng mL ⁻¹)	The detection content (ng mL ⁻¹)	Radionuclide method (ng mL ⁻¹)	Recovery (%)
Patient I	—	5.36	5.41	—
	2.5	8.12	—	108.4
	5	10.28	—	97.4
	10	14.69	—	92.8
Patient II	—	11.36	11.7	—
	2.5	13.96	—	90.4
	5	16.48	—	95.6
	10	22.31	—	106.1
Patient III	—	10.76	10.82	—
	2.5	13.49	—	106.8
	5	15.93	—	102.2
	10	20.68	—	98.6
Patient IV	—	15.21	14.96	—
	2.5	17.69	—	109.2
	5	20.21	—	105
	10	25.18	—	102.2

storage at 4 °C. As demonstrated in Fig. 9D, the current still retained 86.6% after 10 days, indicative of a desirable long-term storage stability. At the same time, the selectivity of the immunosensor is vital to detect the target marker accurately. To probe into the selectivity of the proposed immunosensor toward the detection of 10 ng mL⁻¹ CEA, five interferences of BAS, AFP, IgG and PSA at 10 ng mL⁻¹ were chosen as co-existing interferences. Among the five groups, group 1 was CEA only at 10 ng mL⁻¹, and the other four groups were 10 ng mL⁻¹ CEA containing 10 ng mL⁻¹ interferences, respectively. Fig. 9E shows that the current signal of the proposed immunosensor was not obviously interfered proteins, revealing that the Ag@C NFS-based immunosensor possessed an admirable specificity for CEA detection.

3.8. Real sample analysis

Given its brilliant performance, the newly proposed Ag@C NFS-based immunosensor was used to determine the CEA level in cancerous human serum samples in order to explore its practical application. Here, four blood samples from four cancer patients (I–IV) were detected as shown in Fig. 9F. Obviously, the detected CEA levels in the blood samples from these four patients were much higher than the normal values and in good accordance with that measured by radionuclide methods, as illustrated in Table 3. Additionally, the standard addition method was also used to evaluate the practicability of the proposed immunosensor. Three CEA concentrations of 2.5, 5 and 10 ng mL⁻¹ were added in the four serum samples, respectively with a recovery rate over 90% and up to 109.2%. These results are further proof of the superb practical feasibility of the label-free immunosensor using the Ag@C NFSs as an electrochemical immunosensing platform.

4. Conclusion

In summary, a convenient and effective label-free immunosensor was established for the detection of CEA in clinical

samples with the newly prepared frogspawn-like Ag@C nanocomposites prepared by the one-pot reaction of AgNO₃, CTAB and AA. Compared with the prepared Ag@C NCs, the as-prepared Ag@C NFSs exhibited a larger surface area, faster electron transfer rate and better biocompatibility owing to the different crystal structure and synergistic effect of silver and nanocarbon, which is preferred for the fabrication of electrochemical immunosensors. Using the Ag@C NFSs to capture anti-CEA, the resultant immunosensing platform demonstrated excellent analytical performance for CEA detection in a wide CEA concentration range with a fairly low LOD. With high reproducibility, good stability and preferable interference immunity, the CEA levels in four blood samples from four cancer patients were accurately determined with the proposed method which were highly consistent with that obtained by the radionuclide method, implying its great potential in practical application for early clinical cancer diagnosis. Overall, this simple label-free immunosensor may be considered as a promising alternative to detect other biomarkers in early clinical diagnosis.

Conflicts of interest

There are no conflicts to declare.

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References

- Q. Q. Yan, L. Wang, F. X. Song, Y. J. Zhang, L. Gao, F. J. Liu and Y. X. Shi, *J. Mater. Res.*, 2020, **35**, 591–599.
- P. A. Crosbie, R. Gabe, I. Simmonds, M. Kennedy, S. Rogerson, N. Ahmed, D. R. Baldwin, R. Booton,



- A. Cochrane, M. Darby, K. Franks, S. Hinde, S. M. Janes, U. Macleod, M. Messenger, H. Moller, R. L. Murray, R. D. Neal, S. L. Quaipe, M. Sculpher, P. Tharmanathan, D. Torgerson and M. E. Callister, *BMJ Open*, 2020, **10**, e037075.
- 3 M. N. Fadhel, S. Appak Baskoy, Y. Wang, E. Hysi and M. C. Kolios, *J. Biophotonics*, 2020, e202000209.
- 4 D. Wu, H. Ma, Y. Zhang, H. Jia, T. Yan and Q. Wei, *ACS Appl. Mater. Interfaces*, 2015, **7**, 18786–18793.
- 5 X. Zhang, Y. Li, H. Lv, Z. Gao, C. Zhang, S. Zhang, Y. Wang, Z. Xu and Z. Zhao, *J. Electrochem. Soc.*, 2018, **165**, B931–B938.
- 6 Y. Chen, A. J. Wang, P. X. Yuan, X. Luo, Y. Xue and J. J. Feng, *Biosens. Bioelectron.*, 2019, **132**, 294–301.
- 7 C. Zhang, S. Zhang, Y. Jia, Y. Li, P. Wang, Q. Liu, Z. Xu, X. Li and Y. Dong, *Biosens. Bioelectron.*, 2019, **126**, 785–791.
- 8 N. L. Li, L. P. Jia, R. N. Ma, W. L. Jia, Y. Y. Lu, S. S. Shi and H. S. Wang, *Biosens. Bioelectron.*, 2017, **89**, 453–460.
- 9 H. Yan, L. Jiao, H. Wang, W. Xu, Y. Wu, W. Gu, D. Du, Y. Lin and C. Zhu, *Sens. Actuators, B*, 2019, **297**, 126760.
- 10 Q. Han, R. Wang, B. Xing, T. Zhang, M. S. Khan, D. Wu and Q. Wei, *Biosens. Bioelectron.*, 2018, **99**, 493–499.
- 11 E. Tan, N. Gouvas, R. J. Nicholls, P. Ziprin, E. Xynos and P. P. Tekkis, *Surg. Oncol.*, 2009, **18**, 15–24.
- 12 Y. Jia, Y. Li, S. Zhang, P. Wang, Q. Liu and Y. Dong, *Biosens. Bioelectron.*, 2020, **149**, 111842.
- 13 D. Song, J. Zheng, N. V. Myung, J. Xu and M. Zhang, *Talanta*, 2021, **225**, 122006.
- 14 S. H. Shamsuddin, T. D. Gibson, D. C. Tomlinson, M. J. McPherson, D. G. Jayne and P. A. Millner, *Biosens. Bioelectron.*, 2021, **178**, 113013.
- 15 H. Q. Wang and Z. F. Ma, *Biosens. Bioelectron.*, 2020, **169**, 112644.
- 16 W. J. Li, Y. Yang, C. Y. Ma, Y. J. Song, X. W. Qiao and C. L. Hong, *Talanta*, 2020, **219**, 121322.
- 17 Y. Song, K. Cao, W. Li, C. Ma, X. Qiao, H. Li and C. Hong, *Microchem. J.*, 2020, **155**, 104694.
- 18 F. Li, Y. Li, Y. Dong, L. Jiang, P. Wang, Q. Liu, H. Liu and Q. Wei, *Sci. Rep.*, 2016, **6**, 21281.
- 19 P. Butmee, G. Tumcharern, G. Thouand, K. Kalcher and A. Samphao, *Bioelectrochemistry*, 2020, **132**, 107452.
- 20 S. Xu, R. L. Zhang, W. Zhao, Y. Zhu, W. Wei, X. Y. Liu and J. Luo, *Biosens. Bioelectron.*, 2017, **92**, 570–576.
- 21 A. L. Sun, Q. A. Qi and Z. L. Dong, *Protein Pept. Lett.*, 2008, **15**, 782–788.
- 22 M. Rizwan, S. Elma, S. A. Lim and M. U. Ahmed, *Biosens. Bioelectron.*, 2018, **107**, 211–217.
- 23 W. Li, X. Qiao, C. Hong, C. Ma and Y. Song, *Anal. Biochem.*, 2020, **592**, 113566.
- 24 J. Duan, L. Bai, K. Xu, Q. Fang, Y. Sun, H. Xu, K. C.-F. Leung and S. Xuan, *J. Hazard. Mater.*, 2020, **384**, 121276.
- 25 Y. Wang, D. Fan, D. Wu, Y. Zhang, H. Ma, B. Du and Q. Wei, *Sens. Actuators, B*, 2016, **236**, 241–248.
- 26 C. J. Mao, X. B. Chen, H. L. Niu, J. M. Song, S. Y. Zhang and R. J. Cui, *Biosens. Bioelectron.*, 2012, **31**, 544–547.
- 27 F. Li, J. Feng, Z. Gao, L. Shi, D. Wu, B. Du and Q. Wei, *ACS Appl. Mater. Interfaces*, 2019, **11**, 8945–8953.
- 28 W. Zhang, L. Hao, Q. Lin, C. Lu, Z. Xu and X. Chen, *Mater. Sci. Eng., B*, 2014, **190**, 1–6.
- 29 X. Liu, C. Chen, H. Ye, Y. Jia, Y. Wu, A. Jin, Y. Wang and X. Chen, *Carbon*, 2018, **131**, 213–222.
- 30 H. Niu, X. Wu, M. Qiu, Y. Gao, J. Song, C. Mao, S. Zhang, H. Liu and J. Wu, *J. Mater. Res.*, 2011, **26**, 2780–2794.
- 31 X. M. Sun and Y. D. Li, *Langmuir*, 2005, **21**, 6019–6024.
- 32 Z. Chen, M. Niu, G. Chen, Q. Wu, L. Tan, C. Fu, X. Ren, H. Zhong, K. Xu and X. Meng, *ACS Nano*, 2018, **12**, 12721–12732.
- 33 S. A. Nakhjavani, H. Afsharan, B. Khalilzadeh, M. H. Ghahremani, S. Carrara and Y. Omid, *Biosens. Bioelectron.*, 2019, **141**, 111439.
- 34 L. Yan, Y. Hao, X. Feng, Y. Yang, X. Liu, Y. Chen and B. Xu, *RSC Adv.*, 2015, **5**, 79860–79867.
- 35 Z. Wang, X. Tian, D. Sun, P. Cao, M. Ding, Y. Li, N. Guo, R. Ouyang and Y. Miao, *RSC Adv.*, 2020, **10**, 15870–15880.
- 36 J. Xue, L. Yang, Y. Jia, H. Wang, N. Zhang, X. Ren, H. Ma, Q. Wei and H. Ju, *ACS Sens.*, 2019, **4**, 2825–2831.
- 37 P. Wang, F. Pei, E. Ma, Q. Yang, H. Yu, J. Liu, Y. Li, Q. Liu, Y. Dong and H. Zhu, *Bioelectrochemistry*, 2020, **131**, 107352.
- 38 Z. Chao, S. Govindaraju, K. Giribabu, Y. S. Huh and K. Yun, *Sens. Actuators, B*, 2017, **252**, 616–623.
- 39 J. Miao, K. Du, X. Li, X. Xu, X. Dong, J. Fang, W. Cao and Q. Wei, *Biosens. Bioelectron.*, 2021, **171**, 112713.
- 40 Y. Chen, L. P. Mei, J. J. Feng, P. X. Yuan, X. Luo and A. J. Wang, *Biosens. Bioelectron.*, 2019, **145**, 111638.
- 41 Y. Zhang, R. Yuan, Y. Chai, Y. Xiang, X. Qian and H. Zhang, *J. Colloid Interface Sci.*, 2010, **348**, 108–113.
- 42 K. L. Lin, T. Yang, H. Y. Zou, Y. F. Li and C. Z. Huang, *Talanta*, 2019, **192**, 400–406.
- 43 M. Jozghorbani, M. Fathi, S. H. Kazemi and N. Alinejadian, *Anal. Biochem.*, 2020, **613**, 114017.
- 44 G. Paimard, M. Shahlaei, P. Moradipour, H. Akbari, M. Jafari and E. Arkan, *Sens. Actuators, B*, 2020, **311**, 127928.
- 45 X. Pang, J. Li, Y. Zhao, D. Wu, Y. Zhang, B. Du, H. Ma and Q. Wei, *ACS Appl. Mater. Interfaces*, 2015, **7**, 19260–19267.
- 46 J. Zheng, J. Wang, D. Song, J. Xu and M. Zhang, *ACS Appl. Nano Mater.*, 2020, **3**, 3449–3458.
- 47 X. Tian, P. Cao, D. Sun, Z. Wang, M. Ding, X. Yang, Y. Li, R. Ouyang and Y. Miao, *Appl. Surf. Sci.*, 2020, **528**, 146956.
- 48 J. Lin, K. Y. Li, M. F. Wang, X. H. Chen, J. Y. Liu and H. L. Tang, *RSC Adv.*, 2020, **10**, 38316–38322.
- 49 Q. C. Lan, C. L. Ren, A. Lambert, G. S. Zhang, J. Li, Q. Cheng, X. Y. Hu and Z. J. Yang, *ACS Sustainable Chem. Eng.*, 2020, **8**, 4392–4399.
- 50 T. T. Yang, Y. S. Gao, Z. Liu, J. K. Xu, L. M. Lu and Y. F. Yu, *Sens. Actuators, B*, 2017, **239**, 76–84.
- 51 Z. T. Bao, J. L. Sun, X. Q. Zhao, Z. Y. Li, S. K. Cui, Q. Y. Meng, Y. Zhang, T. Wang and Y. F. Jiang, *Int. J. Nanomed.*, 2017, **12**, 4623–4631.
- 52 Y. Li, Z. Zhang, Y. T. Zhang, D. M. Deng, L. Q. Luo, B. S. Han and C. H. Fan, *Biosens. Bioelectron.*, 2016, **79**, 536–542.

