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# Ultrasensitive electrochemical biosensor for detection of circulating tumor cells based on a highly efficient enzymatic cascade reaction†

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There has been great interest in the enzymatic cascade amplification strategy for the electrochemical detection of circulating tumor cells (CTCs). In this work, we designed a highly efficient enzymatic cascade reaction based on a multiwalled carbon nanotubes–chitosan (MWCNTs–CS) composite for detection of CTCs. A high electrochemical effective surface area was obtained for a MWCNTs–CS-modified glassy carbon electrode (GCE) for loading glucose oxidase (GOD), as well as a high loading rate and high electrical activity of the enzyme. As a ‘power source’, the MWCNTs–CS composites provided a strong driving power for horseradish peroxidase (HRP) on the surface of polystyrene (PS) microspheres, which acted as probes for capturing CTCs and allowed the reaction to proceed with further facilitation of electron transfer. Aptamer, CTCs, and PS microspheres with HRP and anti-epithelial cell adhesion molecule (anti-EpCAM) antibody were assembled on the MWCNTs–CS/GCE to allow for the modulation of enzyme distance at the micrometer level, and thus ultra-long-range signal transmission was made possible. An ultrasensitive response to CTCs was obtained *via* this proposed sensing strategy, with a linear range from 10 cell mL<sup>−1</sup> to 6 × 10<sup>6</sup> cell mL<sup>−1</sup> and a detection limit of 3 cell mL<sup>−1</sup>. Moreover, this electrochemical sensor possessed the capability to detect CTCs in serum samples with satisfactory accuracy, which indicated great potential for early diagnosis and clinical analysis of cancer.

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## Introduction

Circulating tumor cells (CTCs) are tumor cells that disseminate and survive at low concentrations in the peripheral blood during the development of malignant tumors, and are closely related to tumor metastasis and prognosis.<sup>1,2</sup> There are many methods used for the detection of CTCs, such as liquid biopsy,<sup>3,4</sup> fluorometry,<sup>5</sup> photoelectrochemical biosensors,<sup>6,7</sup> ratiometric detection,<sup>8</sup> flow cytometry,<sup>9</sup> mass spectrometry,<sup>10</sup> and colorimetric analysis.<sup>11</sup> Electrochemical biosensors based on enzymatic cascade reactions have exhibited great potential in clinical analysis and diagnosis of CTCs.<sup>12–14</sup>

In a typical case, the first enzyme's catalytic product is directly used as the second enzyme's catalytic substrate in a one-pot reaction with high activity and stereoselectivity without intermediates, which can provide valuable detection information and understanding of enzymatic reactions in living

systems. The enzymatic cascade amplification strategy has been widely applied to detect biomarkers.<sup>15–17</sup> However, numerous studies have demonstrated that specific parameters have an impact on the effectiveness of enzymatic cascade catalysis. In particular, enzyme–enzyme distance limits enzymatic cascade catalysis, although all types of scaffolds, including metal–organic frameworks,<sup>18</sup> metal nanoparticles,<sup>19</sup> DNA tetrahedra,<sup>20</sup> cellular surfaces,<sup>21</sup> and proteins,<sup>22</sup> have been used for enzyme loading,<sup>23</sup> enzyme activating,<sup>24</sup> and modulating enzyme spacing.<sup>25</sup> For example, when the gap between two enzymes was maintained at a maximum of 6 nm, Wilner *et al.*<sup>26</sup> found that the reaction rate increased 16 fold while a DNA scaffold was employed. Fu *et al.*<sup>27</sup> obtained a 15-fold improvement when a glucose oxidase (GOD)–horseradish peroxidase (HRP) pair at 10 nm was co-assembled in a DNA scaffold. Hence, nanomaterial-based systems have been developed to address these issues due to their promising properties.<sup>28–30</sup>

Carbon nanotubes (CNTs), which are one-dimensional nanomaterials with a distinctive hexagonal shape, possess excellent electrical conductivity and can be used as electrode modifiers to accelerate electrode responses, reduce overpotential, and facilitate electron transfer events.<sup>31,32</sup> However, the widespread use of CNTs has been constrained by their poor solubility in solvents and suboptimal dispersion.<sup>33,34</sup> To achieve

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uniform dispersion of CNTs in solvents, chitosan (CS), a natural polymer rich in amino and hydroxyl groups, can be used to interact with the walls of CNTs based on their hydrophobic ends, while their hydrophilic ends would interact with water or other polar solvents.<sup>35,36</sup> Due to their high surface area for loading and immobilizing enzymes as well as their high enzymatic unit activity, multiwalled carbon nanotubes–chitosan (MWCNTs–CS) composites may have potential applications in increasing the mass transit of substrates in enzymatic cascade catalysis.<sup>37–43</sup>

Herein, we developed an extremely effective enzymatic cascade reaction based on a MWCNTs–CS composite to construct an electrochemical biosensing system for detecting CTCs. In our strategy (Scheme 1), MWCNTs–CS composites were modified on the surface of a glassy carbon electrode (GCE) to obtain GCE/MWCNTs–CS. Then, GOD was loaded onto the surface of the GCE/MWCNTs–CS, and the aptamer for CTCs was modified on the GCE/MWCNTs–CS/GOD *via* amide bonds. In addition, anti-epithelial cell adhesion molecule (anti-EpCAM) and HRP were assembled on the surface of polystyrene (PS) to obtain anti-EpCAM/PS/HRP conjugates. The aptamer on the surface of GCE/MWCNTs–CS/GOD specifically recognizes CTCs, and then the prepared anti-EpCAM/PS/HRP was linked to the surface of CTCs by anti-EpCAM. Thus, an enzymatic cascade reaction was obtained.

Glucose in the electrolyte was initially oxidized by GOD to produce gluconic acid and  $\text{H}_2\text{O}_2$ , and then the  $\text{H}_2\text{O}_2$  was further utilized by HRP to accelerate the oxidation of  $\text{Fe}^{2+}$  with amplified electrochemical responses. The produced  $\text{H}_2\text{O}_2$  by GOD is crucial for the subsequent enzymatic cascade reaction so that the GCE/MWCNTs–CS/GOD provides a driving force for the enzymatic cascade reaction. This MWCNTs–CS composite-based electrochemical biosensing technique triggered a highly effective enzymatic cascade reaction for the detection of CTCs that exhibited a wide detection range, high sensitivity, and low detection limit.

## Experimental

### Materials

A glassy carbon electrode (GCE), chitosan (CS), glutaraldehyde (GA), glucose, and multiwalled carbon nanotubes (MWCNTs) were obtained from Aladdin Chemistry Co., Ltd. (Shanghai, China). Glucose oxidase (GOD,  $>180 \text{ U mg}^{-1}$ ) was purchased from Shanghai Ruiyong Biological Technology Co., Ltd. (Shanghai, China). Bovine serum albumin (BSA), phosphate-buffered saline (PBS), potassium chloride,  $\text{K}_3[\text{Fe}(\text{CN})_6]$ ,  $\text{K}_4[\text{Fe}(\text{CN})_6] \cdot 3\text{H}_2\text{O}$ , carboxylated polystyrene microspheres (PS–COOH), horseradish peroxidase (HRP,  $>300 \text{ U mg}^{-1}$ ), and anti-epithelial cell adhesion molecule antibody (anti-EpCAM) were obtained from Abcam Co., Ltd. (Cambridge, MA, USA). Glacial acetic acid, *N*-(3-(dimethylamino)-propyl)-*N*'-ethyl carbodiimide hydrochloride (EDC), *N*-hydroxysulfosuccinimide (NHS), and oligonucleotides were received from Sangon Biotechnology Co. Ltd. (Shanghai, China).

The aptamer sequence for MCF-7 cells: 5'-COOH-AAAAA-CACTACAGAGGTTGCGTCTGTCCACGTTGTCATGGGGGGTTGGCCTG-3'.

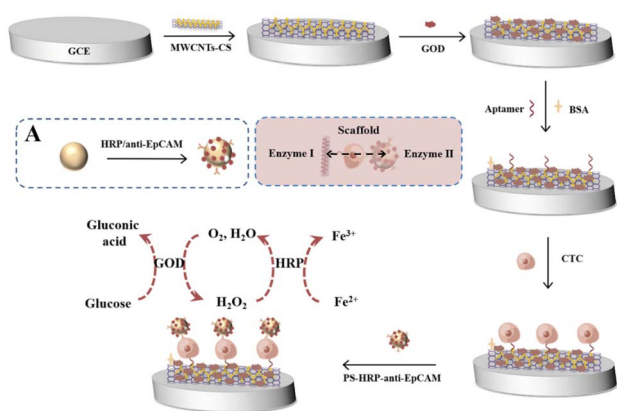
### Apparatus

Scanning electron microscopy (SEM) images were obtained using a SUPRA 55 scanning electron microscope (Carl Zeiss, Germany). Transmission electron microscopy (TEM) images were obtained using an HT7700 transmission electron microscope (Hitachi, Tokyo, Japan). All the electrochemical measurements were carried out on a CHI 760E electrochemical workstation (Shanghai Chenhua Instrument Co., Ltd., China) with a three-electrode system composed of a GCE (3 mm in diameter) as the working electrode, a platinum electrode as the counter electrode, and a saturated Ag/AgCl electrode as the reference electrode. Electrochemical impedance spectroscopy (EIS), cyclic voltammetry (CV), and differential pulse voltammetry (DPV) were performed in 5 mM  $\text{K}_3\text{Fe}(\text{CN})_6/\text{K}_4\text{Fe}(\text{CN})_6$  (0.1 M KCl) containing glucose (1 mM) as the supporting electrolyte.

### Preparation of GCE/MWCNT–CS/GOD

The GCE was polished with 0.3 and 0.05  $\mu\text{m}$   $\text{Al}_2\text{O}_3$  powder, washed with alcohol and distilled water, and then dried at room temperature. Purified MWCNTs (2 mg) were subsequently dispersed into 200  $\mu\text{L}$  CS solution ( $10 \text{ mg mL}^{-1}$ ) and sonicated for 3 h, and a mixed solution of MWCNTs–CS composite was obtained. Then, the MWCNTs–CS mixture (10  $\mu\text{L}$ ) was dropped onto the electrode to form an MWCNT–CS film by physical adsorption at room temperature.

The GOD solution ( $10 \text{ mg mL}^{-1}$ , 5  $\mu\text{L}$ ), BSA solution (2.5  $\mu\text{L}$ ), and GA solution (2.5%, v/v, 2.5  $\mu\text{L}$ ) were then added dropwise to the surface of the GCE in turn and air-dried at room temperature. The GOD was linked onto the surface of the MWCNTs–CS composite through electrostatic forces due to the positive charges of CS and the negative charges of GOD.



**Scheme 1** Schematic diagram of the carbon nanotubes–chitosan composite-driven highly efficient enzymatic cascade reaction for electrochemical biosensing detection of CTCs.



### Preparation of anti-EpCAM/PS/HRP conjugates

PS-COOH (20  $\mu\text{L}$ ) was dispersed in 1000  $\mu\text{L}$  PBS buffer solution, and then EDC (18 mg) and NHS (27 mg) were added. After sonication and subsequent incubation at 37  $^{\circ}\text{C}$  for 15 min, the carboxyl groups on the PS surface were fully activated. Then, anti-EpCAM (1  $\mu\text{g mL}^{-1}$ , 1  $\mu\text{L}$ ) and HRP (10  $\text{mg mL}^{-1}$ , 500  $\mu\text{L}$ ) were dispersed in the above PBS solution and gently stirred at 25  $^{\circ}\text{C}$  for 6 h. The PS-COOH was activated by EDC/NHS, and the HRP and anti-EpCAM were linked on the surface of PS-COOH by amide bonds. After being washed three times with 1 mL PBS solution and concentrated at 5000 rpm for 3 min, the unbound HRP and anti-EpCAM were eliminated. The obtained anti-EpCAM/PS/HRP mixture was stored at 4  $^{\circ}\text{C}$  for the next use.

### Preparation and electrochemical performance of the biosensor

The aptamer solution (10  $\mu\text{M}$ , 5  $\mu\text{L}$ ) was activated by EDC/NHS (20  $\text{mg mL}^{-1}$ ), dropped onto the surface of the GCE/MWCNTs-CS/GOD electrode, and then incubated for 20 h at room temperature. The aptamer was modified on the electrode by -COOH at the bottom of the aptamer and -NH<sub>2</sub> of CS on the GCE/MWCNTs-CS/GOD. After 1% BSA was used to block non-specific sites, the GCE/MWCNTs-CS/GOD/aptamer/BSA capture electrode was obtained. Subsequently, human breast cancer cells (MCF-7, 10  $\mu\text{L}$ ) at different concentrations were dropped and incubated for 1 h at 37  $^{\circ}\text{C}$  to capture the cells. Then, the prepared anti-EpCAM/PS/HRP (8  $\mu\text{L}$ ) was dropped onto the capture electrode (GCE/MWCNTs-CS/GOD/aptamer/MCF-7 cells), and incubated for 1 h. Thus, a heterostructured scaffold for GOD and HRP in the electrochemical biosensing system was obtained. Due to a positive relationship between the DPV value and the MCF-7 cell concentration in a 5 mM K<sub>3</sub>Fe(CN)<sub>6</sub>/K<sub>4</sub>Fe(CN)<sub>6</sub> solution (0.1 M KCl) containing glucose (1 mM) with a 50 mV pulse amplitude from -0.3 to 0.1 V, the electrochemical detection of MCF-7 cells was achieved.

### Preparation of CTCs and electrochemical detection of CTCs in whole blood samples

All experiments were performed in accordance with the guidelines of the "Ethics of Biomedical Research Involving Humans (National Health Commission of People's Republic of China)." The experiments were approved by the ethics committee of Yanshan University. Informed consent was obtained from all the human participants of this study. MCF-7 cells were digested by trypsin and resuspended in PBS buffer solution. The CTCs were quantified by hemocytometer. Then, whole blood samples were diluted ten times, and different concentrations of CTCs were dispersed into the whole blood samples for subsequent experimentation.

The whole blood samples containing different concentrations of CTCs were dropped onto the surface of the GCE/MWCNTs-CS/GOD/aptamer/BSA capture electrode and incubated for 1 h at 37  $^{\circ}\text{C}$  to capture the cells. After that, the prepared anti-EpCAM/PS/HRP (8  $\mu\text{L}$ ) was dropped onto the capture electrode (GCE/MWCNTs-CS/GOD/aptamer/MCF-7

cells) and incubated for 1 h. The electrochemical detection of MCF-7 cells in whole blood samples was conducted in a 5 mM K<sub>3</sub>Fe(CN)<sub>6</sub>/K<sub>4</sub>Fe(CN)<sub>6</sub> solution (0.1 M KCl) containing glucose (1 mM) with a 50 mV pulse amplitude from -0.3 to 0.1 V.

## Results and discussion

### Characterization of GCE/MWCNTs-CS/GOD

As shown in Fig. 1A, the MWCNTs are uniform tubular structures with diameters and lengths of approximately 10 nm and 10  $\mu\text{m}$ , respectively. In the SEM image of the MWCNTs-CS composite (Fig. 1B), the MWCNTs were uniformly coated by CS. As shown in Fig. S1,<sup>†</sup> elemental mapping of the MWCNTs-CS composite was performed using energy-dispersive X-ray spectroscopy (EDX) to verify the uniform distribution of O and N, which also revealed the CS dispersed on the carbon nanotube surface. SEM of the further modification of GOD on the MWCNTs-CS electrode surface is shown in Fig. 1C, in which GOD is seen as dewdrops scattered on the branches of trees. UV-Vis absorption spectra are exhibited in Fig. 1D, and no absorption peak was observed for CS (curve a), MWCNTs (curve b), or the MWCNTs-CS composite (curve d). However, a 280 nm absorption peak appeared for MWCNT-CS/GOD (curve e), which corresponded to the typical protein absorption peak of GOD (curve c).<sup>44,45</sup> These results indicated that GOD was successfully loaded onto the MWCNTs-CS composite.

The electrochemical effective surface area of the electrode would affect the GOD loading rate, which would further affect the catalytic efficiency of the enzymatic cascade reactions. A series of CV experiments was carried out for investigation of the electrochemical effective surface area of the GCE and GCE/MWCNTs-CS, as shown in Fig. 2. The electrochemical effective surface area of the electrode, unlike the physical surface area, was investigated by CV measurements using K<sub>3</sub>Fe(CN)<sub>6</sub>/K<sub>4</sub>Fe(CN)<sub>6</sub> as a probe and the following Randles-Sevcik equation:<sup>46</sup>

$$I_p = kn^{3/2}cAD^{1/2}\nu^{1/2}$$

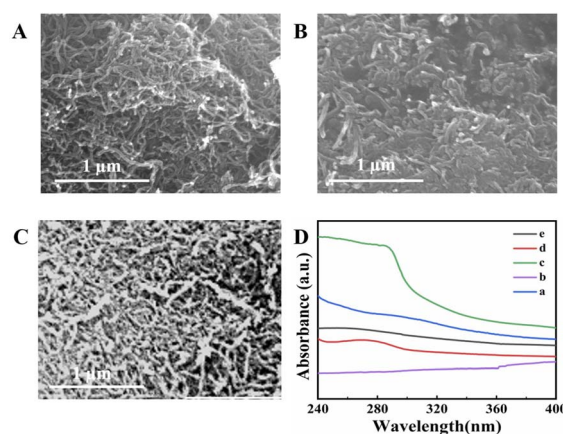


Fig. 1 SEM images of (A) MWCNTs, (B) MWCNTs-CS composite, and (C) MWCNTs-CS/GOD. (D) UV-Vis spectra of (a) CS, (b) MWCNTs, (c) GOD, (d) MWCNTs-CS-GOD, and (e) MWCNTs-CS composite.





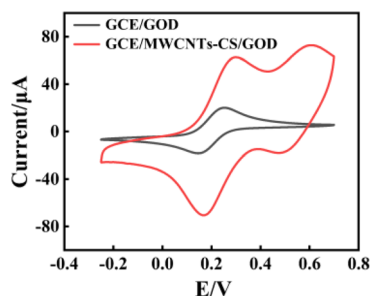


Fig. 2 CV results for GCE/GOD and GCE/MWCNTs-CS/GOD in 0.1 M KCl solution containing 5 mM  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (scan rate of  $100\text{ mV s}^{-1}$ ).

where  $k = 2.69 \times 10^5\text{ C (mol V}^{1/2})^{-1}$ ,  $n$  denotes the number of electron transfers,  $A$  denotes the electrochemical surface area,  $D$  denotes the diffusion coefficient ( $7.6 \times 10^{-7}\text{ cm}^2\text{ s}^{-1}$ ) of  $K_3Fe(CN)_6/K_4Fe(CN)_6$ ,  $\nu$  denotes the scanning rate ( $\text{V s}^{-1}$ ), and  $c$  denotes the concentration of the solution ( $\text{mol m}^{-3}$ ). The electrochemical effective surface area of the GCE and GCE/MWCNTs-CS was  $0.0539\text{ cm}^2$  and  $0.1688\text{ cm}^2$ , respectively. Comparing the GCE, we found that the electrochemical effective surface area of the GCE/MWCNTs-CS increased approximately 3-fold, which would increase the loading rate of the GOD and thus increase the electrochemical response signal.

### Characterization of the anti-EpCAM/PS/HRP conjugates

Carboxylated polystyrene microspheres (PS-COOH) with a large surface area were used as carriers for HRP and anti-EpCAM. HRP and anti-EpCAM were linked by the amino groups in anti-EpCAM and the carboxyl groups in PS. The SEM image of PS-COOH showed that they exhibited a smooth and uniform surface with a diameter of  $1\text{ }\mu\text{m}$  (Fig. 3A). After being covered by HRP and anti-EpCAM, some particles and synapses appeared on the PS surface (Fig. 3B). The FT-IR spectra of PS and anti-EpCAM/PS/HRP are illustrated in Fig. 3C. The absorption peaks of HRP at  $1600\text{ cm}^{-1}$  were attributed to amide groups,

and the absorption peaks at  $1500\text{ cm}^{-1}$  and  $1300\text{ cm}^{-1}$  were assigned to the heme vibration and tertiary amide, respectively (Fig. 3C, curve b). The absorption peaks of anti-EpCAM/PS/HRP at  $1654\text{ cm}^{-1}$  and  $3367\text{ cm}^{-1}$  were attributed to the bending vibrations of  $-\text{CONH}$  and  $-\text{NH}$ , respectively (Fig. 3C, curve c).

These results indicated that anti-EpCAM and HRP had connected on the surface of PS. The zeta potentials of PS-COOH, anti-EpCAM/PS, and anti-EpCAM/PS/HRP were investigated, as shown in Fig. 3D. The zeta potential of PS-COOH was  $-50.7\text{ mV}$  due to the negative charge of  $-\text{COOH}$ . In comparison, the zeta potentials of anti-EpCAM/PS and anti-EpCAM/PS/HRP were  $-52.5\text{ mV}$  and  $-63.3\text{ mV}$ , respectively, after being modified with negatively charged anti-EpCAM and HRP. These results indicated that anti-EpCAM and HRP were successfully loaded on PS-COOH.

### Characterization of the 'heterostructured' scaffold for GOD and HRP

Electrochemical impedance spectroscopy (EIS) and CV were used to study the assembly progress of the heterostructured scaffold for GOD and HRP, as described in Fig. 4. The EIS consisted of two parts, a semicircular part and a linear part. The semicircular diameter of the EIS in the high-frequency region corresponded to the electron transport resistance ( $R_{et}$ ), and the linear counterpart in the low-frequency region was related to the diffusion process. The bare GCE showed a small  $R_{et}$  (Fig. 4A, curve a) and an obvious current peak (Fig. 4B, curve a). The MWCNTs can promote electron transfer to the interface to a large extent due to their satisfactory conductivity, and therefore, when compared with the bare GCE, when the components of the MWCNTs-CS composite were modified on the GCE surface, a lower  $R_{et}$  was exhibited in the EIS of the GCE/MWCNTs-CS (Fig. 4A, curve b), and the CV peak current was significantly increased (Fig. 4B, curve b).

However, electron transfer was hindered when GOD, as a poor conductor, was wrapped in the GCE/MWCNTs-CS electrode surface, resulting in an increase in the  $R_{et}$  (Fig. 4A, curve c)

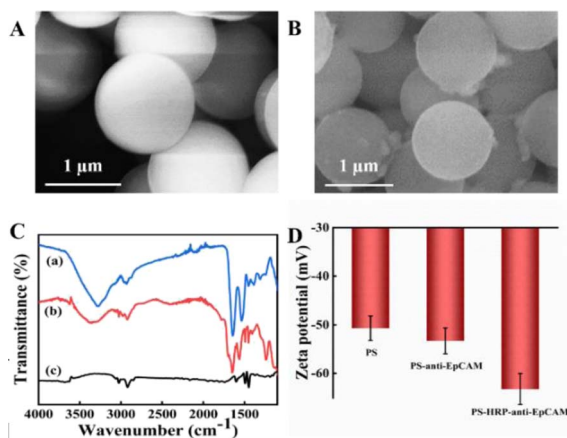


Fig. 3 SEM images of (A) PS and (B) anti-EpCAM/PS/HRP (diameter is  $1\text{ }\mu\text{m}$ ). (C) FT-IR spectra of (a) HRP, (b) anti-EpCAM/PS/HRP, and (c) PS. (D) Zeta potentials of PS, anti-EpCAM/PS, and anti-EpCAM/PS/HRP.

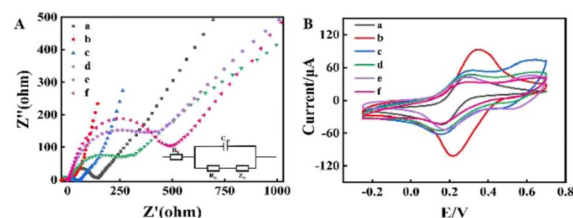


Fig. 4 (A) Electrochemical impedance spectroscopy of (a) bare GCE, (b) GCE/MWCNTs-CS, (c) GCE/MWCNTs-CS/GOD, (d) GCE/MWCNTs-CS/GOD/aptamer, (e) GCE/MWCNTs-CS/GOD/aptamer/CTCs, and (f) GCE/MWCNTs-CS/GOD/aptamer/CTCs/anti-EpCAM/PS/HRP in 0.1 M KCl solution containing 5 mM  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (impedance spectral frequency of  $0.1\text{--}10^5\text{ Hz}$ , amplitude of  $5\text{ mV}$ ). (B) Cyclic voltammograms of (a) bare GCE, (b) GCE/MWCNTs-CS, (c) GCE/MWCNTs-CS/GOD, (d) GCE/MWCNTs-CS/GOD/aptamer, (e) GCE/MWCNTs-CS/GOD/aptamer/CTCs, and (f) GCE/MWCNTs-CS/GOD/aptamer/CTCs/anti-EpCAM/PS/HRP in 0.1 M KCl solution containing 5 mM  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (scan rate of  $100\text{ mV s}^{-1}$ ).

and a decrease in the peak current (Fig. 4B, curve c). Then, the immobilization of aptamer onto GCE/MWCNTs-CS/GOD *via* the terminal carboxyl group and the amino group of CS after activation by EDC and NHS resulted in an increase in the impedance value (Fig. 4A, curve d), indicating that electron transfer was further blocked in the modified electrode. Similarly, the peak current also decreased (Fig. 4B, curve d). Subsequently, when CTCs were captured on the surface of GCE/MWCNTs-CS/GOD/aptamer, the impedance of the system led to a continual increase (Fig. 4A, curve e), and the CV peak current decreased (Fig. 4B, curve e). The isolation between anti-EpCAM/PS/HRP conjugates and the electrode interface resulted in a significant increase in  $R_{et}$  (Fig. 4A, curve f) due to the effect of steric hindrance. Additionally, the CV peak current further decreased as expected (Fig. 4B, curve f), which also indicated that fabrication of the electrochemical biosensor for CTCs was feasible.

### Detection performance of the electrochemical biosensor for CTCs

Glucose in the electrolyte was initially oxidized by GOD to produce gluconic acid and  $H_2O_2$ . Then, the  $H_2O_2$  was further utilized by HRP to accelerate the oxidation of  $Fe^{2+}$  with amplified electrochemical responses. The mechanism was proved by the DPV responses of the biosensor in the absence and presence of glucose (1 mM) and  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (5 mM) in the PBS solution (Fig. S2†). The DPV response clearly increased in the presence of glucose (1 mM) in the PBS solution containing  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (5 mM). This occurred because GOD had catalyzed glucose in the electrolyte to produce gluconic acid and  $H_2O_2$ . Then, the  $H_2O_2$  was further utilized by HRP to accelerate the oxidation of  $Fe^{2+}$  with amplified electrochemical responses. Moreover, compared to the absence of  $K_3Fe(CN)_6/K_4Fe(CN)_6$ , the DPV response clearly increased in the presence of  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (5 mM) in the PBS solution containing glucose (1 mM) (Fig. S3†). This further indicated that the enzymatic cascade reaction accelerated the oxidation of  $Fe^{2+}$ , with amplified electrochemical responses. In addition, some experimental conditions were optimized to obtain the excellent performance of electrochemical biosensing detection, including the concentration of MWCNTs-CS composite, the enrichment time for DNA, the concentration of HRP, and the concentration of glucose (Fig. S4†).

The change in the electrical signal resulting from the cascade reactions of GOD and HRP was proportional to the number of CTCs captured on the electrode. The DPV experiment was performed to verify the signal amplification strategy based on a dual enzymatic cascade reaction for the detection of CTCs. As shown in Fig. 5, compared with the GCE and GCE/MWCNTs-CS/GOD, the DPV peak currents in 5 mM  $K_3Fe(CN)_6/K_4Fe(CN)_6$  (0.1 M KCl) containing glucose (1 mM) significantly increased with the increase in the CTC concentration with a 50 mV pulse amplitude from  $-0.3$  to  $0.1$  V. The intensity of DPV was increased with the cell concentration (Fig. 5A), and the intensity of DPV was linear in the logarithm of cell concentration ranging from  $10$  cells  $mL^{-1}$  to  $10^6$  cells  $mL^{-1}$ , as shown in

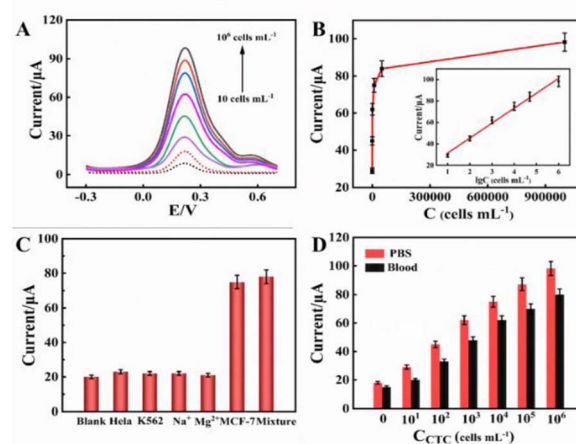


Fig. 5 (A) DPV was obtained with different concentrations of CTCs (the solid lines indicate the cells numbers in the range of  $10^1$  to  $10^6$  cells  $mL^{-1}$ , and the dotted lines denote the GCE (blank line) and GCE/MWCNTs-CS/GOD (red line)). (B) Relationship between the peak current of DPV and the concentration value of CTCs. (C) Peak current in response to various types of cells at a concentration of  $10^4$  cells  $mL^{-1}$ . (D) The relationship between the peak current of DPV and CTC concentration in PBS and whole blood.

Fig. 5B. The linear regression equation was  $y = 13.941 \lg C \text{ cells (cells } mL^{-1}) + 17.438$  with a correlation coefficient of 0.9927 ( $n = 3$ ). The detection limit was calculated to 3 cells  $mL^{-1}$  for CTCs at  $3\sigma$ .

The reproducibility of the electrochemical biosensor was evaluated by measuring the electrochemical response at multiple cell concentrations. The relative standard deviations (RSDs) were 1.45%, 2.25%, 3.1%, 3.75%, 4.2%, and 4.91% at cell concentrations of  $10$  cells  $mL^{-1}$ ,  $10^2$  cells  $mL^{-1}$ ,  $10^3$  cells  $mL^{-1}$ ,  $10^4$  cells  $mL^{-1}$ ,  $10^5$  cells  $mL^{-1}$ , and  $10^6$  cells  $mL^{-1}$ , respectively (the detailed RSD values are listed in Table S1†), which indicated satisfactory repeatability.

To investigate the specificity of the designed electrochemical biosensor, a series of experiments was performed using various types of cells, including HeLa cells, K562 cells, MCF-7 cells,  $Na^+$ , and  $Mg^{2+}$ , as shown in Fig. 5C. Only MCF-7 cells elicited an obvious electrochemical response, which indicated that this electrochemical biosensor exhibited satisfactory selectivity for MCF-7 cells based on the strategy of a dual enzymatic cascade reaction.

Furthermore, we performed whole blood experiments to investigate the potential clinical application of this sensing system. As shown in Fig. 5D, the comparison graphs of PBS and 10-fold diluted whole blood samples indicate that this biosensor exhibited strong resistance to interference, and the relative standard deviations of whole blood samples were 1%, 1.65%, 2.4%, 3.1%, 3.5%, and 4% at cell concentrations of  $10$  cells  $mL^{-1}$ ,  $10^2$  cells  $mL^{-1}$ ,  $10^3$  cells  $mL^{-1}$ ,  $10^4$  cells  $mL^{-1}$ ,  $10^5$  cells  $mL^{-1}$ , and  $10^6$  cells  $mL^{-1}$ , respectively (the detailed RSD values are listed in Table S2†). Moreover, the regression analysis of DPV current *versus* the cell concentration in PBS and whole blood in Fig. S5† showed a linear relationship in the range of  $10$  cells  $mL^{-1}$  to  $10^6$  cells  $mL^{-1}$ . The linear regression equation was



$y = 13.941 \lg C \text{ cells (cells mL}^{-1}) + 17.438$  with a correlation coefficient of 0.9927 and  $y = 5.2736 \lg C \text{ cells (cells mL}^{-1}) + 9.6667$  with a correlation coefficient ( $R^2$ ) of 0.9891. All the results indicated that the MWCNT-CS composite was able to drive the highly efficient enzymatic cascade reaction, and it holds promise for clinical applications of electrochemical biosensing detection of CTCs.

## Conclusions

An MWCNTs-CS composite-driven highly efficient enzymatic cascade reaction for electrochemical biosensing detection of CTCs was developed. In this strategy, MWCNTs-CS composites were adopted to modify a GCE for loading GOD, which achieved a high electrochemical effective surface area of the electrode, high loading rate, and high electrical activity of the enzyme. As a 'power source', the MWCNTs-CS composites provided strong driving power for the subsequent enzymatic cascade reactions. Aptamer, CTCs, and PS microspheres with HRP and anti-EpCAM were assembled on MWCNTs-CS/GCE. GOD catalyzed the oxidation of glucose to produce gluconic acid and  $\text{H}_2\text{O}_2$ , while HRP catalyzed the decomposition of  $\text{H}_2\text{O}_2$ , allowing the reaction to proceed and further facilitating electron transfer. This resulted in a highly efficient enzymatic cascade reaction for electrochemical biosensing detection of CTCs that exhibited a wide linear concentration range ( $10 \text{ cells mL}^{-1}$  to  $10^6 \text{ cells mL}^{-1}$ ) with high sensitivity, selectivity, reproducibility, and detection limit ( $3 \text{ cells mL}^{-1}$  for CTCs at  $3\sigma$ ).

Based on these advantages, great potential was indicated for the MWCNT-CS composite-driven highly efficient enzymatic cascade strategy for clinical electrochemical biosensing detection of CTCs. However, the stability of natural enzymes is easily affected by temperature, pH, and salt concentrations, which also affects the catalytic efficiency of the enzymatic cascade reaction in the electrochemical biosensors. Nanozymes have been discovered with high catalytic efficiency, selectivity, and stability, and they have been developed as potentially viable alternatives to natural enzymes. In our next study, we will use nanozymes to replace the natural enzymes in the enzymatic cascade reaction.

## Author contributions

Min Dong: conceptualization, methodology, data curation, writing – original draft preparation. Zhihong Gao: visualization, investigation. Jian Li, Yating Zhang: supervision, investigation. Jiahui Cai: investigation. Hong Jiang, Panpan Xu: software, validation. Jidong Wang: supervision, funding acquisition, writing – review and editing.

## Conflicts of interest

There are no conflicts to declare.

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