



Ultrasensitive electrochemical biosensors based on zinc sulfide/graphene hybrid for rapid detection of SARS-CoV-2

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Abstract

The coronavirus disease 2019 (COVID-19) is a highly contagious and fatal disease caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). In general, the diagnostic tests for COVID-19 are based on the detection of nucleic acid, antibodies, and protein. Among different analytes, the gold standard of the COVID-19 test is the viral nucleic acid detection performed by the quantitative reverse transcription polymerase chain reaction (qRT-PCR) method. However, the gold standard test is time-consuming and requires expensive instrumentation, as well as trained personnel. Herein, we report an ultrasensitive electrochemical biosensor based on zinc sulfide/graphene (ZnS/graphene) nanocomposite for rapid and direct nucleic acid detection of SARS-CoV-2. We demonstrated a simple one-step route for manufacturing ZnS/graphene by employing an ultrafast (90 s) microwave-based non-equilibrium heating approach. The biosensor assay involves the hybridization of target DNA or RNA samples with probes that are immersed into a redox active electrolyte, which are detectable by electrochemical measurements. In this study, we have performed the tests for synthetic DNA samples and, SARS-CoV-2 standard samples. Experimental results revealed that the proposed biosensor could detect low concentrations of all different SARS-CoV-2 samples, using such as S, ORF 1a, and ORF 1b gene sequences as targets. This microwave-synthesized ZnS/graphene-based biosensor could be reliably used as an on-site, real-time, and rapid diagnostic test for COVID-19.

Keywords DNA biosensor · COVID-19 · Microwave · Graphene · Zinc sulfide · Nanocomposite

1 Introduction

The coronavirus disease 2019 (COVID-19) has been striking the world in the past 2 years [1–5]. People who got infected will have various symptoms including fever, cough, headache, loss of smell, loss of taste, fatigue, and breathing difficulties [6–8]. The major diagnostic methods to detect the

disease include computed tomography (CT) scan, nucleic acid amplification test (NAAT), and serological techniques [9]. Today, NAAT using the quantitative reverse transcription polymerase chain reaction (qRT-PCR) is the most widely used method for detecting the COVID-19. However, the qRT-PCR is a time-consuming and expensive test which also requires trained personnel. Therefore, the development of highly sensitive and quick detection methods is urgently needed. And the electrochemical DNA biosensors have received a lot of attraction for being a good detecting device [10–13].

Biosensor is an analytical device detecting biological or chemical reactions and then generating reliable data for several fields such as gene analysis [12, 14], food safety [15, 16], disease diagnosis [17], environmental monitoring [18], drug discovery [19], and water quality screening [20]. A typical biosensor consists of biological receptor, transducer, amplification, and signal processing. Based on different transducers, biosensors can be divided into electrochemical, optical, thermal, electronic, acoustic, and

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mass-based biosensor [21]. Among different transducers, the electrochemical biosensor is the most widely studied and applied biosensor due to the advantages of low-cost, simplicity, portability, sensitivity, and rapid detection [14, 22]. It is critical to develop an appropriate composite for the signal amplification platform.

In practice, materials such as metals, metal oxides, graphene, conducting polymer, and their composite have been studied as signal amplifier for the electrochemical biosensors [23, 24]. For instance, Saber et al. fabricated a gold electrode decorated by gold nanorods (GNRs) for detecting hepatitis B virus [25]. Due to the large surface area of the GNRs, the modified gold electrode can assemble more probe DNA and exhibit higher sensitivity with a detection limit of 2×10^{-12} M. Similarly, a hybrid nanocomposite containing copper oxide nanowires and carboxyl-functionalized single-walled carbon nanotubes was developed by Chen et al. for detecting the target DNA [26]. The performance of the modified electrode was benefited by the nanocomposite which provides a large surface area, fast electron transfer, and good electrical conductivity with the limit detection of 3.5×10^{-15} M. Graphene also shows a great potential in DNA detection due to its astonishing properties in electrical, optical, mechanical, and chemical aspects [27]. Pumera et al. reported a graphene-based electrode immobilized with hairpin-shaped probe DNA for detecting Alzheimer's disease [28]. The triple and quadruple layer graphene electrodes presented the best sensitivity with the detection limit of 6.6×10^{-12} M. All cases above show that the modified electrode provides a larger surface area and low charge transfer resistance for superior sensitivity compared to the raw electrode.

In the detection of COVID-19, the antibody testing, also known as serology testing, can check the antibodies to COVID-19 after a person has been infected or vaccinated. For instance, a paper-based biosensor modified by ZnO nanowires showed enhanced sensitivity for detecting antibody of COVID-19 with human serum samples [29]. In addition, Seo et al. reported a graphene-based biosensor functionalized with SARS-CoV-2 spike antibody as a receptor which demonstrated highly sensing detection for the target SARS-CoV-2 antigen protein with a detection limit of 1×10^{-15} g/mL [30]. On the other hand, the molecular test detects the virus's genetic material to determine if a person has an active infection of COVID-19. Song et al. utilized polyaniline nanowires to modify the glassy carbon electrode for the detection of the COVID-19 N gene, resulting in a detection limit of 3.5×10^{-15} M [31]. This demonstrated that the electrochemical biosensor has the potential as a detection method for COVID-19. In many cases, the process of making the composite takes multiple steps with long preparation time and high cost. Therefore, developing a quick and one-step process is in need.

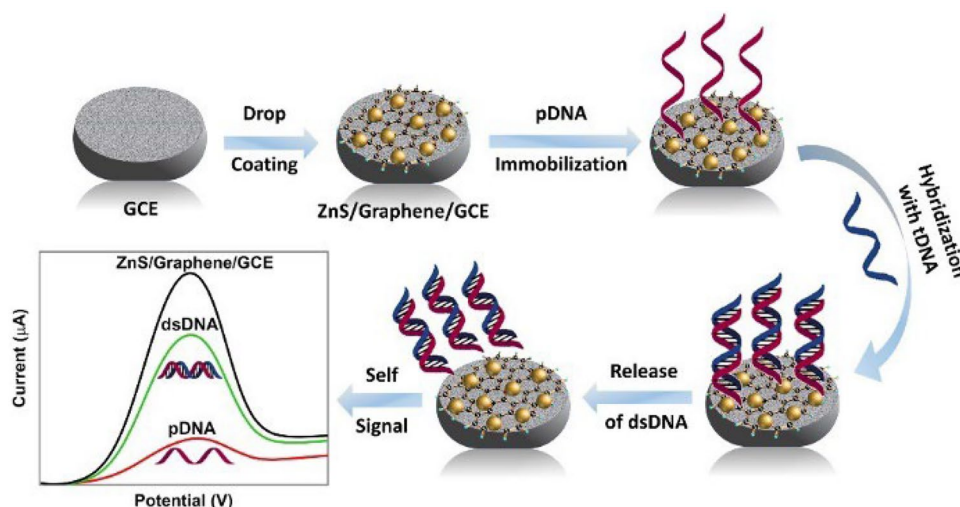
In this paper, we present a one-step and ultrafast microwave-initiated manufacturing process to prepare the electrical interface containing zinc sulfate and graphene (ZnS/graphene). ZnS has attracted a lot of interest due to their superior chemical, physical, and electrochemical properties [32, 33]. For example, a system with urease and CdSe/ZnS quantum dots (QDs) had been developed for the detection of urea by Huang et al. [34]. By the enhancement of QD photoluminescence, the novel system can analyze the urea concentration ranging from 0.01 to 100 mM with good sensitivity, and no enzyme immobilization is required. In addition, Rajesh et al. studied the electrical and DNA sensing properties of ZnS decorated single-walled carbon nanotube (SWNT) to investigate the sensing mechanism by measuring the transfer characteristics of I-V and field-effect transistors (FET) [35]. The result demonstrated that the improvement of the sensitivity is attributed to the large surface-to-volume ratio and the superior charge transfer ability of the ZnS. The graphene which accelerates the reaction rate by absorbing the electromagnetic energy acts as a heat resource during the microwave preparation. Besides, graphene has been applied for different biosensors due to its superior sensing performance including good conductivity, large surface area, excellent thermal conductivity, and mechanical strength [36–39]. Compared with the conventional heating method under thermodynamic equilibrium conditions, the microwave reaction method under non-equilibrium conditions can provide low-cost and time-saving process [40–42]. Most importantly, the ZnS/graphene DNA biosensor provides highly sensitivity. Different ratios of zinc precursors were investigated to obtain the optimized conditions and compared with commercial ZnS mixed with graphene without microwave reaction. The obtained ZnS/graphene sample was drop cast onto a glassy carbon electrode (GCE) surface for further electrochemical experiment. The probe DNA (pDNA) was successfully immobilized with GCE and then hybridized with different concentrations of target DNA (tDNA) to perform self-signal regeneration during the releasing process (Scheme 1). Three-electrode system was utilized while the ZnS/graphene modified electrode acted as the working electrode. The performance of the modified electrode was characterized by cyclic voltammetry (CV) and differential pulse voltammetry (DPV).

2 Methodology

2.1 Materials

Zinc acetate (GR, 99.5% purity) was obtained from Bean-Town Chemical Inc. Graphene substrate was obtained from Magnolia Ridge Inc. Sulfur powder (GR, 99.5% purity) was provided by Alfa Aesar, America. The detection samples

Scheme 1 Schematic illustration of detecting the SARS-CoV-2 by microwave-synthesized ZnS/graphene nanocomposite



included artificial DNA probes, artificial DNA targets, and SARS-CoV-2 standard were obtained from Eurofins Genomics LLC (Louisville, Kentucky).

2.2 Preparation of ZnS/graphene composites

For the preparation of ZnS/graphene nanocomposites, $\text{ZnAc}_2 \cdot 2\text{H}_2\text{O}$, graphene, and sulfur powder were transferred into a 20-mL scintillation vial and mixed thoroughly to get a homogeneous mixture. The precursor ratios and synthesis parameters are described in Table S1. Subsequently, the vial was placed inside microwave oven and irradiated with a high-power level (~ 1250 W) for 90 s. After the vial was cooled down to room temperature, the obtained powder was taken out. For comparison, the mixture of graphene and commercial ZnS nanoparticles were also prepared under the same microwave experimental conditions.

The surface of glassy carbon electrode (GCE) was sequentially polished with 0.3- and 0.05-mm alumina slurries, then immersed in Piranha solution (conc. H_2SO_4 :30% H_2O_2 ; 3:1) for 30 min, followed by rinsing with DI water, and vacuum dried at 60 °C. To prepare the coating paste, 100 mg of hybrid powder was suspended in 5 mL of isopropyl alcohol. Then, 0.1 mL of Nafion was added into the suspension following 20 min of sonication. Finally, 20 μL of the suspension was added to the prepared GCE surface and vacuum dried at 60 °C.

2.3 Nanocomposite-based DNA biosensor probe and target sequences

The entire genetic sequence of SARS-CoV-2 was uploaded to the Global Initiative on Sharing All Influenza Data (GISAID) platform on January 10, 2020. The sequence data was used as reference for the design of oligos/probes needed for the development of SARS-CoV-2-specific testing. As

shown in Table 1, the probe and target DNA oligonucleotides were complementary to the specific target gene sequences and purchased from Eurofin Genomics.

2.4 Immobilization and hybridization of DNA

A 20 μL of probe-DNA solution (1.0×10^{-6} M pDNA) was added dropwise to the surface of modified GCE and dried at temperature below the melting T_m (temperature of melting) of corresponding probe sequences. After pDNA immobilization, following by rinsing with DI water, the GCE was dried again. Then, 20 μL of target DNA (tDNA) was added onto the pDNA-modified GCE surface, and the corresponding electrode was dried at room temperature to form the hybrid double-stranded DNA (dsDNA). Subsequently, the electrode was kept at -0.5 V for 300 s in 1.0 M KCl solution containing 0.2 M $\text{K}_3[\text{Fe}(\text{CN})_6]$ to release the dsDNA into the electrolyte. After rinsing the surface of GCE with DI water, the electrode was ready for electrochemical tests.

2.5 Material characterization

The surface morphology of the ZnS/graphene nanocomposite was characterized using scanning electron microscope (SEM) on JEOL JSM-7000F coupled with an energy dispersive X-ray spectrometer (EDS, EDAX Instruments).

Table 1 List of probe and target DNA sequences

Gene (s)	Synthetic base-pairs of DNA templates
S	Probe GACATGTATAGCATGGAACC Target GGTTCATGCTATACATGTCTC
ORF 1b	Probe CTTTAATGTTTATTCTCTA Target TAGAGAATAAAACATTAAAG
Mis-matched negative control	Probe TTCACAAGTGCCGTGCCTAC Target GGGTACTTAACAATGATTATT

The crystalline phase characteristics of the microwave samples were examined using the X-ray diffraction (XRD) (Philips X'pert MPD diffractometer) under Cu K α radiation ($\lambda = 1.5405 \text{ \AA}$) at 2θ range of $20 - 70^\circ$.

2.6 Electrochemical measurements

All the electrochemical tests were conducted in a mixture of 1.0 M KCl and 0.2 M K₃[Fe(CN)₆] at room temperature. Cyclic voltammetry (CV) experiments were carried out at a scan rate of 100 mV s^{-1} from -0.4 to 0.8 V for different ZnS/graphene composites. The differential pulse voltammetry (DPV) tests were taken by using parameters as follows: pulse amplitude, 0.05 V; pulse width, 0.05 s; and pulse period, 0.5 s. The peak current difference (ΔI_p) was calculated according to the relation of $\Delta I_p = I - I_0$, where I_0 represents the peak current for the modified electrodes immobilized with pDNA and I represents the peak current for the resulting dsDNA after being released into the electrolyte. The calibration plot, ΔI_p vs. $-\log C$ was drawn by calculating the negative logarithmic values of target samples concentrations, where C is the concentration of target sample. Furthermore, the LOD (limit of detection) is measured using the following equation:

$$\text{LOD} = \frac{3 \times \text{Standard deviation of blank solution}, \sigma}{\text{Slope of the calibration plot}} \quad (1)$$

In Eq. (1), the standard deviation of blank solution was measured by running the DPV tests on pDNA-modified GCE for 10 times.

3 Results and discussion

3.1 Optimizing the precursor ratio

The electrochemical properties of different modified electrodes with different mass of precursors were investigated

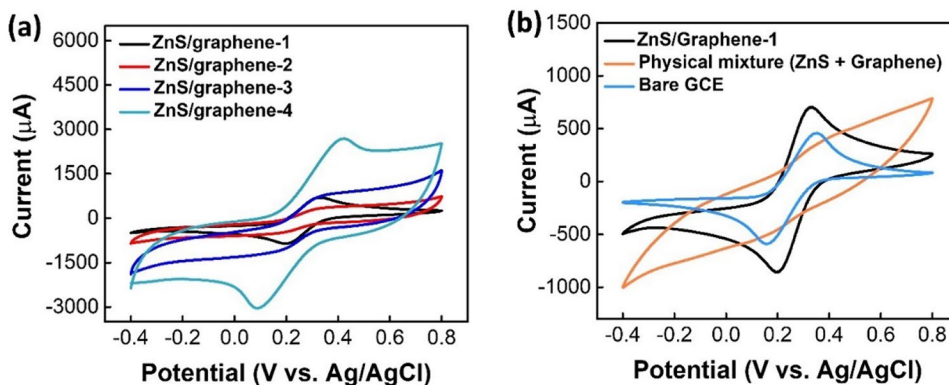
by CV technique. Among the four different mass ratios (see Table S1), as displayed in Fig. 1a, ZnS/graphene-1 has the prominent redox peaks with the anodic peak to cathodic peak separation of potential, ΔE_p ($\Delta E_p = E_{\text{anodic peak}} - E_{\text{cathodic peak}}$) of 130 mV. For the samples of ZnS/graphene-2 and ZnS/graphene-3, the redox peaks are not prominent and may be due to the incomplete reaction taking place during the microwave-assisted synthesis with the less amount of Zn-precursor. In addition, although ZnS/graphene-4 provides much higher current due to the presence of high amount of conductive graphene, the ΔE_p is as much as 330 mV, which indicates a sluggish kinetic transfer process of electrons.

Moreover, the electrochemical property of ZnS/graphene-1 was compared with a physical mixture of ZnS and graphene with a mass ratio of 6:1, and with the bare GCE. From Fig. 1b, it can be clearly observed that the physical mixture does not show prominent redox peaks. For ZnS/graphene-1 and bare GCE, ΔE_p values are 130 mV and 190 mV, respectively. Moreover, the ZnS/graphene-1 yields the maximum peak current, possibly because of the accelerated $[\text{Fe}(\text{CN})_6]^{3-}$ diffusion towards the surface by a layer of graphene. Apparently, the ZnS/graphene-1 sample exhibited much higher electrochemical activity and faster electron-transfer kinetics (reflected in the lower peak to peak separation) compared to all other modified electrodes including bare GCE. Therefore, for further studies, the detection of DNA samples was carried out on the surface of ZnS/graphene-1 modified GCE which was denoted as ZnS/graphene/GCE thereafter.

3.2 Characterization of the modified electrode

The morphology and distribution of the ZnS/graphene composite are shown in Fig. 2a–e. The SEM image was utilized to characterize the prepared ZnS/graphene composite (Fig. 2a). It can clearly be seen that the graphene substrate was fully decorated by ZnS. The EDS spectrum of ZnS/graphene exhibited the existence of the Zn and S elements. Figure 2b–e illustrated the EDS elemental mapping of ZnS/

Fig. 1 **a** CVs of different electrodes in a solution containing 1.0 M KCl and 0.2 M K₃[Fe(CN)₆] as a redox probe at scan rate of 100 mV s^{-1} . Zn(Ac)₂·2H₂O, graphene, and S precursors have a mass ratio of 6:1:1, 4:1:1, 2:1:1, and 6:2:1, respectively. **b** CV curves of different electrodes in a solution containing 1.0 M KCl and 0.2 M K₃[Fe(CN)₆]



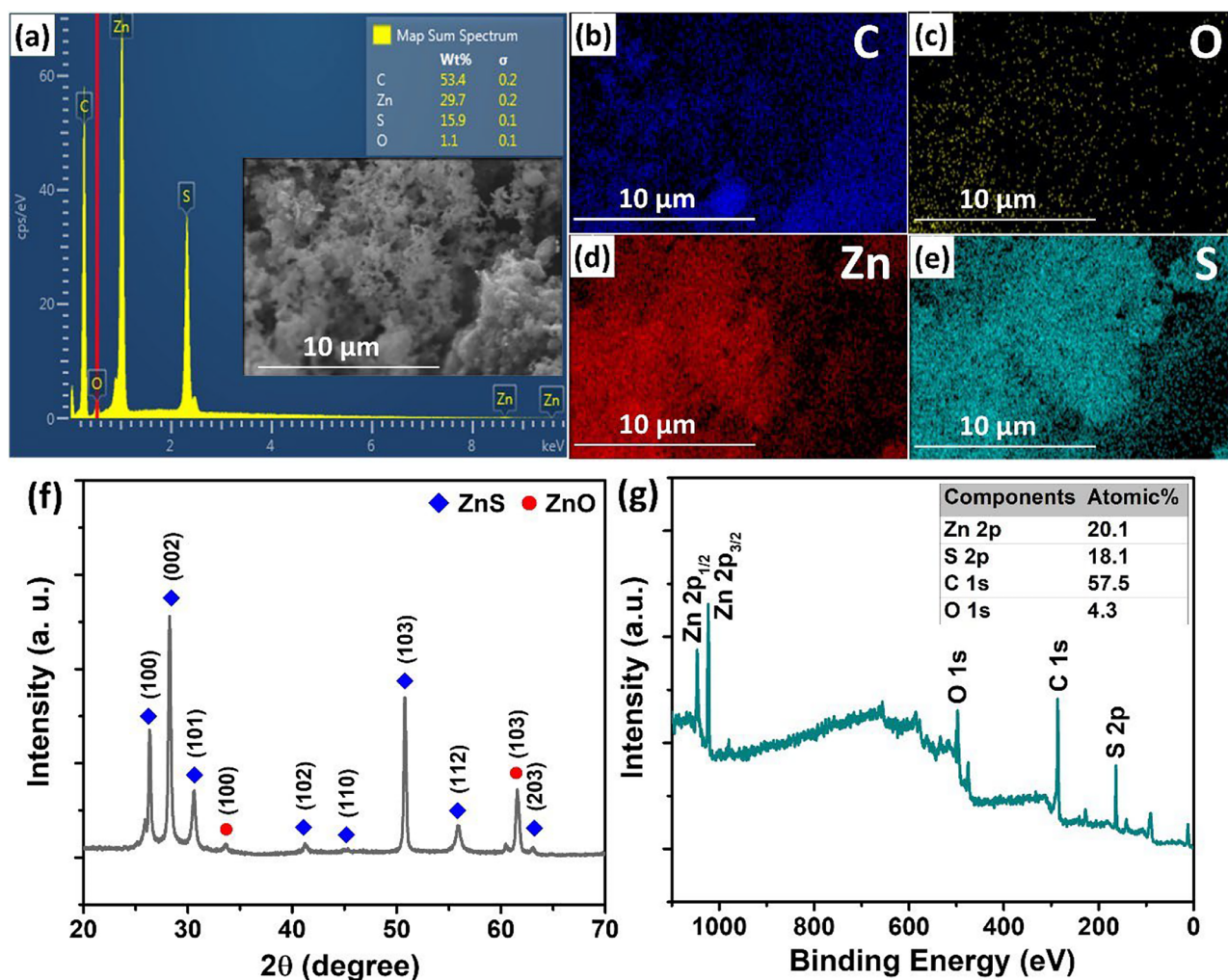


Fig. 2 a SEM image and EDS spectrum; b – e the EDS elemental mapping; f XRD pattern and g XPS spectrum of ZnS/graphene composite

graphene composite which clearly showed the evenly dispersed elements of Zn and S on the graphene surface. In addition, the C, Zn, and S components were closely stacked, indicating the formation of the hybrid composite. The XRD diffractogram of ZnS/graphene is shown in Fig. 2f. The strong peaks appeared at 2θ around 26.68° , 28.56° , 30.86° , and 51.1° are assigned to (100), (002), (101), and (103) reflections of hexagonal wurtzite ZnS (JCPDS 36–1450), respectively. The overall XPS spectrum for the ZnS/graphene composite is shown in Fig. 2g. The spectrum showed the presence of C 1s, O 1s, Zn 2p, and S 2p peaks. The characterized results confirmed the successful synthesis of ZnS/graphene composite. Since microwave-initiated synthesis does not have any inert gas protection during the heating process, the EDS spectrum showed the presence of ZnO. However, the ZnO does not contribute much to DNA detection. The CV and DPV comparison between ZnO and ZnS have been discussed and shown in Fig. S1.

3.3 Detection of synthetic target DNA samples

To explore the analytical characteristics of the optimized ZnS/graphene/GCE for the detection of synthetic gene DNA sequences, DPVs were performed in an electrolyte solution of 1.0 M KCl containing 0.2 M $K_3[Fe(CN)_6]$. Initially, the sensor electrode of ZnS/graphene/GCE was attached with the 1.0×10^{-6} M probe DNA (pDNA) and treated for different concentrations of target DNA (tDNA) solutions as depicted in Fig. 3a. After immobilizing the pDNA on the surface of ZnS/graphene/GCE, the oxidation or anodic peak current (I_{pa}) decreases significantly because the non-conductive pDNA blocks the effective electron transfer through the coating surface. However, when the pDNA was hybridized with the tDNA, the peak currents were enhanced linearly with the enhancing concentrations of tDNA from 1.0×10^{-18} M to 1.0×10^{-7} M. The release of the resulted dsDNA from the ZnS/graphene/GCE surface occurs because

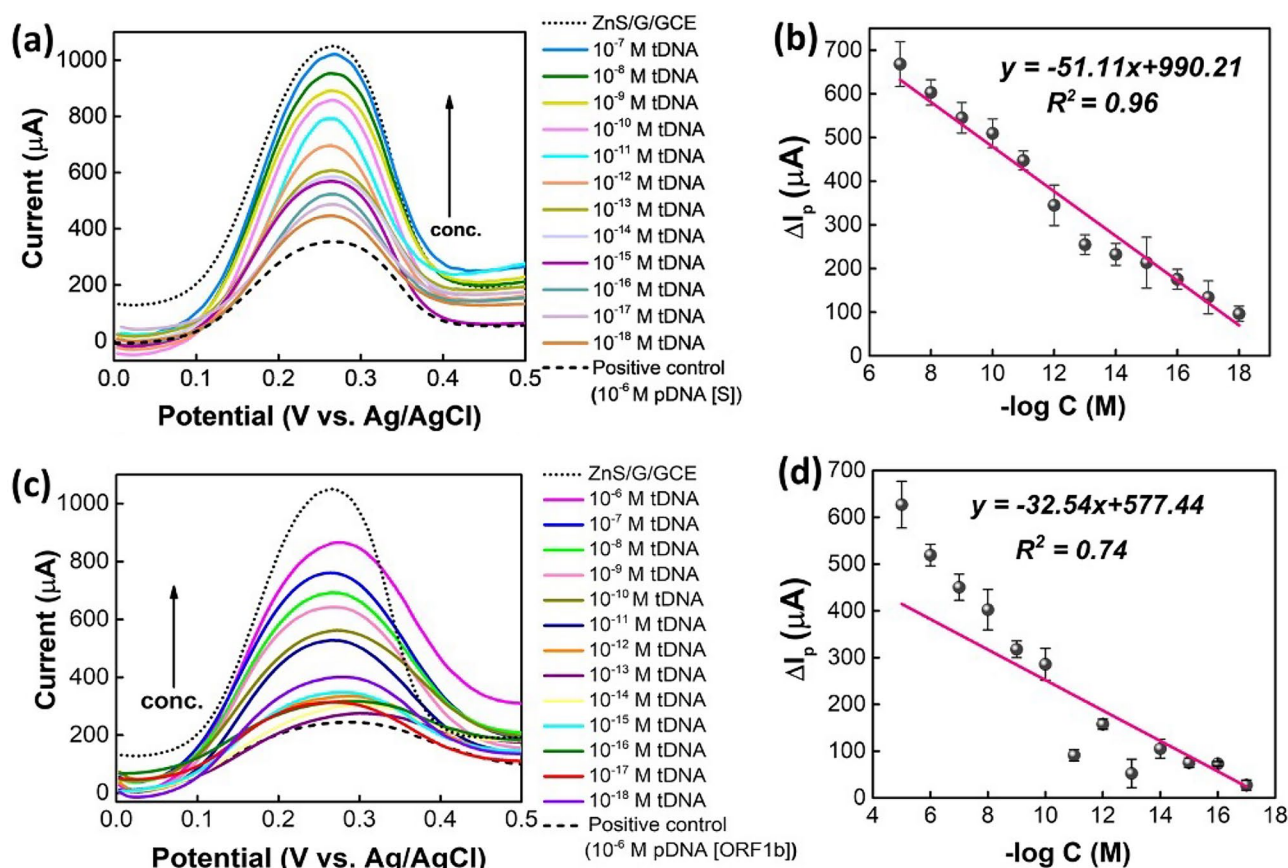


Fig. 3 **a** DPV plots and **b** the regression calibration plots (note: probe and target DNA samples are for S-gene). **c** DPV plots and **d** the regression calibration plots (Note: probe and complementary DNA samples are for ORF 1b-gene)

of hybridization with complementary targets, accompanied with the self-signal regeneration of the ZnS/graphene/GCE. With an increase in tDNA concentration, more of the pDNA can get hybridized and then release into the electrolyte. Therefore, the electron transfers as well as the peak currents increase. Furthermore, the regression calibration plot was

drawn from ΔI_p vs. $-\log C$ with the correlation coefficient estimated as $R^2 = 0.96$ as shown in Fig. 3b. Thus, the limit of detection ($\text{LOD} = 3\sigma/\text{slope}$) was determined, which was found to be 4.453×10^{-20} M.

In addition, the DPV tests were performed for the ORF 1b-gene. As displayed in Fig. 3c, the results follow a similar

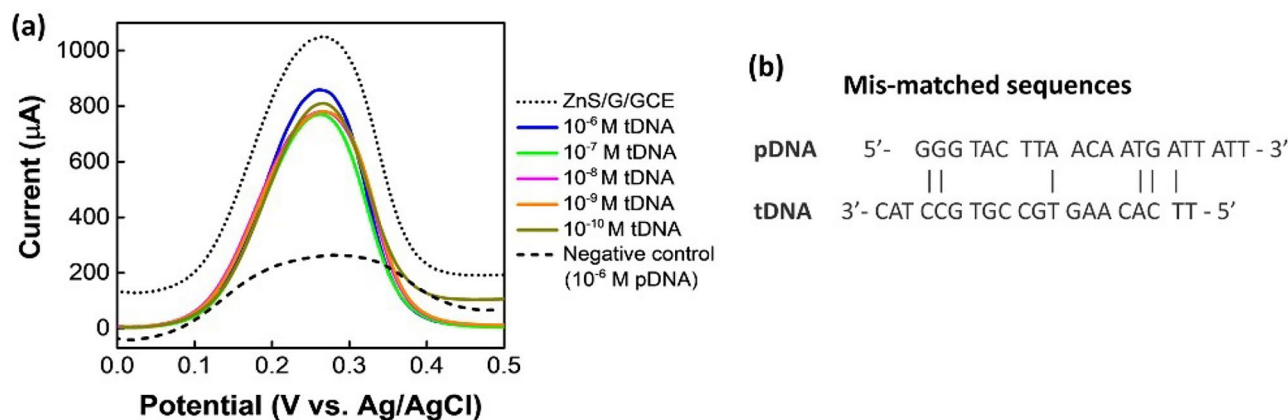
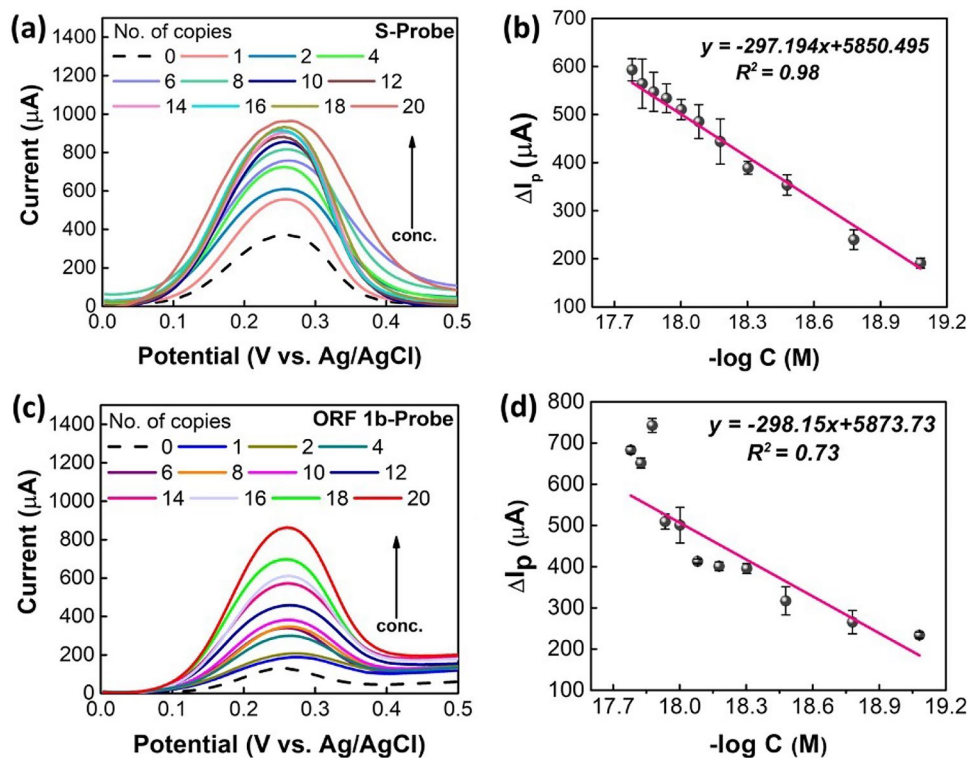


Fig. 4 **a** DPV plots of synthetic samples. **b** Mismatched sequences of pDNA and tDNA (Note: probe and target DNA samples are negative control)

Fig. 5 **a, b** DPV and calibration plots for the detection of SARS-CoV-2 standard using S-probe. **c, d** DPV and calibration plots for the detection of SARS-CoV-2 standard solutions using ORF 1b-probe in a solution containing 1.0 M KCl and 0.2 M $K_3[Fe(CN)_6]$



trend like S-gene. However, at the lower concentrations of cDNA samples from 10^{-12} M to 10^{-18} M, results showed arbitrary currents, and therefore the regression coefficient is only 0.74 (Fig. 3d). Based on these findings, the LOD was measured to be 2.013×10^{-18} M.

Following the same steps and procedures, DPV tests were also performed for mismatched negative control sample, as displayed in Fig. 4a. Since there is only a few of matched sequences present in these samples (Fig. 4b), the tDNA cannot hybridize successfully with the pDNA. Therefore, although after hybridization the currents get increased from the pDNA-modified electrode, there is no particular trend found for these samples. The matched sequence of S and ORF-1b genes are shown in Fig. S2.

3.4 Detection of SARS-CoV-2 standard samples

Following the similar DPV characterization technique, the detection of SARS-CoV-2 standard samples was performed. The positive control from TaqPath™ COVID-19 Fast PCR Combo Kit 2.0 was diluted to a working stock of 20 copies/ μ L on ice (ThermoFisher). This Fast PCR kit Positive Control consists of several different gene templates (200,000 copies/1 mL), such as S, N, and ORF1ab, which is widely used by CDC CLIA certified laboratories. Any of the probe templates targeting these genes can be used to perform the validation of the detection tests; therefore, we tested the S and ORF1b probes used in Fig. 2. As

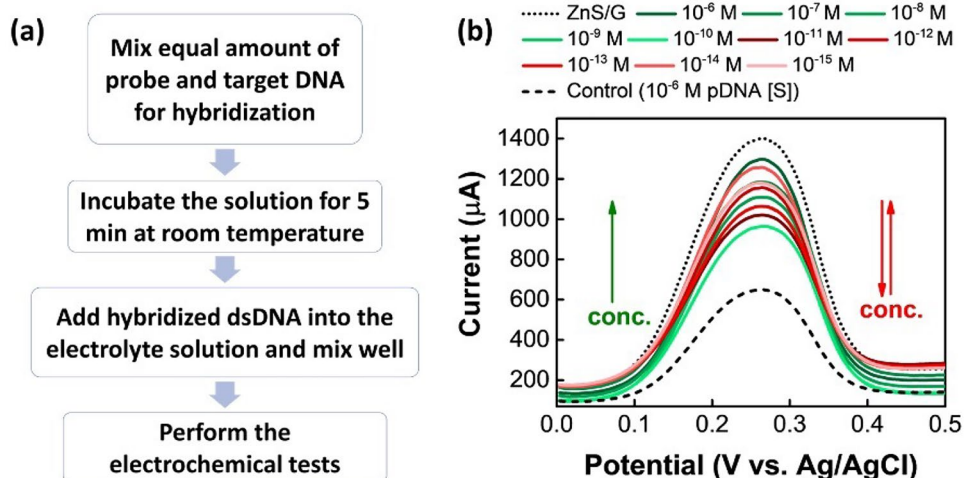
shown in Fig. 5a, S-probe can detect from 20 copies up to 1 copy. The corresponding concentrations of different number of copies are displayed in supporting information, Table S2. From these results, one thing is noticeable that the low concentrations at the range of 10^{-18} M to 10^{-20} M can be detected. Despite of the presence of all different genes in a standard solution, only a single probe template was used to detect the hybridized dsDNA. Hence, there is a huge chance of some target templates remain immobilized

Table 2 Comparison of the designed biosensors for the detection of SARS-CoV-2

Platform	Detection Method	LOD	Reference
Pt/Ti/glass wafer	Impedance	1×10^{-8} M	[43]
Co-TNTs	Amperometry	7×10^{-10} M	[44]
Graphene	SWV	2.6×10^{-7} M	[45]
Au IDE	EIS	3.9×10^{-16} M	[46]
CSPE/CNF-AuNP	EIS	7×10^{-13} M	[47]
rGO-Au	EIS	13×10^{-15} M	[48]
PANI/GCE	DPV	3.5×10^{-15} M	[31]
MIP/Au-TFE	DPV	15×10^{-15} M	[49]
AuNPs/FTO	DPV	10×10^{-15} M	[50]
ZnS/graphene/GCE	DPV	2.1×10^{-20} M	This work

Co-TNTs cobalt functionalized TiO_2 nanotubes, *IDE* interdigitated electrode, *CSPE* carbon-based screen-printed electrode, *CNF* carbon nanofibers, *MIP* molecular imprinted polymer, *TFE* thin film electrode, *FTO* fluorine doped tin oxide electrode

Fig. 6 **a** Detection steps of synthetic DNA samples via one-step hybridization method. **b** DPV plots to detect S-gene by one-step method [note: probe and complementary DNA samples are for S-gene]



on the surface of ZnS/graphene/GCE even after rinsing with DI water, which eventually may hinder the electron transfer process resulting arbitrary change in peak currents. Similar to the results for synthetic base-pair DNA samples, the detection of SARS-CoV-2 standard sample also provides the similar trend of increase in peak currents by increasing the concentration of target samples. Moreover, the regression calibration plot (Fig. 5b) was drawn and LOD values were calculated to be 2.068×10^{-20} M ($R^2 = 0.98$) for S-probes. These results clearly indicate a good performance from S-probe to detect the SARS-CoV-2 standard solutions.

Moreover, the SARS-CoV-2 standard solutions were also detected using the ORF 1b-probes. For ORF 1b-probe (Fig. 5c, d), the results followed the trend from 1 to 20 copies, which has superior sensitivity as RT-PCR detection method using the TaqPath™ COVID-19 Combo Kit.

Additionally, the performance of the DNA biosensor for the detection of SARS-CoV-2 was compared with previously published studies. According to Table 2, the proposed ZnS/graphene modified GCE achieved a lower detection limit which has 5 orders of magnitude less than the other reports.

3.5 Investigating the detection of synthetic DNA samples by one-step hybridization

Besides investigating the detection of DNA samples following the multiple-step procedure, the one-step hybridization technique was also applied in this study. The major steps are displayed in Fig. 6a, where instead of drop coating the DNA samples onto the surface of ZnS/G-modified GCE, the probe and target DNA samples were directly mixed into the electrolyte solution. Subsequently, the DPV tests were performed using the ZnS/graphene/GCE as the working electrode. From the results displayed in Fig. 6b, at high concentration range from 1×10^{-6} M to 1×10^{-10} M of cDNA,

the results showed the similar trend as before. However, the samples with low concentrations from 1×10^{-11} M to 1×10^{-15} M, the results are inconsistent. These results can be improved by handling the samples more carefully. Moreover, this one-step hybridization can save a lot of sample-preparation time comparing to the multiple-step procedure. Further optimizations, specifically for incubation time and temperature are still required to establish a stable detection method using one-step hybridization.

4 Conclusions

In conclusion, we report the construction of novel electrochemical sensing nanohybrid of ZnS/graphene for detecting SARS-CoV-2 nucleic acids by a one-pot and ultrafast microwave-controlled synthesis approach. The ZnS/graphene modified GCE exhibited promising electrochemical property and faster electron transfer kinetic which improved the detection performance. When the pDNA was immobilized on the surface of ZnS/graphene/GCE, the peak current decreased due to the obstruction of electron transfer process. Hybridization between the probe and target DNA caused the regeneration of the self-signal of ZnS/graphene. The peak current showed linear trend with decreasing the concentration of target DNA. Among different pDNA sequences, S-gene provides a better detection result when hybridized with target DNA or standard SARS-CoV-2 controls where the limits of detection are 4.453×10^{-20} M and 2.068×10^{-20} M, respectively. In addition, we presented a one-step hybridization to save the multiple-step procedure. A similar result was obtained only at high concentrations from 1×10^{-6} M to 1×10^{-10} M. The investigation of the one-step hybridization could be further studied. Overall, the microwave-synthesized composite of ZnS/graphene coating on the GCE electrode showed superior sensitivity

and low-concentration detection limit. Our strategy and assays developed here could be useful for rapid virus infection screening, public health surveillance, and in the settings with limited laboratory resources and utilized by users with limited training. Our platform will enable robust, low-cost, multiplex, and quantitative DNA/RNA analysis tools for field-deployable detection devices.

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Author contribution All authors contributed to the study conception and design. Etinosa Iyayi and Haseena Polk contributed to sample preparation. Material preparation, data collection, and analysis were performed by Shatila Sarwar and Carolina Ameza. Zhen Wei performed the SEM, XPS, and XRD characterization. Both Shatila Sarwar and Mao-Chia Lin aided in interpreting the results and worked on the manuscript. Ruigang Wang, Honghe Wang, and Xinyu Zhang supervised the project. All authors read and approved the final manuscript.

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Declarations

Conflict of interest The authors declare no competing interests.

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