Non-Enzymatic Amperometric Glucose Sensor Based on Carbon Nanodots and Copper Oxide Nanocomposites Electrode

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Abstract: In this research work, a non-enzymatic amperometric sensor for the determination of glucose was designed based on carbon nanodots (C-dots) and copper oxide (CuO) nanocomposites (CuO-C-dots). The CuO-C-dots nanocomposites were modified on the surface of a screen-printed carbon electrode (SPCE) to increase the sensitivity and selectivity of the glucose sensor. The as-synthesized materials were further analyzed for physico-chemical properties through characterization tools such as transmission electron microscopy (TEM) and Fourier-transform infrared spectroscopy (FTIR); and their electrochemical performance was also studied. The SPCE modified with CuO-C-dots possess desirable electrocatalytic properties for glucose oxidation in alkaline solutions. Moreover, the proposed sensing platform exhibited a linear range of 0.5 to 2 and 2 to 5 mM for glucose detection with high sensitivity (110 and 63.3 μA mM⁻¹ cm⁻²), and good selectivity and stability; and could potentially serve as an effective alternative method of glucose detection.

Keywords: non-enzymatic; carbon nanodots; copper oxide; glucose sensor; amperometry

1. Introduction

Diabetes is a disease that occurs when blood glucose level in the body is extremely high, reducing the cells’ ability to absorb sugar and convert it into energy. Normal blood glucose levels in humans should be less than 5.5 mM, with diabetics recording 7.0 mM or more (National Institute for Health and Care Excellence, NICE) [1]. Diabetics are required to check their glucose levels several times a day and take insulin to maintain stability. Hence, the determination of glucose concentration in body fluids is important for the effective diagnosis, monitoring, and treatment of diabetic patients. The challenge in controlling diabetes is strongly associated with the accurate, rapid, and sensitive monitoring of glucose levels and extensive efforts have been devoted to this end in recent years [2–5].

Glucose biosensors have greatly contributed to the monitoring of glucose levels in diabetic patients [6–9]. Enzymatic electrochemical biosensors, for example, offer good selectivity and
sensitivity [6,10–13]. However, they also have drawbacks, such as tedious immobilization steps, constrained operational conditions, and poor stability/reproducibility [6,14]. To overcome these shortcomings, enzyme-free (non-enzymatic) glucose biosensing platforms are being designed and developed on the basis of direct oxidation of glucose [15–18].

Another available and widely used technique for glucose detection is the amperometry. Amperometry offers several advantages, such as ease of use, short analysis time, low detection limit, and lower cost when compared to other sensors [19–21]. The traditional amperometric glucose sensor uses an enzyme—glucose oxidase (GOx)—to catalyze the oxidation of glucose and realize high performance (high selectivity/sensitivity). The GOx acts as a catalyst for the oxidation of glucose at the electrode, forming gluconolactone and hydrogen peroxide. The detection of hydrogen peroxide generated at the electrode relates to the concentration of glucose in the sample. However, the enzyme has limitations, such as high cost, poor stability, and low reproducibility. Enzyme activity is also easily affected by temperature, pH, and chemicals [6]. A non-enzymatic sensor, thus, is a suitable alternative, using nanostructured materials that offer enzyme-like activity or nanozymes modified on the surface of the electrode. The nanozymes act as electrocatalysts to generate electrical current by directly oxidizing glucose on the surface of the electrode. They offer several advantages, such as longer storage life, simple modification process, and high selectivity/sensitivity. Therefore, extensive research efforts have been devoted to developing an enzyme-free glucose sensor [22–26]. Non-enzymatic glucose sensors form an oxidation layer, which catalyzes a glucose oxidation reaction, similar to that of the enzyme activity. Various metal nanostructures and their oxides are used as catalysts, including noble metals such as platinum (Pt), gold (Au), and silver (Ag), which exhibit high sensitivity and low detection limits for glucose detection. However, noble metal electrodes have high production costs further limiting their potential applications. Therefore, low-cost materials such as nickel (Ni) and copper (Cu) and their oxides can be used instead [27–32].

In this work, we used copper oxide (CuO) as a catalyst because of its superior capabilities—low cost, low toxicity, facile preparation, and remarkable electrochemical catalytic property [33–36]. The electrocatalysis process of glucose oxidation is mediated by Cu(II) in an alkaline solution, which oxidizes into Cu(III) on the surface electrode. Cu(III) is a highly oxidizing state and oxidizes glucose into gluconolactone before reverting back to Cu(II). The current signals acquire electrons by the redox process of Cu(II) and Cu(III) [37–39]. Researchers have focused on employing carbon-based nanostructures—such as carbon nanotube (CNT), graphene or graphene oxide (GO)—as supporting materials, with the aim to modify working electrodes, improve sensitivity of the electrode, enhance the surface area, and transfer charge to the nanozyme. Carbon nanodots (C-dots), thus, are a newly emerging class of carbon materials that consist of carbon nanoparticles with diameter less than 10 nm and several hydrophilic surface groups exhibiting excellent water dispersibility, high surface area, and chemical stability [40–42]. In addition, a screen-printed carbon electrode (SPCE) has been conveniently used for glucose detection owing to advantages such as flexible design and scope for modification. In this study, a non-enzymatic amperometric glucose sensor based on carbon nanodots (C-dots) and copper oxide (CuO) nanocomposites acting as nanozymes was judiciously designed and developed. The as-developed nanocomposites were appropriately characterized and subsequently used to modify SPCE to construct an enzyme-free glucose sensor. The as-constructed non-enzymatic sensing platform possesses unique properties with high active surface area and electrocatalytic activity. In comparison to other materials reported previously, the CuO-C-dots provide primary advantages of improving sensitivity due to the synergistic effects of the nanocomposite (good catalytic activity for glucose oxidation of CuO nanoparticles), increase in surface area of the electrode, and prevention of agglomeration of CuO nanoparticles from C-dots. Besides, the as-developed sensor platform requires facile fabrication and do not involve complicated production processes.
2. Materials and Methods

2.1. Materials, Chemicals, and Instrumentation

The materials and chemicals used in this work are listed as follows: graphite rod (99.999%, St. Louis, MO, USA, Sigma-Aldrich), ultrapure water (18.2 MΩ cm−1), ethanol (99.9%, Bangkok, Thailand, RCi Labscan Limited), sodium hydroxide (Lobachemie, Mumbai, India), copper(II) oxide nanopowder (Sigma-Aldrich, St. Louis, USA), poly-(dimethyldiallylammonium chloride) (PDDA, MW = 200,000–350,000, 20 wt% in water, Sigma-Aldrich, St. Louis, MO, USA), D-glucose (Fisher scientific, New Hampshire, USA), sucrose (Ajax Finechem, Prospect, South Australia), lactose (Fluka, St. Louis, MO, USA), dopamine (Sigma-Aldrich, St. Louis, MO, USA), ascorbic acid (Ajax Finechem, Prospect, South Australia), and uric acid (Sigma-Aldrich, St. Louis, MO, USA). All the chemicals used were of analytical reagent grade.

Transmission electron microscopy (TEM) and scanning electron microscopy (SEM) images were obtained using a transmission electron microscope (JEM 2010, Jeol, Japan) and a field emission scanning electron microscope (JSM 6335 F, Jeol, Japan), respectively. FTIR spectrum was recorded on a FTIR spectrophotometer (Thermo Scientific, Massachusetts, USA) from 400 to 4000 cm−1. EmStat potentiostats (PalmSens, Houten, Netherlands) was employed for electrochemical detection.

2.2. Synthesis of carbon nanodots (C-dots)

C-dots were synthesized with an electrochemical exfoliation method, using two graphite rods (diameter 3 mm) as the working electrode and the counter electrode placed at a distance of 1.0 cm. Typically, ethanol (140 mL), ultrapure water (10 mL), and NaOH (0.12 g) were mixed to obtain an electrolyte precursor solution. Static potential of 60 V from a direct current (DC) power supply was applied to the two electrodes for 3 h under continuous stirring and N2 atmosphere. After 3 h, the excess precipitates were removed by centrifugation at 6000 rpm for 10 min. Consequently, a homogeneous supernatant containing C-dots dispersion was obtained [43,44].

2.3. Synthesis of carbon nanodots and copper oxide nanocomposite (CuO-C-dots)

Firstly, 45 mL C-dots and 0.020 g CuO were mixed for 30 min with continuous stirring and the solution was then heated to 100 °C. Secondly, 5 mL of 0.5 M NaOH was added to the boiling solution and kept at that temperature for 5 min. After cooling down to room temperature, the suspension was copiously washed with water and ethanol. Finally, the product was subjected to evaporate and CuO-C-dots were collected [45].

2.4. Preparation of the CuO-C-dots Nanocomposite Modified Electrode

The prepared 10 mg of CuO-C-dots nanocomposite was dispersed in 1.00 mL of ultrapure water by ultrasonic treatment for 30 min. Then 3 µL of the suspension was dropped on the surface of the SPCE (diameter 2 mm) and dried under an infrared lamp. Finally, 5 µL of 0.5% PDDA solution was coated onto the electrode surface and dried under an incandescent lamp.

2.5. Electrochemical Measurement of Glucose

EmStat potentiostats was used for chronoamperometric glucose detection with a three-electrode system using the modified SPCE as a working electrode, carbon as an auxiliary electrode, and Ag/AgCl as a reference electrode using 50 µL of 0.1 M NaOH solution as a total volume of electrolyte solution. The current response was recorded for 100 s for all amperometric experiments.
3. Results and Discussion

3.1. Characterizations of C-dots and CuO-C-dots

C-dots were synthesized by the electrochemical exfoliation method. A graphite rod acted as the carbon source and alkaline alcohol as an electrolyte. After 3 h, the colorless solution changed into a dark yellow solution, indicating successful preparation of C-dots. The morphology of the synthesized C-dots was further characterized by TEM. It was found that well-dispersed C-dots were of uniform (small spherical) shape with an average diameter of 2 nm, as shown in Figure 1A,B, respectively. Moreover, the lattice fringe of C-dots was analyzed through high-resolution TEM (HRTEM), as shown in Figure 1C. The lattice fringe of about 0.367 nm can be assigned to the (002) reflection plane of graphite [46,47].

![Transmission electron microscopy image (A), size distribution histogram (B) and high-resolution transmission electron microscopy image (C) of the as-synthesized carbon nanodots.](image)

FTIR was then used to analyze the functional groups of C-dots, as shown in Figure 2. The presence of the oxygen-containing groups can be confirmed by the stretching vibration bands of broad peak of O−H at 3300 cm\(^{-1}\), C=O at 1648 cm\(^{-1}\), and the epoxide group at 1087 cm\(^{-1}\). Moreover, there were a few absorption peaks, including C−H stretching at 2974 cm\(^{-1}\), bending vibrations of aromatic C=C, and aromatic =C−H around 1378 cm\(^{-1}\) and 879 cm\(^{-1}\), respectively. Owing to the presence of polar functional groups, the synthesized C-dots showed a highly hydrophilic property and excellent dispersibility in water [43,44,48,49].

![Fourier-transform infrared spectroscopy spectrum of the synthesized carbon nanodots.](image)
X-ray photoelectron spectroscopy (XPS) of the CuO-C-dots nanocomposite was done to further confirm the product’s composition and chemical states. The high-resolution XPS spectrum of C 1s showed the highest intensity, demonstrating the presence of carbon in the nanocomposite structure, as shown in Figure 3A. The main peak at 285.0 eV corresponds to sp² (C-C, C=C) bonding. The shoulders at about 288.5 eV and 290.0 eV correspond to C=O and O=C=O binding energies (BEs), respectively. The BEs for the chemical states of C-OH and C-O typically lies at 286.6 eV. The high high-resolution O 1s spectrum in Figure 3B further confirms the presence of CuO and Cu₂O. The other two peaks are tentatively assigned at 532.6 eV, ascertained by to the presence of Cu-O-C bonding and the C=O groups in C-dots. The BE at 533.7 eV can be attributed to the original oxygen in C-dots and chemisorbed water molecules (H₂O). Besides, the high-resolution XPS spectrum of Cu 2p shows two characteristic peaks with BEs at 932.1 and 951.7 eV, which can be assigned to Cu 2p₃/₂ and Cu 2p₁/₂, respectively (Figure 3C). The binding energy of the Cu 2p₃/₂ peaks lies in the range of 932.2 eV and 933.3–934.5 eV for Cu⁺ in Cu₂O and Cu²⁺ in CuO, respectively, suggesting the presence of CuO in the nanocomposite catalyst. The BEs at 932.1, 933.3, and 954.5 eV correspond to Cu 2p₁/₂ and Cu 2p₃/₂, respectively, in CuO. The BEs at 951.7, 952.9, and 954.1 eV correspond to Cu 2p₁/₂ and Cu 2p₃/₂ in Cu₂O, respectively [35,47].

The nanocomposites of PDDA/CuO-C-dots were then prepared for application as nanozymes for the detection of glucose. The SEM image of the surface of CuO-C-dots modified on SPCE is shown in Figure 4A. It was found that CuO-C-dots were well covered on the electrode surface with larger surface area, and carbon, oxygen, and copper was found in the EDS spectrum, as shown in Figure 4B. In addition, reticular structures were obtained on incorporation of PDDA onto CuO-C-dots, as presented in Figure 4C. The EDS spectrum shown in Figure 4D confirms the existence of carbon, copper, oxygen, nitrogen, and chloride (from PDDA), indicating that the PDDA/CuO-C-dots were successfully modified.

![Figure 3](#)

**Figure 3.** The high-resolution core-level X-ray photoelectron spectroscopy spectra of C 1s (A), O 1s (B), and Cu 2p (C).
When C-dots were modified on SPCE, a slightly higher background current was observed because (2) good catalytic activity for glucose oxidation of CuO nanoparticles, and (3) enhanced surface area of C-dots. Similarly, the current response was significantly increased on the CuO/SPCE, indicating that CuO had ample electrocatalytic activity to electron transfer between glucose and the electrode surface, and reduce the electrocatalytic active site of glucose’s oxidation reaction, as shown in Figure 5 (purple line) [50].

3.2. Electrochemical Characterization of the Proposed Electrode

Cyclic voltammetry was used to investigate the electrochemical characteristics of glucose oxidation on various modified electrodes in 0.10 M NaOH solution with 5 mM glucose. Cyclic voltammograms are shown in Figure 5. There was no obvious signal response obtained from bare SPCE and PDDA/SPCE. When C-dots were modified on SPCE, a slightly higher background current was observed because of a large surface area of C-dots. Similarly, the current response was significantly increased on the CuO/SPCE, indicating that CuO had ample electrocatalytic activity to glucose oxidation. In addition, the PDDA/CuO-C-dots exhibited a markedly improved current response of glucose oxidation with oxidation peak of about 0.50 V vs. Ag/AgCl. The probable reason for this might be the synergistic effects of nanocomposites, which include (1) good electrical conductivity and high stability of PDDA, (2) good catalytic activity for glucose oxidation of CuO nanoparticles, and (3) enhanced surface area of the electrode and prevention of agglomeration of CuO nanoparticles from C-dots.

Figure 5. CVs of bare SPCE, C-dots/SPCE, CuO/SPCE, PDDA/SPCE, and PDDA/CuO-C-dots/SPCE for detection of 5 mM glucose in 0.10 M NaOH at a scan rate of 90 mV s⁻¹.
The electrocatalytic oxidation of glucose in alkaline solution at the PDDA/CuO-C-dots/SPCE can be described as follows: CuO is firstly oxidized to CuOOH as a strong oxidizing agent. The Cu(III) species then electrochemically oxidized glucose to gluconolactone, responding to the oxidation peak of glucose’s oxidation reaction, as shown in Figure 5 (purple line) [50].

3.3. Effect of CuO-C-dots Modified on Screen-printed Carbon Electrode

A higher current response of glucose oxidation was obtained from a large catalytic active site on the modified electrode surface. Therefore, the amount of CuO-C-dots nanocomposite was optimized by depositing different layers of CuO-C-dots (each layer is 3 µL of CuO-C-dots suspension) before coating the electrode surface with PDDA. The as-coated multiple layers of suspension were studied by constructing calibration graphs of glucose in a range of 0.5 to 2 mM using amperometric detection at a constant potential of +0.50 V vs. Ag/AgCl; the slopes obtained indicate sensitivity of different layer-modified electrodes. Sensitivity obviously increased with rise in CuO-C-dots layer up to four layers and decreased after continuous deposition, as shown in Figure 6. Thick films can hinder electron transfer between glucose and the electrode surface, and reduce the electrocatalytic active area of the modified electrode. Thus, four sequential layers of CuO-C-dots were selected for further study.

![Figure 6](image_url)

Figure 6. Sensitivity of the electrode achieved with different layers of CuO-C-dots on SPCE for glucose determination in 0.1 M NaOH.

3.4. Amperometric Detection of Glucose by the Proposed Electrode

Chronoamperometry was employed as a detection technique to determine glucose using PDDA/CuO-C-dots/SPCE as a working electrode at a constant oxidation potential of +0.50 V. The steady current from 80–100 s was used as a current response, and current increased with increasing glucose concentration, as shown in Figure 7A. The calibration graph in Figure 7B shows two linear ranges: 0.5 to 2 mM and 2 to 5 mM with a limit of detection of 0.2 mM. The current began to saturate at a glucose concentration higher than 5 mM due to limitations of the active surface of the modified electrode. Their corresponding linear equations were $I (\mu A) = 3.4708C \text{ (mM)} + 0.3777$ with $R^2 = 0.9994$ and $I (\mu A) = 1.9552C \text{ (mM)} + 2.6148$ with $R^2 = 0.9977$, respectively. It was found that sensitivity at high concentration was lower than that at low concentration due to the saturated adsorption dynamics of the former. A summary of the analytical performance of various glucose sensors, including enzymatic and non-enzymatic ones, is shown in Table 1. The developed non-enzymatic glucose sensor exhibited good analytical characteristics such as good linearity and high sensitivity. Moreover, the preparation and detection procedures of the proposed method were also simple, quick, and cost-effective.
At a concentration of 0.1 mM, ascorbic acid, uric acid, dopamine, sucrose, and lactose were investigated. Storing the modified electrode at room temperature, it was found that the fabricated sensor had excellent selectivity toward glucose. The fabricated sensor showed a negligible current response of other species, compared to glucose, demonstrating that the sensor had excellent selectivity toward glucose. The amperometric responses of the proposed sensor with 1 mM glucose and other interfering species were evaluated. The repeatability of the modified electrode was found to be 2.6% RSD (n = 7). Storing the modified electrode for 12 days resulted in no obvious change in current response, highlighting good stability.

3.5. Selectivity and Stability of the Glucose Sensor

The physiological level of glucose concentration is normally about 3–8 mM, which is higher than that of main interferences such as ascorbic acid (0.1 mM) and uric acid (0.1 mM) [56]. Therefore, the amperometric responses of the proposed sensor with 1 mM glucose and other interfering species (ascorbic acid, uric acid, dopamine, sucrose, and lactose) at a concentration of 0.1 mM were investigated. Figure 8 shows a negligible current response of other species, compared to glucose, demonstrating that the fabricated sensor had excellent selectivity toward glucose.

| Sensing Electrode | Detection Method | Potential (V) | Sensitivity (µA mM⁻¹cm⁻²) | Linear Range (mM) | LOD (mM) | Ref. |
|-------------------|-----------------|---------------|---------------------------|-------------------|---------|------|
| GOx/Cds/Gr on GCE | CV              | -             | 1.76                      | 2–16              | 0.7     | [51] |
| PDDA/Ch/GOx/PtAuNPs/PtZn on Pt | Amp            | +0.60         | 17.85                     | 0.01–8            | 0.001   | [52] |
| Au/GO on GCE      | Amp             | +0.0          | 5.20, 4.56                | 0.1–2, 2–16       | 0.025   | [53] |
| Cu/Cu₂O/CsS on GCE | Amp            | +0.65         | 63.8, 22.6                | 0.01–0.69, 3.19–3.69 | 0.005   | [54] |
| Nafion/NPC-CB on GCE | Amp            | +0.64         | 33.75                     | 0.006–3.369       | 0.002   | [55] |
| PDDA/CuO-C-dot on SPCE | Amp            | +0.50         | 110, 62.3                 | 0.5–2, 2–5        | 0.2     | This work |

Amp = amperometry, Gr = graphene, GCE = glassy carbon electrode, PtAuNPs = platinum and gold nanoparticles, GO = graphene oxide, CsS = carbon spheres, and NPC-CB = nanoporous copper and carbon back

Figure 7. The chronoamperometric responses of PDDA/CuO-C-dots/SPCE for various glucose concentrations (0.5–5 mM) in 0.1 M NaOH at +0.50 V (A) and their calibration curve of current response vs. glucose concentration (B).

Figure 8. Amperometric response of the proposed sensor to interfering species.
there was no obvious change in current response after 12 days, thus highlighting good stability of the proposed sensor. Batch variation of the CuO-C-dots-based amperometric sensor was evaluated. The repeatability of the modified electrode was found to be 2.6% RSD (n = 7).

3.6. Applicability of the Prepared Sensor for Glucose Measurement in Real Time

To evaluate the practical usefulness of the as-fabricated sensor, the proposed sensor was used to determine glucose levels in human blood serum samples. Although many proteins in serum samples can act as insulators and block electron transfer between the solution and the electrode surface, the serum samples should be significantly diluted (100 times with 0.1 M NaOH solution) before analysis. The samples were spiked with standard glucose to obtain final concentrations at 0.5, 1.0, and 3.0 mM. The recoveries of the spiked samples were found from 88 to 94%, as shown in Table 2. The analytical results indicate that the PDDA/CuO-C-dots nanocomposite modified on SPCE is reliable for determining glucose, and nanocomposites could be a possible alternative nanzyme in constructing new kind of glucose sensors. However, to enhance the applicability of the sensor for low-level detection, sample preparation should be optimized to further separate other type of interferences.

| Sample | Spiked Glucose (mM) | Detected Glucose (mM) | %Recovery |
|--------|---------------------|-----------------------|-----------|
| 1      | 0.5                 | 0.47 ± 0.05           | 94        |
| 2      | 1.0                 | 0.88 ± 0.08           | 88        |
| 3      | 3.0                 | 2.71 ± 0.26           | 90        |

4. Conclusions

In summary, we successfully prepared a CuO-C-dots nanocomposite and fabricated a non-enzymatic glucose sensor based on PDDA/CuO-C-dots, modified on an SPCE electrode. Due to the synergistic effects of the high surface area of C-dots, good electrical conductivity of PDDA, and remarkable catalytic active sites of CuO nanoparticles, the as-prepared nanocomposites showed excellent electrochemical properties in catalyzing the oxidation reaction of glucose. The developed sensor (CuO-C-dots/SPCE) offers several advantages, which include simple procedures for preparation and detection, fast analysis, high sensitivity, and good selectivity and stability. Moreover, the satisfied recoveries indicate that the sensor is reliable and can be applied to the determination of glucose in real samples.

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References
1. Blood Sugar Level Ranges. Available online: https://www.diabetes.co.uk/diabetes_care/blood-sugar-level-ranges.html (accessed on 15 November 2019).
2. Wang, J. Electrochemical glucose biosensors. Chem. Rev. 2008, 108, 814–825. [CrossRef] [PubMed]
3. Makaram, P.; Owens, D.; Aceros, J. Trends in nanomaterial-based non-invasive diabetes sensing technologies. Diagnostics 2014, 4, 27–46. [CrossRef] [PubMed]
4. Tian, K.; Prestgard, M.; Tiwari, A. A review of recent advances in nonenzymatic glucose sensors. *Mater. Sci. Eng. C* 2014, 41, 100–118. [CrossRef] [PubMed]

5. Bruen, D.; Delaney, C.; Florea, L.; Diamond, D. Glucose Sensing for Diabetes Monitoring: Recent Developments. *Sensors* 2017, 17, 1866. [CrossRef] [PubMed]

6. Gopalan, A.I.; Muthuchamy, N.; Lee, K.P. A novel bismuth oxychloride-graphene hybrid nanosheets based non-enzymatic photoelectrochemical glucose sensing platform for high performances. *Biosens. Bioelectron.* 2017, 89, 352–360. [CrossRef] [PubMed]

7. Gopalan, A.I.; Muthuchamy, N.; Komathi, S.; Lee, K.P. A novel multicomponent redox polymer nanobead based high performance non-enzymatic glucose sensor. *Biosens. Bioelectron.* 2016, 84, 53–63. [CrossRef] [PubMed]

8. Muthuchamy, N.; Gopalan, A.; Lee, K.-P. Highly selective non-enzymatic electrochemical sensor based on a titanium dioxide nanowire–poly(3-aminophenyl boronic acid)–gold nanoparticle ternary nanocomposite. *RSC Adv.* 2018, 8, 2138–2147. [CrossRef]

9. Sai-Anand, G.; Gopalan, A.-I.; Kang, S.-W.; Komathi, S.; Lee, K.-P. One pot synthesis of new gold nanoparticles dispersed poly (2-aminophenyl boronic acid) composites for fabricating an affinity based electrochemical detection of glucose. *Sci. Adv. Mater.* 2014, 6, 1356–1364. [CrossRef]

10. Gopalan, A.I.; Komathi, S.; Sai Anand, G.; Lee, K.P. Nanodiamond based sponges with entrapped enzyme: A novel electrochemical probe for hydrogen peroxide. *Biosens. Bioelectron.* 2013, 46, 136–141. [CrossRef]

11. Manesh, K.M.; Santhosh, P.; Uthayakumar, S.; Gopalan, A.I.; Lee, K.P. One-pot construction of mediatorless bi-enzymatic glucose biosensor based on organic-inorganic hybrid. *Biosens. Bioelectron.* 2010, 25, 1579–1586. [CrossRef]

12. Santhosh, P.; Manesh, K.M.; Uthayakumar, S.; Komathi, S.; Gopalan, A.I.; Lee, K.P. Fabrication of enzymatic glucose biosensor based on palladium nanoparticles dispersed onto poly(3,4-ethylenedioxythiophene) nanofibers. *Biosens. Bioelectron.* 2019, 75, 61–66. [CrossRef]

13. Anand, G.S.; Gopalan, A.I.; Kang, S.-W.; Lee, K.-P. Development of a surface plasmon assisted label-free calorimetric method for sensitive detection of mercury based on functionalized gold nanorods. *J. Anal. Atom. Spectrom.* 2013, 28, 488. [CrossRef]

14. Haldorai, Y.; Hwang, S.K.; Gopalan, A.I.; Huh, Y.S.; Han, Y.K.; Voit, W.; Sai-Anand, G.; Lee, K.P. Direct electrochemistry of cytochrome c immobilized on titanium nitride/multi-walled carbon nanotube composite for amperometric nitrite biosensor. *Biosens. Bioelectron.* 2016, 79, 543–552. [CrossRef] [PubMed]

15. Kavitha, T.; Gopalan, A.I.; Lee, K.-P.; Park, S.-Y. Glucose sensing, photocatalytic and antibacterial properties of graphene–ZnO nanoparticle hybrids. *Carbon* 2012, 50, 2994–3000. [CrossRef]

16. Ragupathy, D.; Gopalan, A.I.; Lee, K.-P. Synergistic contributions of multiwall carbon nanotubes and gold nanoparticles in a chitosan–ionic liquid matrix towards improved performance for a glucose sensor. *Electrochem. Commun.* 2009, 11, 397–401. [CrossRef]

17. Anantha-Iyengar, G.; Shanmugasundaram, K.; Nallal, M.; Lee, K.-P.; Whitcombe, M.J.; Lakshmi, D.; Sai-Anand, G. Functionalized conjugated polymers for sensing and molecular imprinting applications. *Prog. Polym. Sci.* 2009, 34, 1–129. [CrossRef]

18. Sai-Anand, G.; Sivanesan, A.; Benzigar, M.R.; Singh, G.; Gopalan, A.-I.; Baskar, A.V.; Ilbeygi, H.; Ramadass, K.; Kambala, V.; Vinu, A. Recent Progress on the Sensing of Pathogenic Bacteria Using Advanced Nanostructures. *Bull. Chem. Soc. Jpn.* 2019, 92, 216–244. [CrossRef]

19. Komathi, S.; Gopalan, A.I.; Kim, S.-K.; Anand, G.S.; Lee, K.-P. Fabrication of horseradish peroxidase immobilized poly(N-[3-(trimethoxy silyl)propyl]aniline) gold nanorods film modified electrode and electrochemical hydrogen peroxide sensing. *Electrochim. Acta* 2013, 92, 71–78. [CrossRef]

20. Thunkhamrak, C.; Chuntib, P.; Ounnunkad, K.; Banet, P.; Aubert, P.H.; Saiand, G.; Gopalan, A.I.; Jakmunee, J. Highly sensitive voltammetric immunosensor for the detection of prostate specific antigen based on silver nanoprobe assisted graphene oxide modified screen printed carbon electrode. *Talanta* 2020, 208, 120389. [CrossRef]

21. Shanmugasundaram, K.; Sai-Anand, G.; Gopalan, A.-I.; Lee, H.-G.; Yeo, H.K.; Kang, S.-W.; Lee, K.-P. Direct electrochemistry of cytochrome c with three-dimensional nanoarchitected multicomponent composite electrode and nitrite biosensing. *Sens. Actuators B Chem.* 2016, 228, 737–747. [CrossRef]

22. Chen, X.; Wu, G.; Cai, Z.; Oyama, M.; Chen, X. Advances in enzyme-free electrochemical sensors for hydrogen peroxide, glucose, and uric acid. *Microchim. Acta* 2014, 181, 689–705. [CrossRef]
23. Niu, X.; Li, X.; Pan, J.; He, Y.; Qiu, F.; Yan, Y. Recent advances in non-enzymatic electrochemical glucose sensors based on non-precious transition metal materials: Opportunities and challenges. *RSC Adv.* 2016, 6, 84893–84905. [CrossRef]

24. Zhu, H.; Li, L.; Zhou, W.; Shao, Z.; Chen, X. Advances in non-enzymatic glucose sensors based on metal oxides. *J. Mater. Chem. B* 2016, 4, 7333–7349. [CrossRef]

25. Amala, G.; Gowtham, S.M. Recent advancements, key challenges and solutions in non-enzymatic electrochemical glucose sensors based on graphene platforms. *RSC Adv.* 2017, 7, 36949–36976.

26. Hwang, D.W.; Lee, S.; Seo, M.; Chung, T.D. Recent Advances in Electrochemical Non-Enzymatic Glucose Sensors—A Review. *Anal. Chim. Acta* 2018, 1033, 1–34. [CrossRef] [PubMed]

27. Jothi, L.; Jayakumar, N.; Jaganathan, S.K.; Nageswaran, G. Ultrasensitive and selective non-enzymatic electrochemical glucose sensor based on hybrid material of graphene nanosheets/graphene nanoribbons/nickel nanoparticle. *Mater. Res. Bull.* 2018, 98, 300–307. [CrossRef]

28. Yang, J.; Lin, Q.; Yin, W.; Jiang, T.; Zhao, D.; Jiang, L. A novel nonenzymatic glucose sensor based on functionalized PDDA-graphene/CuO nanocomposites. *Sens. Actuators B* 2017, 253, 1087–1095. [CrossRef]

29. Zhu, J.; Yin, H.; Cui, Z.; Qin, D.; Gong, J.; Nie, Q. Amorphous Ni (OH)2/CQDs microspheres for highly sensitive non-enzymatic glucose detection prepared via CQDs induced aggregation process. *Appl. Surf. Sci.* 2017, 420, 323–330. [CrossRef]

30. Baghayeri, M.; Amiri, A.; Farhadi, S. Development of non-enzymatic glucose sensor based on efficient loading Ag nanoparticles on functionalized carbon nanotubes. *Sens. Actuators B Chem.* 2016, 225, 354–362. [CrossRef]

31. Scandurra, A.; Ruffino, F.; Sanzaro, S.; Grimaldi, M.G. Laser and thermal dewetting of gold layer onto graphene paper for non-enzymatic electrochemical detection of glucose and fructose. *Sens. Actuators B Chem.* 2019, 301, 127113. [CrossRef]

32. Baghayeri, M.; Amiri, A.; Alizadeh, Z.; Veisi, H.; Hasheminejad, E. Non-enzymatic voltammetric glucose sensor made of ternary NiO/Fe3O4-SH/para-amino hippuric acid nanocomposite. *J. Electroanal. Chem.* 2018, 810, 69–77. [CrossRef]

33. Saianand, G.; Gopalan, A.I.; Lee, J.C.; Sathish, C.I.; Gopalakrishnan, K.; Unni, G.E.; Shanbhag, D.; Dasireddy, V.; Yi, J.; Xi, S.; et al. Mixed Copper/Copper-Oxide Anchored Mesoporous Fullerene Nanohybrids as Superior Electrocatalysts toward Oxygen Reduction Reaction. *Small* 2019, e1903937. [CrossRef] [PubMed]

34. Bui, Q-T.; Yu, I-K.; Gopalan, A.I.; Saianand, G.; Kim, W.; Choi, S-H. Facile Fabrication of Metal Oxide Based Catalytic Electrodes by AC Plasma Deposition and Electrochemical Detection of Hydrogen Peroxide. *Catalysts* 2019, 9, 888. [CrossRef]

35. Zhang, Y.; Li, N.; Xiang, Y.; Wang, D.; Zhang, P.; Wang, Y.; Lu, S.; Xu, R.; Zhao, J. A flexible non-enzymatic glucose sensor based on copper nanoparticles anchored on laser-induced graphene. *Carbon* 2020, 156, 506–513. [CrossRef]

36. Liu, X.; Yang, W.; Chen, L.; Jia, J. Three-Dimensional Copper Foam Supported CuO Nanowire Arrays: An Efficient Non-enzymatic Glucose Sensor. *Electrochim. Acta* 2017, 235, 519–526. [CrossRef]

37. Namdari, P.; Negahdari, B.; Eatemadi, A. Synthesis, properties and biomedical applications of carbon-based quantum dots: An updated review. *Biomed. Pharmacother.* 2017, 87, 209–222. [CrossRef]

38. Shi, L.; Zhu, X.; Liu, T.; Zhao, H.; Lan, M. Encapsulating Cu nanoparticles into metal-organic frameworks for nonenzymatic glucose sensing. *Sens. Actuators B Chem.* 2016, 227, 583–590. [CrossRef]

39. Wang, X.; Liu, E.; Zhang, X. Non-enzymatic glucose biosensor based on copper oxide-reduced graphene oxide nanocomposites synthesized from water-isopropanol solution. *Electrochim. Acta* 2014, 130, 253–260. [CrossRef]

40. Miao, P.; Han, K.; Tang, Y.; Wang, B.; Lin, T.; Cheng, W. Recent advances in carbon nanodots: Synthesis, properties and biomedical applications. *Nanoscale* 2015, 7, 1586–1595. [CrossRef]

41. Yulong, Y.; Xinsheng, P. Recent advances in carbon-based dots for electroanalysis. *Analyst* 2016, 141, 2619–2628. [CrossRef]

42. Tuerhong, M.; Yang, X.U.; Xue-Bo, Y. Review on carbon dots and their applications. *Chin. J. Anal. Chem.* 2017, 45, 139–150. [CrossRef]

43. Ming, H.; Ma, Z.; Liu, Y.; Pan, K.; Yu, H.; Wang, F.; Kang, Z. Large scale electrochemical synthesis of high quality carbon nanodots and their photocatalytic property. *Dalton Trans.* 2012, 41, 9526–9531. [CrossRef] [PubMed]
44. Liu, M.; Xu, Y.; Niu, F.; Gooding, J.J.; Liu, J. Carbon quantum dots directly generated from electrochemical oxidation of graphite electrodes in alkaline alcohols and the applications for specific ferric ion detection and cell imaging. *Analyst* 2016, 141, 2657–2664. [CrossRef] [PubMed]

45. Zhang, L.; Hai, X.; Xia, C.; Chen, X-W.; Wang, J.-H. Growth of CuO nanoneedles on graphene quantum dots as peroxidase mimics for sensitive colorimetric detection of hydrogen peroxide and glucose. *Sens. Actuators B Chem.* 2017, 248, 374–384. [CrossRef]

46. Liu, Y.; Liu, C.-y.; Zhang, Z.-y. Graphitized carbon dots emitting strong green photoluminescence. *J. Mater. Chem. C* 2013, 1, 4902. [CrossRef]

47. Wu, J.; Wang, P.; Wang, F.; Fang, Y. Investigation of the Microstructures of Graphene Quantum Dots (GQDs) by Surface-Enhanced Raman Spectroscopy. *Nanomaterials* 2018, 8, 864. [CrossRef] [PubMed]

48. Aoun, S.B. Nanostructured carbon electrode modified with N-doped graphene quantum dots–chitosan nanocomposite: A sensitive electrochemical dopamine sensor. *R. Soc. Open Sci.* 2017, 4, 171199. [CrossRef]

49. Deng, J.; Lu, Q.; Mi, N.; Li, H.; Liu, M.; Xu, M.; Tan, L.; Xie, Q.; Zhang, Y.; Yao, S. Electrochemical synthesis of carbon nanodots directly from alcohols. *Chem. A Eur. J.* 2014, 20, 4993–4999. [CrossRef]

50. Wei, H.; Sun, J.J.; Guo, L.; Li, X.; Chen, G.N. Highly enhanced electrocatalytic oxidation of glucose and shikimic acid at a disposable electrically heated oxide covered copper electrode. *Chem. Commun. (Camb.)* 2009, 20, 2842–2844. [CrossRef]

51. Wang, K.; Liu, Q.; Guan, Q.M.; Wu, J.; Li, H.N.; Yan, J.J. Enhanced direct electrochemistry of glucose oxidase and biosensing for glucose via synergy effect of graphene and CdS nanocrystals. *Biosens. Bioelectron.* 2011, 26, 2252–2257. [CrossRef]

52. Hossain, M.F.; Park, J.Y. Fabrication of sensitive enzymatic biosensor based on multi-layered reduced graphene oxide added PtAu nanoparticles-modified hybrid electrode. *PLoS ONE* 2017, 12, e0173553. [CrossRef] [PubMed]

53. Shu, H.; Chang, G.; Su, J.; Cao, L.; Huang, Q.; Zhang, Y.; Xia, T.; He, Y. Single-step electrochemical deposition of high performance Au-graphene nanocomposites for nonenzymatic glucose sensing. *Sens. Actuators B* 2015, 220, 331–339. [CrossRef]

54. Yin, H.; Cui, Z.; Wang, L.; Nie, Q. In situ reduction of the Cu/Cu2O/carbon spheres composite for enzymaticless glucose sensors. *Sens. Actuators B* 2016, 222, 1018–1023. [CrossRef]

55. Ling, M.; Pengchong, Z.; Jiuyun, C.; Dandan, C.; Ying, Q.; Ning, G.; Genhua, Z.; Rongjing, C. Non-enzymatic sensing of glucose and hydrogen peroxide using a glassy carbon electrode modified with a nanocomposite consisting of nanoporous copper, carbon black and nafion. *Microchim. Acta* 2016, 183, 1359–1365.

56. Safavi, A.; Maleki, N.; Farjami, E. Fabrication of a glucose sensor based on a novel nanocomposite electrode. *Biosens. Bioelectron.* 2009, 24, 1655–1660. [CrossRef] [PubMed]