A Non-Enzymatic Sensor Based on Trimetallic Nanoalloy with Poly (Diallyldimethylammonium Chloride)-Capped Reduced Graphene Oxide for Dynamic Monitoring Hydrogen Peroxide Production by Cancerous Cells

Jun Jiao, Meixin Pan, Xinran Liu, Binshuai Li, Jian Liu and Qiang Chen *

The Key Laboratory of Bioactive Materials Ministry of Education, College of Life Science, Nankai University, Tianjin 300071, China
* Correspondence: qiangchen@nankai.edu.cn

Received: 17 November 2019; Accepted: 19 December 2019; Published: 21 December 2019

Abstract: Catching cancer at an early stage is necessary to make it easier to treat and to save people’s lives rather than just extending them. Reactive oxygen species (ROS) have sparked a huge interest owing to their vital role in various biological processes, especially in tumorigenesis, thus leading to the potential of ROS as prognostic biomarkers for cancer. Herein, a non-enzymatic biosensor for the dynamic monitoring of intracellular hydrogen peroxide ($H_2O_2$), the most important ROS, via an effective electrode composed of poly (diallyldimethylammonium chloride) (PDDA)-capped reduced graphene oxide (RGO) nanosheets with high loading trimetallic AuPtAg nanoalloy, is proposed. The designed biosensor was able to measure $H_2O_2$ released from different cancerous cells promptly and precisely owing to the impressive conductivity of RGO and PDDA and the excellent synergistic effect of the ternary alloy in boosting the electrocatalytic activity. Built upon the peroxidase-like activity of the nanoalloy, the developed sensor exhibited distinguished electrochemical performance, resulting in a low detection limit of 1.2 nM and a wide linear range from 0.05 µM to 5.5 mM. Our approach offers a significant contribution toward the further elucidation of the role of ROS in carcinogenesis and the effective screening of cancer at an early stage.

Keywords: non-enzymatic biosensors; nanosheets; trimetallic nanoalloy; reactive oxygen species; living cancer cells

1. Introduction

Cancer is a global issue and has a major impact on public health. It is estimated that there were 18.1 million new cancer cases and 9.6 million deaths from cancer in 2018 around the world [1]. As the technology evolves, researchers are focusing on ways to detect, treat, and prevent cancer earlier to level off or decrease the incidence of and mortality due to cancer. Although cancer remains one of the leading causes of death, diagnosing it early offers the opportunity to improve survival rates [2]. The broader application of advanced techniques like X-ray imaging, computed tomography (CT), positron emission tomography (PET), and biopsy in cancer diagnosis has accelerated progress against cancer [3–5]. However, the accuracy of such typical methods is reduced if carcinogens are present at the very beginning. The sooner cancer is diagnosed, the higher the survival rate will be. Therefore, if identifying precancerous lesions or tumors in the early initiation phase can be accomplished, there is more chance of administering more precise therapies.
It has been proven that reactive oxygen species (ROS), including superoxide (O$_2^-$), hydrogen peroxide (H$_2$O$_2$), hydroxyl radicals, and peroxynitrite, are messenger molecules, which can turn various biological processes on and off [6]. Balanced ROS metabolism increases antioxidant ability, thereby establishing a barrier against tumorigenesis [7,8]. However, DNA damage caused by ROS results in genomic instability and altered gene expression, while ROS-mediated signaling pathways also drive cell proliferation and apoptosis suppression, thus promoting tumor formation [9,10]. Among the ROS, H$_2$O$_2$ has sparked huge interest in anticancer therapies as not only is it the main substance in the cellular response to oxidative stress, but it is also able to pass through cell membranes over a relatively long lifetime. In addition, cancer cells have metabolic and signaling aberrations, and thus exhibit an enhanced H$_2$O$_2$ level that differs from that in normal cells [11,12]. Therefore, the measurement of intracellular H$_2$O$_2$ could be of great value in clarifying its role in oncogenesis and exploring advanced therapeutic strategies for cancer and other ROS-related diseases.

Although various analytical techniques for measuring H$_2$O$_2$, such as spectrophotometry and chemiluminescence, have been developed, they still suffer drawbacks such as low detection limits and complex processing when faced with the challenge of achieving detection limits in the nanomolar range in living cells [13,14]. In this regard, electrochemical techniques with high sensitivity and simplicity to improve the timely detection of H$_2$O$_2$ in vivo have gained increasing attention. Enzymes like horseradish peroxidase have been widely used for biosensor fabrication as they own impressive specificity and notable sensitivity towards target analytes [15–17]. Dai et al. [18] immobilized horseradish peroxidase (HRP) on a boronic acid-functionalized metal–organic framework to fabricate a sensitive sensor for the real-time detection of H$_2$O$_2$ released from living cells. However, such enzymatic sensors have been gradually replaced by non-enzymatic ones owing to their deficiency in enzyme immobilization and instability [19–21]. Noble metals, alloys, and metal oxides have played remarkable roles as robust inorganic catalysts in non-enzymatic biosensors for bimolecular detection on account of their intrinsic peroxidase-like activity [22–26]. For instance, platinum revealed admirable catalytic activity for reduction of hydrogen peroxide, which effectively enhances the analytical performances of biosensors [27,28]. Muhammet et al. reported a non-enzymatic biosensor with outstanding electrochemical properties by decorating Pd@Au bimetallic nanoparticles and reduced graphene oxide on the surface of electrode for determination of H$_2$O$_2$. Meanwhile, the constructed sensor achieved the detection in a real milk sample [29]. Compared with their bimetallic or monometallic counterparts, trimetallic catalysts with an enlarged surface area, abundant active sites, and fast mass transfer present a pronounced synergistic effect, resulting in superior electrocatalytic activity and stability [30,31]. Hence, trimetallic catalysts hold considerable promise as competitive candidates for the real-time cellular monitoring of ROS levels after oxidative stress. Reduced graphene oxide (RGO) is a widely used competitor for the construction of sensing interfaces by virtue of its great electrical conductivity, its extensive surface area, and the fact that it is conveniently functionalizable [32–34]. However, it has a strong inclination to agglomerate irreversibly owing to van der Waals interactions, leading to poor dispersibility in aqueous solutions [35]. The cationic polymer poly (diallyldimethylammonium chloride) (PDDA) was introduced to tackle this problem as the electrostatic charge in RGO can be altered by PDDA, thus avoiding the agglomeration of RGO and revealing its more engaging characteristics such as hydrophilicity and modifiability [36,37].

In this work, we utilized a one-pot and cost-effective method for synthesizing PDDA-capped AuPtAg/RGO nanohybrids. The obtained nanocomposites presented stronger catalytic and conductive capacity via incrementing the electroactive surface area and enhancing the electron transport efficiency. By analyzing the electrochemical responses from the PDDA–RGO nanocomposites and nanoalloy particles, the dynamic levels of H$_2$O$_2$ in different living cancerous cells were determined almost instantly with satisfactory results (Scheme 1). The as-prepared sensor exhibited a limit of detection (LOD) as low as 1.2 nM, thus indicating its usefulness as a platform for monitoring differences in H$_2$O$_2$ in cancerous tissues and opening the door to identify its role in tumor formation.
Scheme 1. Schematic illustration for the preparation of poly (diallyldimethylammonium chloride) (PDDA)-AuPtAg/reduced graphene oxide (RGO)-modified glassy carbon electrode (GCE) applied for the detection of H$_2$O$_2$ released from SKOV3, MCF-7, and A431 cells stimulated with phorbol-12-myristate-13-acetate (PMA). ROS, reactive oxygen species.

2. Materials and Methods

2.1. Reagents and Apparatus

Graphene oxide (GO) was acquired from Nanjing XFNANO Materials Tech Co. (China). Platinic chloride hydrate, hydrogen tetrachloroaurate trihydrate (HAuCl$_4$·3H$_2$O, 99%), silver nitrate, poly (diallyldimethylammonium chloride) (PDDA), glucose, ascorbic acid (AA), uric acid (UA), xanthine, and phorbol-12-myristate-13-acetate (PMA) to liberate H$_2$O$_2$ from live cells were received from Sigma Aldrich Co. Cell culture medium (Dulbecco’s modified essential medium (DMEM), McCoy’s 5A) and Amplex Red Hydrogen Peroxide Assay kit was purchased from Thermo Fisher. These were used as such without any further purification. All working aqueous solutions were prepared with double distilled water.

All electrochemical measurements were conducted using 283 Potentiostat-Galvanostat electrochemical workstation (EG&GPARC with M270 software) with an ordinary triple-electrode system. It consisted of an Ag/AgCl electrode saturated with KCl, a platinum wire, and a modified glassy carbon electrode as the reference, counter, and adopted working electrode, respectively. All experiments were performed at room temperature. In addition, transmission electron microscopy (TEM) image analysis was obtained from Tecnai G2 F20 instrument (Philips Holland), while an energy-dispersive X-ray spectroscopy (EDX) analyzer on the Tecnai G2 F20 carried out the EDX spectrum and mapping analysis. The X-ray diffraction (XRD) patterns were performed on Rigaku D/max-rA (Rigaku, Japan) using Cu Kα radiation ($\lambda = 1.5418$ Å).

2.2. One-Pot Synthesis of PDDA-AuPtAg/RGO Nanohybrids

Firstly, 20 mg of GO was dispersed in 20 mL double distilled water containing 1% poly (diallyldimethylammonium chloride) (PDDA) solution followed by 2 h ultrasonication for the formation
of homogeneous PDDA–GO aqueous solution. Then, HAuCl₄ solution (100 µL, 1 M), H₂PtCl solution (100 µL, 1 M), and 10 mg AgNO₃ in 20 mL double distilled water were mixed with PDDA–GO aqueous solution. Then, 10 mL NaBH₄ solution was added dropwise under magnetic stirring at room temperature. With constant stirring lasting 24 h, the reaction was terminated until dark black products formed. Subsequently, PDDA-AuPtAg/RGO nanoalloy was obtained after centrifugation and washing by ethanol three times. Finally, the product was dried in a vacuum oven at 80 °C and prepared for use. RGO, PDDA-AuPt/RGO, PDDA-AuAg/RGO, and PDDA-PtAg/RGO were prepared through using a similar method.

2.3. Preparation of Modified Electrodes

The glassy carbon (GC) electrode was polished with 0.3 and 0.05 µm α-alumina powder before modification, and then washed by double distilled water and ethanol, respectively, followed by drying with high-purity nitrogen stream. Then, 10 mg of PDDA-AuPtAg/RGO was re-dispersed in 10 mL of double distilled water with sonication for 2 h, after which 6 µL of the suspension was immobilized on the surface of the GC electrode by dip-coating and dried in air.

2.4. Real-Time Detection of H₂O₂ Released From Living Cells

The cells tested in this work were SKOV3 cells (human ovarian cancer cell line), MCF-7 cells (breast cancer cell line), and A431 cells (epidermoid carcinoma cell line) obtained from School of Basic Medical Science, Tianjin Medical University, Tianjin, China (come from ATCC). SKOV3 cells and A431 cells were maintained in McCoy’s 5A (Modified) Medium and DMEM containing 10% fetal bovine serum, respectively. For MCF-7 cells, they were maintained in Dulbecco’s modified essential medium (DMEM) containing 10% fetal bovine serum, 0.01 mg/ml human recombinant insulin, 100 units/mL penicillin, and 100 µg/mL streptomycin. All the cells were incubated under ambient conditions with 37 °C, 5% CO₂, 95% humidified atmosphere, and sub-cultured after every second day.

After sub-cultured to 90% confluence, the cells were washed three times with phosphate-buffered saline (PBS) (0.1 M, pH 7.0) and collected by centrifugation. The cell number was calculated by using a cell counter. A total of 1×10⁶ cells were resuspended into 10 mL 0.1 M PBS (pH 7.0) as cell samples and the control group containing catalases (300 U/mL) was also prepared. For electrochemical measurements, injection of 10 µL PMA (1 µg mL⁻¹) was applied into the tested system and current responses on the PDDA-AuPtAg/RGO/GCE were recorded at the potential of 0.13 V, as shown in Scheme 1.

3. Results and Discussion

3.1. Characterization of PDDA-AuPtAg/RGO Nanocomposites

To identify the morphology of the as-prepared nanocomposites, transmission electron microscopy (TEM) was used for structural analysis. In Figure 1A, a typical wrinkled pattern of reduced graphene oxide sheets can be observed. Under different magnification (Figure 1B,C), it can be clearly seen that AuPtAg alloy nanoparticles (as pointed out in Figure 2B) distributed well on the PDDA–RGO sheets with less aggregation owing to excellent dispersing performance of poly dimethyl diallyl ammonium chloride (PDDA). The PDDA-AuPtAg alloy nanocrystals displayed paralleled lattice fringes along the same orientation (Figure 1D), indicating there were single AuPtAg nanoalloy particles locating on the PDDA-capped reduced graphene oxide layers.
A TEM elemental mapping analysis (Figure 2A–F) and energy dispersive X-ray (EDX) spectroscopy (Figure S2) were conducted to verify the components of the PDDA-AuPtAg/RGO composites. The results indicate that the obtained nanocomposites consisted of C, O, Au, Pt, and Ag. The C and O elements were evenly distributed as a film, while Au, Pt, and Ag were relatively concentrated, implying the formation of a trimetallic alloy. Figure 2G shows X-ray diffraction (XRD) spectra of GO, PDDA-Pt/RGO, PDDA-Ag/RGO, PDDA-AuAg/RGO, and PDDA-AuPtAg/RGO. GO presented a typical XRD diffraction pattern with a sharp peak at around 10° and a wider peak at 23.99°. The peak at 10° disappeared with the formation of the various nanocomposites, thus indicating that GO was reduced to RGO [38]. The diffraction peaks at around 39.8°, 46.3°, and 67.8° for PDDA-Pt/RGO were associated with the (111), (200), and (220) facets of the Pt nanoparticles, respectively [39,40]. In addition, peaks at 38.1°, 44.1°, 64.4°, and 77.3° for PDDA-Ag/RGO were attributed to the reflections on the (111), (200), (220), and (311) planes of Ag, respectively. Meanwhile, the characteristic peaks at 38.3°, 44.3°, 64.9°, and 77.69° for the bimetallic PDDA-AuAg/RGO were indexed to diffraction peaks of the (111), (200), (220), and (311) lattice planes of Au, respectively, according to Bragg’s reflection with no obvious diffraction peaks of silver, indicating that the surfaces of the Ag nanoparticles were replaced with Au, which led to the formation of a core–shell structure [41]. Compared with its counterparts, all of the peaks in the diffraction pattern for PDDA-AuPtAg/RGO were broader owing to the superposition of lattice planes, which suggests the formation of a trimetallic alloy composite.
AuPtAg/RGO were broader owing to the superposition of lattice planes, which suggests the formation of a trimetallic alloy composite.

Figure 2. TEM elemental mapping for (A) PDDA-AuPtAg/RGO and (B) C, (C) O, (D) Au, (E) Pt, and (F) Ag in the nanocomposite. (G) X-ray diffraction (XRD) spectra of (a) GO, (b) PDDA-Pt/RGO, (c) PDDA-Ag/RGO, (d) PDDA-AuAg/RGO, and (e) PDDA-AuPtAg/RGO.

3.2. Electrochemical Behavior of Obtained Materials

Electrochemical characterization using ferro/ferricyanide as the redox probe was accomplished in cyclic voltammograms (CVs) for a bare glassy carbon electrode (GCE), PDDA-AuAg/RGO, PDDA-PtAg/RGO, PDDA-AuPt/RGO, and PDDA-AuPtAg/RGO, modified GCE electrodes (Figure 3A). PDDA-AuPtAg/RGO/GCE revealed a pair of well-defined redox peaks at around +300 mV and +172 mV. Compared with bare GCE and the modified GCE electrodes with bimetal composites, an obvious increment in the peaks was observed for PDDA-AuPtAg/RGO/GCE owing to its fast electron-transfer kinetics. To further characterize the obtained material, microscopic electroactive areas were estimated according to the Randles–Sevcik equation [42]:

\[ I_p = 2.69 \times 10^5 \times A \times D^{1/2} \times n^{3/2} \times v^{1/2} \times c, \]  

(1)
were much lower, suggesting that the trimetallic alloy composite responded well to the reduction of peroxidase-like activity of single Au, Pt, and Ag nanoparticles and the synergistic effects after integration [43–45]. Besides this, PDDA–RGO nanosheets also play a compelling part in boosting the integration [43–45].

3.3. Electrochemical Response to H2O2

In addition, a comparison on the performance of our designed sensor with previously reported results of which are shown in Figure 3B. A distinctly higher response signals could be contributed to the outstanding peroxidase-like activity of single Au, Pt, and Ag nanoparticles and the synergistic effects after integration [43–45]. Besides this, PDDA–RGO nanosheets also play a compelling part in boosting the performance of the proposed electrode, offering more arching sites for trimetallic alloy nanocomposites.
as a supporting platform. Therefore, the electrode modified with PDDA-AuPtAg/RGO exhibited excellent catalytic activity toward the reduction of H$_2$O$_2$.

3.4. Performance of the Proposed Sensor

3.4.1. Optimization of the Experimental Variables

As shown in Figure 3C,D, typical CV was adopted to investigate the effects of the scan rate at PDDA-AuPtAg/RGO/GCE. It can be clearly observed that the oxidation and reduction currents for H$_2$O$_2$ increased linearly along with the rising scan rate. In addition, there was a clear linear relationship between the redox peak current and the square root of scan rate ($v^{1/2}$) using the regression equation Ip ($\mu$A) = $-111.17v$ (mV/s) + 147.84 ($R^2 = 0.997$), indicating a diffusion-controlled process [46]. Other experimental parameters including concentration and deposition volume of PDDA-AuPtAg/RGO and pH were also investigated to determine the optimal sensor response (Figure S2, S3). In summary, the PDDA-AuPtAg/RGO/GCE demonstrated the best response performance toward H$_2$O$_2$ under the following conditions: 6 $\mu$L of PDDA-AuPtAg/RGO (1 mg/mL), phosphate-buffered saline (PBS) at pH 7.0, and a scan rate of 50 mV/s.

3.4.2. Amperometric Response towards H$_2$O$_2$

The constructed sensor based on PDDA-AuPtAg/RGO was used to determine the H$_2$O$_2$ concentration via amperometric i-t curves under optimal conditions. Figure 4A shows the amperometric response toward different concentrations of H$_2$O$_2$. In pace with the rising concentration of H$_2$O$_2$, the current response increased gradually, with 95% of the steady-state current being achieved within 5 s. The calibration curve (Figure 4B) was fitted after five independent repetitive experiments and the PDDA-AuPtAg/RGO modified GCE revealed an exceptional linear relationship between current response and H$_2$O$_2$ concentration in two segments: from 0.05 $\mu$M to 1 mM and from 1 mM to 5.5 mM. As shown in the insert in Figure 4B, the corresponding linear equation is $I = -244.15C - 7.83$ ($R^2 = 0.9985$) with a sensitivity of 2838.95 $\mu$A·mM$^{-1}$·cm$^{-2}$ in the range from 0.05 $\mu$M to 1 mM. In the meantime, another linear relationship, $I = -66.18C - 184.12$ ($R^2 = 0.9978$), existed from 1 mM to 5.5 mM. The calculated detection limit for H$_2$O$_2$ was 1.2 nM at a signal-to-noise ratio of 3. These results suggest that the PDDA-AuPtAg/RGO fabricated sensor achieved a wide detection range and a low detection limit when determining H$_2$O$_2$ concentration, thus showing it to be a credible and powerful platform for monitoring cellular H$_2$O$_2$ levels.

In addition, a comparison on the performance of our designed sensor with previously reported H$_2$O$_2$ biosensors is summarized in Table S1, in which it can be clearly seen that the fabricated sensor based on PDDA-AuPtAg/RGO displayed noticeably improved properties, including a wider linear range and a lower detection limit for H$_2$O$_2$. These improvements could be attributed to the excellent synergetic effects of the trimetallic alloy and the larger working surface provided by the PDDA-capped RGO nanosheets.
synergetic effects of the trimetallic alloy and the larger working surface provided by the PDDA-capped RGO nanosheets.

Figure 4. (A) Amperometric i-t curves for PDDA-AuPtAg/RGO/GCE at 0.13 V (vs. Ag/AgCl) in 0.1 M PBS (pH 7.0) with successive additions of H$_2$O$_2$ (the insert shows a close-up of the low concentration of H$_2$O$_2$ region from 0.005 mM to 0.05 mM). (B) The corresponding calibration curve of response current vs. the concentration of H$_2$O$_2$.

3.4.3. Interference Immunity, Repeatability, and Stability

Specificity, reproducibility, and long-term stability are important indicators when evaluating the accuracy of the designed electrochemical sensors. An anti-interference study was performed with typical probes, including ascorbic acid, glucose, uric acid, and xanthine. As shown in Figure 5, variations compared with the current changes caused by H$_2$O$_2$ caused by the interference were almost non-existent (less than 2%), indicating that the constructed sensor had admirable selectivity and specificity.

To investigate the stability, the fabricated sensor was tested periodically. After storage at 4°C for one month, 92% of the initial current response of the as-prepared electrodes remained, illustrating acceptable long-term stability. Five electrodes were utilized under optimal conditions for exploring the reproducibility. As shown in Figure S4, favorable reproducibility can be observed with a relative standard deviation (RSD) of 2.45%.
was monitored with the as-prepared PDDA-AuPtAg which is of great importance in various biological processes, and many disorders like cancer result from the unbalanced metabolism of ROS [47]. The dynamic process of releasing \( \text{H}_2\text{O}_2 \) upon oxidative stress was monitored with the as-prepared PDDA-AuPtAg/RGO/GCE after adding phorbol myristate acetate (PMA) to SKOV3, MCF-7, and A431 cell lines. PMA is a well-known stimulus for triggering \( \text{H}_2\text{O}_2 \) production to mimic oxidative metabolism in vivo. After stimulation of PMA, respiratory burst will occur and \( \text{H}_2\text{O}_2 \) will be the end product released from cells through fusion pores [48,49]. Determining the cytotoxicity of the obtained materials PDDA-AuPtAg/RGO was completed by MTT assays before the electrochemical measurements. As shown in Figure 6A, MCF-7, SKOV3, and A431 cells maintained 95.1%, 95.8%, and 96.7% viability, respectively, after 12 h incubation with PDDA-AuPtAg/RGO, suggesting the favorable biocompatibility of the biosensor.

Figure 5. Amperometric i-t response of PDDA-AuPtAg/RGO/GCE upon the successive addition of \( \text{H}_2\text{O}_2 \) (0.1 mM), ascorbic acid (AA; 1 mM), glucose (1 mM), uric acid (UA; 1 mM), xanthine (1 mM), and \( \text{H}_2\text{O}_2 \) (0.1 mM) to 0.1 M PBS (pH 7.0) at an applied potential of 0.13 V with stirring.

3.5. In Situ Monitoring of \( \text{H}_2\text{O}_2 \) Released From Living Cells

The excellent performance of the PDDA-AuPtAg/RGO sensor provided the opportunity to explore its application in monitoring \( \text{H}_2\text{O}_2 \) in living cancer cells. It is well proven that \( \text{H}_2\text{O}_2 \) is a stable ROS, which is of great importance in various biological processes, and many disorders like cancer result from the unbalanced metabolism of ROS [47]. The dynamic process of releasing \( \text{H}_2\text{O}_2 \) upon oxidative stress was monitored with the as-prepared PDDA-AuPtAg/RGO/GCE after adding phorbol myristate acetate (PMA) to SKOV3, MCF-7, and A431 cell lines. PMA is a well-known stimulus for triggering \( \text{H}_2\text{O}_2 \) production to mimic oxidative metabolism in vivo. After stimulation of PMA, respiratory burst will occur and \( \text{H}_2\text{O}_2 \) will be the end product released from cells through fusion pores [48,49]. Determining the cytotoxicity of the obtained materials PDDA-AuPtAg/RGO was completed by MTT assays before the electrochemical measurements. As shown in Figure 6A, MCF-7, SKOV3, and A431 cells maintained 95.1%, 95.8%, and 96.7% viability, respectively, after 12 h incubation with PDDA-AuPtAg/RGO, suggesting the favorable biocompatibility of the biosensor.

Figure 6B exhibits amperometric response curves for PDDA-AuPtAg/RGO/GCE in the presence of SKOV3, MCF-7, and A431 cells triggered by the addition of PMA (10 µL, 1 µg mL\(^{-1}\)), after which the currents changed significantly, indicating \( \text{H}_2\text{O}_2 \) released from these cancerous cell lines was reduced on the PDDA-AuPtAg/RGO modified electrode in real-time. PBS buffer without cells and cells with catalase were used as the control. There was no current response in the absence of cells and the current remained almost constant in the presence of 300 U/ml catalase (an \( \text{H}_2\text{O}_2 \) scavenger) in the PBS buffer. These results reveal that the current responses can undoubtedly be ascribed to the reduction of \( \text{H}_2\text{O}_2 \) released from the cancerous cells upon oxidative stress [50]. After the stimulation by 10 µL PMA, the current change in the SKOV3 cells was around 8.27 µA, and according to the calibration curve in Figure 4B, the amount of \( \text{H}_2\text{O}_2 \) was calculated as ~1.98 µM. Similarly, the current changes for the MCF-7 and A431 cells were 8.36 and 8.4 µA, respectively, which correspond to ~2.21 and ~2.34 µM \( \text{H}_2\text{O}_2 \) released from the cells. To confirm the accuracy, an Amplex Red Hydrogen Peroxide Assay kit (Molecular Probes) was used according to the manufacturer’s instructions (Table S2), which revealed a relative deviation between the two methods of less than 2.5%. These results suggest the excellent practicality of the proposed biosensor and its huge potential for the investigation of the mechanisms in ROS-enhanced disorders.
4. Conclusions

An ultrasensitive electrochemical sensor based on PDDA-capped AuPtAg/RGO was designed for the dynamic monitoring of intracellular H₂O₂. The synergetic effects of the trimetallic alloy ensured exceptional electrochemical catalysis toward H₂O₂ and definitely accelerated electron transfer, while the PDDA–RGO nanosheets employed as a supporting framework effectively enlarged the active surface area. The proposed sensor demonstrated admirable performance toward H₂O₂ detection in the concentration range from 0.05 μM to 5.5 mM with a low detection limit of 1.2 nM (signal-to-noise ratio of 3). In addition, the developed sensor was capable of real-time H₂O₂ measurements in three cancerous cell lines, thereby indicating its huge potential for the in-depth study of the role of ROS in tumor formation and the screening of cancer at an early stage.

Supplementary Materials: The following are available online at http://www.mdpi.com/1424-8220/20/1/71/s1, Figure S1. EDX spectrum of PDDA-AuPtAg/RGO; Figure S2. Effects of concentration of PDDA-AuPtAg/RGO nanocomposite (0.3, 0.5, 0.8, 1.0, 1.5 mg/mL) and the deposition volume of PDDA-AuPtAg/RGO nanocomposite (3, 4, 5, 6, 7 μL) in 0.1 mM H₂O₂ and 0.1 M PBS (pH 7.0); Figure S3. The effect of pH on the reduction peak current of 5 mM H₂O₂ in a 0.1 M PBS at a scan rate of 50 mV/s; Figure S4. Column graph of CV signals of 0.1 mM H₂O₂ in 0.1 M PBS (pH 7.0) at five different electrodes prepared under the same conditions; Table S1 comparison of the reported sensors for H₂O₂ determination; Table S2. Comparison on detection of H₂O₂ released from living cells (1 × 10⁶) by different methods (n = 3).
**Author Contributions:** J.J. performed the experiments and analyzed the data, and also proposed the fabrication methods and wrote the manuscript; M.P. and X.L. performed the morphological analysis; B.L. and J.L. contributed reagents/materials/analysis tools. Q.C. managed the research and revised the manuscript. All authors have read and agreed to the published version of the manuscript.

**Funding:** This work was supported by the National Natural Science Foundation of China with Grant Nos. 81671779 and 81871451.

**Acknowledgments:** We are grateful to all laboratory members for their technical advice and helpful discussion and also thanks Li Zhao in School of Basic Medical Science, Tianjin Medical University, Tianjin, China for supporting us with cell lines.

**Conflicts of Interest:** The authors declare no conflict of interest.

**References**

1. Bray, F.; Ferlay, J.; Soerjomataram, I.; Siegel, R.L.; Torre, L.A.; Jemal, A. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J. Clin.* 2018, 68, 394–424. [CrossRef] [PubMed]

2. Wild, C.P.; Espina, C.; Bauld, L.; Bonanni, B.; Brenner, H.; Brown, K.; Dillner, J.; Forman, D.; Kampman, E.; Nilbert, M. Cancer Prevention Europe. *Mol. Oncol.* 2019, 13, 528–534. [CrossRef] [PubMed]

3. Hossain, M.; Luo, Y.; Sun, Z.; Wang, C.; Zhang, M.; Fu, H.; Qiao, Y.; Su, M. X-ray enabled detection and eradication of circulating tumor cells with nanoparticles. *Biosens. Bioelectron.* 2012, 38, 348–354. [CrossRef] [PubMed]

4. Ceci, F.; Castellucci, P.; Graziani, T.; Farolfi, A.; Fonti, C.; Lodi, F.; Fanti, S. 68 Ga-PSMA-11 PET/CT in recurrent prostate cancer: efficacy in different clinical stages of PSA failure after radical therapy. *Eur. J. Nucl. Med. Mol. Imaging* 2019, 46, 31–39. [CrossRef] [PubMed]

5. Tan, Y.; Gao, J.; Wu, C.; Zhao, S.; Yu, J.; Zhu, R.; Zhang, Q.; Wu, G.; Xue, X.; Wu, J. CT characteristics and pathologic basis of solitary cystic lung cancer. *Radiology* 2019, 291, 495–501. [CrossRef] [PubMed]

6. Mizutani, H.; Tada-Oikawa, S.; Hiraku, Y.; Kojima, M.; Kawanishi, S. Mechanism of apoptosis induced by doxorubicin through the generation of hydrogen peroxide. *Life Sci.* 2005, 76, 1439–1453. [CrossRef]

7. Gorrini, C.; Harris, I.S.; Mak, T.W. Modulation of oxidative stress as an anticancer strategy. *Nat. Rev. Drug Discovery* 2013, 12, 931. [CrossRef]

8. Wang, Y.; Zhang, F.; Miao, P.; Li, H.; Tu, Y. An Electrochemiluminescent Platform for Living Cell Oxygen Metabolism Monitoring. *J. Anal. Test.* 2018, 2, 184–189. [CrossRef]

9. Benhar, M.; Engelberg, D.; Levitzki, A. ROS, stress-activated kinases and stress signaling in cancer. *EMBO Rep.* 2002, 3, 420–425. [CrossRef]

10. Valko, M.; Rhodes, C.; Moncol, J.; Izakovic, M.; Mazur, M. Free radicals, metals and antioxidants in oxidative stress-induced cancer. *Chem. Biol. Interact.* 2006, 160, 1–40. [CrossRef]

11. Lennicke, C.; Rahn, J.; Lichtenfels, R.; Wessjohann, L.A.; Seliger, B. Hydrogen peroxide-production, fate and role in redox signaling of tumor cells. *Cell Commun. Signal.* 2015, 13, 39. [CrossRef] [PubMed]

12. Guo, Z.; Xu, J.; Zhang, J.; Hu, Y.; Pan, Y.; Miao, P. Facile Strategy for Electrochemical Analysis of Hydrogen Peroxide Based on Multifunctional Fe3O4@Ag Nanocomposites. *ACS Appl. Bio Mater.* 2018, 1, 367–373. [CrossRef]

13. Yamaguchi, S.; Kishikawa, N.; Ohyama, K.; Ohba, Y.; Kohno, M.; Masuda, T.; Takadate, A.; Nakashima, K.; Kuroda, N. Evaluation of chemiluminescence reagents for selective detection of reactive oxygen species. *Anal. Chim. Acta* 2010, 665, 74–78. [CrossRef] [PubMed]

14. Li, N.; Huang, J.; Wang, Q.; Gu, Y.; Wang, P. A reaction based one-and two-photon fluorescent probe for selective imaging H2O2 in living cells and tissues. *Sens. Actuators B Chem.* 2018, 254, 411–416. [CrossRef]

15. Bollenna, P.; Gorton, L. Enzyme based amperometric biosensors. *Curr. Opin. Electrochem.* 2018, 10, 157–173. [CrossRef]

16. Fan, Z.; Lin, Q.; Gong, P.; Liu, B.; Wang, J.; Yang, S. A new enzymatic immobilization carrier based on graphene capsule for hydrogen peroxide biosensors. *Electrochim. Acta* 2015, 151, 186–194. [CrossRef]

17. Zhu, C.; Yang, G.; Li, H.; Du, D.; Lin, Y. Electrochemical sensors and biosensors based on nanomaterials and nanostructures. *Anal. Chem.* 2014, 87, 230–249. [CrossRef]
18. Dai, H.; Lü, W.; Zuo, X.; Zhu, Q.; Pan, C.; Niu, X.; Liu, J.; Chen, H.; Chen, X. A novel biosensor based on boronic acid functionalized metal-organic frameworks for the determination of hydrogen peroxide released from living cells. Biosens. Bioelectron. 2017, 95, 131–137. [CrossRef]

19. Sun, Y.; Luo, M.; Meng, X.; Xiang, J.; Wang, L.; Ren, Q.; Guo, S. Graphene/intermetallic PtPb nanoplates composites for boosting electrochemical detection of $\text{H}_2\text{O}_2$ released from cells. Anal. Chem. 2017, 89, 3761–3767. [CrossRef]

20. Zhang, C.; Li, L.; Ju, J.; Chen, W. Electrochemical sensor based on graphene-supported tin oxide nanoclusters for nonenzymatic detection of hydrogen peroxide. Electrochim. Acta 2016, 210, 181–189. [CrossRef]

21. Bai, Z.; Li, G.; Liang, J.; Su, J.; Zhang, Y.; Chen, H.; Huang, Y.; Sui, W.; Zhao, Y. Non-enzymatic electrochemical biosensor based on Pt NPs/RGO-CS-Fc nano-hybrids for the detection of hydrogen peroxide in living cells. Biosens. Bioelectron. 2016, 82, 185–194. [CrossRef] [PubMed]

22. Palanisamy, S.; Lee, H.F.; Chen, S.-M.; Thirumalraj, B. An electrochemical facile fabrication of platinum nanoparticle decorated reduced graphene oxide; application for enhanced electrochemical sensing of H$_2$O$_2$. Rsc Adv. 2015, 5, 105567–105573. [CrossRef]

23. Karuppiah, C.; Palanisamy, S.; Chen, S.-M. An ultrahigh selective and sensitive enzyme-free hydrogen peroxide sensor based on palladium nanoparticles and nafion-modified electrode. Electrocatalysis 2014, 5, 177–185. [CrossRef]

24. Palanisamy, S.; Chen, S.-M.; Sarawathi, R. A novel nonenzymatic hydrogen peroxide sensor based on reduced graphene oxide/ZnO composite modified electrode. Sens. Actuators B Chem. 2012, 166, 372–377. [CrossRef]

25. He, W.; Wamer, W.; Xia, Q.; Yin, J.-J.; Fu, P.P. Enzyme-like activity of nanomaterials. J. Environ. Sci. Health Part C Environ. Carcinog. Ecotoxicol. Rev. 2014, 32, 186–211. [CrossRef] [PubMed]

26. Si, P.; Huang, Y.; Wang, T.; Ma, J. Nanomaterials for electrochemical non-enzymatic glucose biosensors. RSC Adv. 2013, 3, 3487–3502. [CrossRef]

27. Chen, T.-W.; Palanisamy, S.; Chen, S.-M. Non-enzymatic sensing of hydrogen peroxide using a glassy carbon electrode modified with a composite consisting of chitosan-encapsulated graphite and platinum nanoparticles. Microchim. Acta 2016, 183, 2861–2869. [CrossRef]

28. Chen, Y.; Li, Q.; Jiang, H.; Wang, X. Pt modified carbon fiber microelectrode for electrochemically catalytic reduction of hydrogen peroxide and its application in living cell H$_2$O$_2$ detection. J. Electroanal. Chem. 2016, 781, 233–237. [CrossRef]

29. Guler, M.; Turkoğlu, V.; Bulut, A.; Zahmakiran, M. Electrochemical sensing of hydrogen peroxide using Pd@Ag bimetallic nanoparticles decorated functionalized reduced graphene oxide. Electrochim. Acta 2018, 263, 118–126. [CrossRef]

30. Kang, J.-X.; Chen, T.-W.; Zhang, D.-F.; Guo, L. PtNiAu trimetallic nanoalloys enabled by a digestive-assisted process as highly efficient catalyst for hydrogen generation. Nano Energy 2016, 23, 145–152. [CrossRef]

31. Han, S.H.; Liu, H.M.; Chen, P.; Jiang, J.X.; Chen, Y. Porous trimetallic PtRhCu cubic nanoboxes for ethanol electrooxidation. Adv. Energy Mater. 2018, 8, 1801326. [CrossRef]

32. Singh, S.; Tuteja, S.K.; Sillu, D.; Deep, A.; Suri, C.R. Gold nanoparticle-reduced graphene oxide based electrochemical immunosensor for the cardiac biomarker myoglobin. Microchim. Acta 2016, 183, 1729–1738. [CrossRef]

33. Pasinszki, T.; Krebsz, M.; Tung, T.T.; Losic, D. Carbon nanomaterial based biosensors for non-invasive detection of cancer and disease biomarkers for clinical diagnosis. Sensors 2017, 17, 1919. [CrossRef] [PubMed]

34. Tung, T.T.; Nine, M.J.; Krebsz, M.; Pasinszki, T.; Coghlan, C.J.; Tran, D.N.; Losic, D. Recent advances in sensing applications of graphene assemblies and their composites. Adv. Funct. Mater. 2017, 27, 1702891. [CrossRef]

35. Liu, W.; Zhang, J.; Li, C.; Tang, L.; Zhang, Z.; Yang, M. A novel composite film derived from cysteic acid and PDDA-functionalized graphene: enhanced sensing material for electrochemical determination of metronidazole. Talanta 2013, 104, 204–211. [CrossRef] [PubMed]

36. Zhang, S.; Shao, Y.; Liao, H.; Engelhard, M.H.; Yin, C.; Lin, Y. Polyelectrolyte-induced reduction of exfoliated graphite oxide: A facile route to synthesis of soluble graphene nanosheets. ACS Nano 2011, 5, 1785–1791. [CrossRef]

37. Xiang, Y.; Banks, M.K.; Wu, R.; Xu, W.; Chen, S. Synthesis of thermo-sensitive PDDA-co-PNIPAM/graphene hybrid via electrostatic interactions and its thermal modulated phase transition. Mater. Chem. Phys. 2018, 220, 58–65. [CrossRef]
38. Hu, C.; Zhai, X.; Liu, L.; Zhao, Y.; Jiang, L.; Qu, L. Spontaneous reduction and assembly of graphene oxide into three-dimensional graphene network on arbitrary conductive substrates. Sci. Rep. 2013, 3, 2065. [CrossRef]
39. Ensafi, A.A.; Jafari-Asl, M.; Rezaei, B. A new strategy for the synthesis of 3-D Pt nanoparticles on reduced graphene oxide through surface functionalization, Application for methanol oxidation and oxygen reduction. Electrochim. Acta 2014, 130, 401–405. [CrossRef]
40. Guan, Y.; Dai, M.; Liu, T.; Liu, Y.; Liu, F.; Liang, X.; Suo, H.; Sun, P.; Lu, G. Effect of the dispersants on the performance of fuel cell type CO sensor with Pt–C/Nafion electrodes. Sens. Actuators B Chem. 2016, 230, 61–69. [CrossRef]
41. Wang, D.; Markus, J.; Kim, Y.-J.; Wang, C.; Pérez, Z.E.J.; Ahn, S.; Aceituno, V.C.; Mathiyalagan, R.; Yang, D.C. Coalescence of functional gold and monodisperse silver nanoparticles mediated by black Panax ginseng Meyer root extract. Int. J. Nanomed. 2016, 11, 6621. [CrossRef] [PubMed]
42. Ramanathan, M.; Patil, M.; Epur, R.; Yun, Y.; Shanov, V.; Schulz, M.; Heineman, W.R.; Datta, M.K.; Kumta, P.N. Gold-coated carbon nanotube electrode arrays: Immunosensors for impedimetric detection of bone biomarkers. Biosens. Bioelectron. 2016, 77, 580–588. [CrossRef] [PubMed]
43. Peng, X.; Wan, G.; Wu, L.; Zeng, M.; Lin, S.; Wang, G. Peroxidase-like activity of Au@ TiO2 yolk-shell nanostructure and its application for colorimetric detection of H$_2$O$_2$ and glucose. Sens. Actuators B Chem. 2018, 257, 166–177. [CrossRef]
44. Zhang, T.; Xing, Y.; Song, Y.; Gu, Y.; Yan, X.; Lu, N.; Liu, H.; Xu, Z.; Xu, H.; Zhang, Z. AuPt/MOF–Graphene: A Synergistic Catalyst with Surprisingly High Peroxidase-Like Activity and Its Application for H$_2$O$_2$ Detection. Anal. Chem. 2019, 91, 10589–10595. [CrossRef]
45. Jiang, H.; Chen, Z.; Cao, H.; Huang, Y. Peroxidase-like activity of chitosan stabilized silver nanoparticles for visual and colorimetric detection of glucose. Analyst 2012, 137, 5560–5564. [CrossRef]
46. Thanh, T.D.; Balamurugan, J.; Tuan, N.T.; Jeong, H.; Lee, S.H.; Kim, N.H.; Lee, J.H. Enhanced electrocatalytic performance of an ultrafine AuPt nanoalloy framework embedded in graphene towards epinephrine sensing. Biosens. Bioelectron. 2017, 89, 750–757. [CrossRef]
47. Khan, M.U.; Qurashi, N.A.; Khan, M.S.; Ijabeen, F.; Umar, A.; Yaqoob, J.; Wajid, M. Generation of reactive oxygen species and their impact on the health related parameters: A critical review. Int. J. Biosci. 2016, 9, 303–323.
48. Griffith, A.W.; Cooper, J.M. Single-cell measurements of human neutrophil activation using electrorotation. Anal. Chem. 1998, 70, 2607–2612. [CrossRef]
49. Wu, P.; Qian, Y.; Du, P.; Zhang, H.; Cai, C. Facile synthesis of nitrogen-doped graphene for measuring the releasing process of hydrogen peroxide from living cells. J. Mater. Chem. 2012, 22, 6402–6412. [CrossRef]
50. Dou, B.; Yang, J.; Yuan, R.; Xiang, Y. Trimetallic hybrid nanoflower-decorated MoS2 nanosheet sensor for direct in situ monitoring of H$_2$O$_2$ secreted from live cancer cells. Anal. Chem. 2018, 90, 5945–5950. [CrossRef]