Hollow-Channel Paper Analytical Devices Supported Biofuel Cell-Based Self-Powered Molecularly Imprinted Polymer Sensor for Pesticide Detection

Yanhu Wang 1,2,*, Huihui Shi 2, Jiantao Sun 3, Jianjian Xu 4, Mengchun Yang 1 and Jinghua Yu 2

1 Shandong Analysis and Test Center, Qilu University of Technology (Shandong Academy of Sciences), Jinan 250014, China
2 School of Chemistry and Chemical Engineering, University of Jinan, Jinan 250022, China
3 Shandong Institute for Product Quality Inspection, Jinan 250102, China
4 Department of Food and Drug, Weihai Ocean Vocational College, Weihai 264300, China
* Correspondence: chm_wangyh@qlu.edu.cn

Abstract: Herein, a paper-based glucose/air biofuel cell (BFC) was constructed and implemented for self-powered pesticide detection. Our developed paper-based chip relies on a hollow-channel to transport fluids rather than capillarity, which reduces analysis times as well as physical absorption. The gold nanoparticles (Au NPs) and carbon nanotubes (CNTs) were adapted to modify the paper fibers to fabricate the flexible conductive paper anode/cathode electrode (Au–PAE/CNT–PCE). Molecularly imprinted polymers (MIPs) using 2,4-dichlorophenoxyacetic acid (2,4-D) as a template were synthesized on Au–PAE for signal control. In the cathode, bilirubin oxidase (BOD) was used for the oxygen reduction reaction. Based on a competitive reaction between 2,4-D and glucose-oxidase-labeled 2,4-D (GOx-2,4-D), the amount of GOx immobilized on the bioanode can be simply tailored, thus a signal-off self-powered sensing platform was achieved for 2,4-D determination. Meanwhile, the coupling of the paper supercapacitor (PS) with the paper-based chip provides a simple route for signal amplification. Combined with a portable digital multi-meter detector, the amplified signal can be sensitively readout. Through rational design of the paper analytical device, the combination of BFC and PS provides a new prototype for constructing a low-cost, simple, portable, and sensitive self-powered biosensor lab-on-paper, which could be easily expanded in the field of clinical analysis and drug delivery.

Keywords: self-powered; biofuel cell; hollow-channel; paper-based analytical device; molecularly imprinted polymers

1. Introduction

Paper, as the most accessible medium for messages and knowledge transport throughout human history, has attracted considerable attention in the field of point-of-care testing (POCT) [1]. The distinct advantages of being abundant, low-cost, and easily disposable, enable paper to be a promising solution to develop biosensors with different detection techniques in resource-limited regions [2,3]. In recent years, microfluidics paper-based analytical devices (µ-PADs) for POCT have become increasingly popular and have attracted more and more interest, and various analytical methods have been established on µ-PADs for quantitative analysis [4]: such as colorimetric [5], fluorescence [6], electrochemical [7], chemiluminescent [8], electrochemiluminescent [9], and photoelectrochemical [10,11]. Although, the developed µ-PADs still featured the advantages of paper, most of them relied on the capillary action of porous paper channels to transport fluids that was time-consuming and allowed physical absorption: therefore, restricting its popularization [12,13]. Recently, the emergence of hollow–channels through removing the cellulose matrix opens up new insights to improve mass transfer efficiency and alter physical absorption, which is beneficial to their popularization in the field of POCT [14–16].
Pesticides have been widely used in the agricultural industry to improve crop yields [17]. However, the abuse of pesticides has also brought great detriment to the ecosystem and human health due to their toxicity [18]. Especially, 2,4-dichlorophenoxyacetic acid (2,4-D) has been widely utilized to eliminate broadleaf weeds in the field crops. Their structural stability makes them resistant to degradation and more likely to accumulate, which may lead to cancer in humans, and endocrine-disrupting issues [19]. Therefore, it is desirable to develop a simple and sensitive method to realize 2,4-D monitoring. Until now, the common methods for 2,4-D assays have been gas chromatography–mass spectrometry and liquid chromatography–mass spectrometry [20,21]. However, the time-consuming procedures, sophisticated instruments, and limited selectivity have restricted their extensive application. As synthesized artificial receptor molecules, molecularly imprinted polymers (MIPs) possess instinctive mechanical/chemical stability, a low-cost that has distinguished MIPs from other biometric recognition molecules, and they have attracted considerable attention among the various biosensors [22–29].

The introduction of MIPs into µ-PADs provides a new analytical approach for pesticide detection [30]. However, most methods fabricated on µ-PADs such as electrochemical and electrochemiluminescence, usually need to be excited by external energy sources causing them to depart from on-site monitoring [31,32]. Considering the stated problems, great efforts have been devoted to exploit portable power sources that simplify complicated operation [33]. Great achievements have been made, but there are still great improvements possible. Biofuel cells (BFCs), harvest electric from chemical energy, representing a new type of green energy conversion device based on biocatalysts [34,35]. Furthermore, the mild operation condition, miniaturization and easy integration of energy conversion with analytes offer a promising approach for self-powered on-site analysis [36]. Various self-powered biosensors based on BFCs have been fabricated and successfully implemented in the field of diagnosis and environmental monitoring [37–39].

Despite great progress having been achieved, most of the BFC-based self-powered sensors rely on sophisticated electrochemical workstations for signal collection that departs from the development of portable, integration and low-cost systems. Fortunately, coupling BFCs with capacitors provides an effective way to enable high power charge/discharge cycles as well as for the output signal amplification [40,41]. Based on this, we demonstrated a self-powered biosensor that integrated glucose/O\(_2\) BFC and an all-solid-state paper supercapacitor (PS) within a signal piece of paper based on origami for 2,4-D detection (Scheme 1). The fabricated BFC is composed of a glucose oxidase (GOx) bioanode and a bilirubin oxidase (BOD) biocathode. MIPs grafted on the bioanode of BFC through electropolymerization of polypyrrole (PPy) acted as biological receptors for specific recognition of 2,4-D. The competition between free 2,4-D and GOx labeled 2,4-D (Gox–2,4-D), further influence the electric output of BFC. The generated current was temporarily collected and stored within the PS realizing an amplified current output detected by a digital multimeter (DMM). On the other hand, hollow channels were designed on the paper device to eliminate the barriers regarding low fluid flow rates and significant nonspecific adsorption. Overall, all these features accounted for excellent analytical performance toward 2,4-D. Our developed method paves a new way for developing integrated, portable, and versatile paper-based analytical devices, as well as provides a new choice for diagnosis and environmental monitoring in resource-limited regions.
2. Experimental Section

2.1. Fabrication of the Origami Biofuel Cell-Based Lab-on-Paper Device (BFCPD)

In this work, the sketch of the origami biofuel cell-based lab-on-paper device (BFCPD) was designed on Adobe illustrator CS6. Different functional zones were differentiated by color. The configuration of the constructed BFCPD comprised an anodic tab (blue), a hollow-channel tab, a cathodic tab (dark blue), a hemichannel tab (green), and two paper supercapacitor tabs (Figure S1). The square hydrophilic zone (2.5 × 2.5 mm) on the hollow-channel tab and cathodic tab, named as the “via hole”, was modified by the Au NPs’ decorated multiwalled carbon nanotubes (Au–MWCNTs) that enable the “via hole” electrical conduction from bottom to top. Each tab was folded based on the unprinted line. The unprinted rectangle region was used for the fabrication of PS (Figure S2). After wax printing, the obtained paper sheet was baked at 130 °C until the wax melted and penetrated through the paper to form the hydrophobic and insulating patterns (Figures S3 and S4). Subsequently, the paper sheet was ready for carbon electrodes and the silver conductive wires screen-printing (Figures S5 and S6). A light yellow hemichannel on the hemichannel tab was obtained through varying the values of C, M, Y, and K. The hydrophilic paper fibers on the cathodic tab and the hollow-channel tab and reservoirs were removed. Finally, the PS was fabricated based on reported work [3]. Specifically, the hydrophilic zone of the anodic/cathodic tab (part a in Scheme S1A,B) was functionalized with Au nanoparticles and ionic liquid CNTs endowing the paper electrode with superior conductivity to fabricate Au–PAE (part b in Scheme S1A) and CNT-PCE (part b in Scheme S1B). The PS was fabricated based on the reported work [42,43].

2.2. Fabrication of Bioanode

The MIPs grafted on Au-PAE were fabricated through electropolymerization of polypyrrole (PPy) MIPs with the presence of the template molecule (2,4-D) (part c in Scheme S1A). An electrolyte solution containing 0.1 M KCl, 0.5 mM pyrrole, and 10 mM 2,4-D in 0.05 M phosphate buffer solution (PBS, pH 7.0) was first prepared and deoxygenized in bubbling nitrogen gas for 10 min. Subsequently, the electropolymerization was performed at a constant voltage of 0.8 V (vs. Ag/AgCl). After a 10-min reaction, the
template molecules were removed and rinsed through a methanol and acetic acid mixture (70% methanol, v/v). Afterwards, the obtained MIP–Au–PAE was rinsed with ultrapure water and dried in an air stream. Meanwhile, the non-imprinted polymers (NIPs)-grafted Au–PAE was prepared with the same procedure except for the introduction of template molecules into the non-imprinted polymers (NIPs)-grafted Au–PAE.

2.3. Fabrication of Biocathode

This process involved 10 µL of BOD (10 µg·mL⁻¹, dissolved in 0.1 M pH 7.4 PBS) being cast onto the CNT-PCE and maintained for 2 h at room temperature (part c in Scheme S1B). Subsequently, physically absorbed molecules were removed through washing with PBS (0.01 M, pH 7.4). The resulting biocathodes (BOD–CNT–PCE) were stored at 4 °C prior to use.

2.4. Assay Procedure

Prior to assay, the designed BFCPD was first folded as indicated in Figure S7, and then clamped by a device-holder guaranteeing that the BFCPD was stacked closely. After that, 50 µL of a 10 mM 2,4-D solution was introduced into the BFCPD through the inlet, and the fluid transported along the designed hollow channel finally reached the bioanode. After a 25-min incubation, the recognition cavities on the MIPs were fully blocked. Excess solution was removed from the inlet. Subsequently, 50 µL of 10 mM GOx-2,4-D solution was added and incubated for another 25 min. During this process, the GOx-2,4-D occupied the binding sites leading to replacement of 2,4-D (part d in Scheme S1A). Part of the 2,4-D could not be replaced due to the stereoscopic hindrance effect from GOx. To assess the sample detection, a 30 µL aqueous solution with different concentrations of 2,4-D was added and incubated for 25 min. A competitive reaction occurred, the GOx2,4-D was replaced by the free 2,4-D (part e in Scheme S1A). After washing with PBS, 50 µL of 30 mM glucose solution was added and migrated to the bioanode and biocathode to initiate the reaction. The generated current was collected and stored within the PS temporarily. After 60 s charging, a high current intensity could be detected by the DMM. Based on the amplified current intensity, the 2,4-D could be quantified.

3. Results and Discussion

3.1. Morphology Characterization

Figure 1A,B shows the scanning electron microscope (SEM) images of the bare paper electrode which possesses porous architecture and a rough surface. Compared with the bare paper, continuous and dense AuNPs assembled on the fibers can be seen (Figure 1C), and the paper still maintains the original porous microstructure (Figure 1D). The SEM image in Figure 1E,F presents well-dispersed CNTs on the surface of the paper fibers, also demonstrating the successful fabrication of CNT–PCE. The decoration of AuNPs and CNT would improve the conductivity that accounted for the superior performance of BFC. To investigate the morphology of the fabricated MIPs–Au–PAE, the SEM images were derived and the results are shown in Figure 1G,H. Compared with Au–PAE (shown in Figure 1C,D), the surface of fibers is rougher after the decoration of PPy MIPs, and the diameter increases with the electro-polymerized of PPy MIPs, which also illustrates a successful decoration of PPy MIPs on the surface of Au-paper fibers.
These results verified the successful construction of the bioanode. The results obtained versus Ag/AgCl. A low glucose oxidation potential of around 0.19 V (vs. Ag/AgCl) was investigated through cyclic voltammetry (CV) toward a 10.0 mM \([\text{Fe(CN)}_6]^{3–/4–}\) solution containing 0.5 M KCl. As shown in Figure 2A, compared with the bare PAE (curve a), a higher current response could be observed after the coating of a Au layer on paper fibers (curve b), which was attributed to the factor that the Au layer could greatly increase the conductivity. After the MIPs were grafted on the Au–PAE (curve c), a remarkable decrease in the peak current was detected due to the reduced conductivity of MIPs layers that acted as a definite kinetic barrier for the charge transfer. The current increased accordingly after the removal of the template (curve d), which suggested that the \([\text{Fe(CN)}_6]^{3–/4–}\) molecules were more easily available for the electrode. After template rebinding, the current decreased again (curve e). This may be caused by the molecules rebinding having blocked the diffusion of \([\text{Fe(CN)}_6]^{3–/4–}\) to the surface of the electrode. These results verified the successful construction of the bioanode. The results obtained from the electrochemical impedance spectroscopy (EIS) (Figure 2B) also supported the above results.

Meanwhile, CV (Figure 2C) and EIS (Figure 2D) were also selected to explain the fabrication procedure of the biocathode. Specifically, the CV response of different electrodes in the sequence of PCE < BOD–CNT–PCE < CNT–PCE, implied the successful construction of the BOD–CNT–PCE. Moreover, the variation of the semicircle in EIS (Figure 2D) also reflected the successful fabrication of the biocathode.

Electrochemical measurements were carried out to evaluate the performance of fabricated bioanode and biocathode. Figure 3A shows that the bioelectrocatalytic current resulted from the enzymatic oxidation of glucose within a voltage range from −0.6 to 0.4 V versus Ag/AgCl. A low glucose oxidation potential of around 0.19 V (vs. Ag/AgCl) was observed, which is consistent with the fact that the GOx comprises two subunits that each include the flavin adenine dinucleotide (FAD) cofactor [44]. Figure 3B displays the polarization curves of the prepared bioanode (incubation with 100 ng mL−1 2,4-D) in the presence of 30 mM glucose. With the potential scanning from −0.6 V to 0.8 V, the anodic currents increased correspondingly. It should be observed that an obvious catalytic electrooxidation potential at −0.2 V (vs. Ag/AgCl) for glucose was observed, and reached a plateau current density of about 10 mA cm−2 near −0.1 V (vs. Ag/AgCl) (curve b) compared to the bare electrolyte (curve a). This result reveals the high electrocatalytic activity of GOx toward glucose oxidation at the MIP-Au-PAE.
3.4. Optimization of Experimental Conditions

The optimized absorption time in this work was investigated. As shown in Figure S8A, the residual 2,4-D in solution decreased with the absorption time. Similarly, based on the results in Figure S8B, 25 min was also used as the optimal GOx–2,4-D labeling time and the competitive reaction time between GOx–2,4-D and 2,4-D.

Figure 2. CV (A) and EIS (B) responses of (a) PAE, (b) Au–PAE, (c) MIPsAu–PAE, (d) MIPs–Au–PAE after template removal, (e) MIPs–Au–PAE after template rebinding; CV (C) and EIS (D) responses of PCE (a), CNT–PCE (b), and BOD–CNT–PCE (c).

Figure 3. (A) CVs and (B) polarization curves for the bioanode in 0.10 M PBS (pH 7.4) with (curve a) and without (curve b) 30 mM glucose; (C) CVs and (D) polarization curves for the biocathode in 0.10 M PBS (pH 7.4) under N₂-atmosphere (curve a) and O₂-atmosphere (curve b). The scan rate for polarization curves is 1 mV·s⁻¹.
The performance of the fabricated biocathode was also studied in 0.1 M PBS (pH 7.4, containing 0.1 M NaCl) saturated by O₂ or N₂ (Figure 3C). From this comparison, the onset potential around 0.5 V for O₂ reduction is similar to the thermodynamic equilibrium potential of E°O₂/H₂O (0.61 V at pH 7.0) with the absence of O₂ (curve b) [34]. However, there was no peak observed for BOD–CNT–PCE in N₂-saturated solution (curve a). Figure 3D shows the polarization curves of BOD–CNT–PCE in a N₂ (curve a) -saturated solution, and an O₂ (curve b) -saturated solution. It could be clearly seen that the BOD–CNT–PCE displays a higher current density in an O₂-saturated surrounding than that in N₂-saturated condition. The above results clearly demonstrate the superior performance of the BOD–CNT–PCE toward oxygen reduction reaction.

3.3. General Working Principle of the Assembled BFC

The general working principle of the assembled BFC is illustrated in Scheme 1. Briefly, the GOx fix on the MIP–Au–PAE could efficiently catalyze the oxidation of glucose selectively to generate gluconolactone. At the same time, the FAD within GOx was reduced to GOx–FADH₂ [45]. The GOx–FADH₂ automatically released two H⁺ and electrons, and regenerated to GOx–FAD smoothly, leading to the efficient electron transfer from GOx to Au–PAE and the electron flow from bioanode to biocathode through the external circuit. The O₂ molecules are reduced to H₂O under a four proton-assisted electron transfer process at the biocathode with the participation of H⁺, eventually realizing the energy conversion from chemical energy to electric energy. The whole process can be illustrated in the following reaction equations:

\[
\text{GOx} - \text{FAD} + \text{glucose} \rightarrow \text{GOx} - \text{FADH}_2 + \text{gluconolactone}
\]

\[
\text{GOx} - \text{FADH}_2 \rightarrow \text{GOx} - \text{FAD} + 2H^+ + 2e^-
\]

\[
O_2 + 4H^+ + 4e^- \rightarrow 2H_2O
\]

3.4. Optimization of Experimental Conditions

In order to obtain optimal analytical performance and efficiency, the incubation time was optimized. First, the absorption time between the eluted MIPs–Au–PAE and 2,4-D was investigated. As shown in Figure S8A, the residual 2,4-D in solution decreased quickly with extending the absorption time, and reached a plateau after 25 min, indicating the equilibrium rebinding between the MIPs and the template. Therefore, 25 min was selected as the optimal absorption time in this work. Similarly based on the results in Figure S8B, 25 min was also used as the optimal GOx–2,4-D labeling time and the competitive reaction time between GOx–2,4-D and 2,4-D.

3.5. Analytical Performance

Under optimal conditions, the quantitative analysis and dynamic range of the developed BFCPD toward 2,4-D were evaluated by varying 2,4-D concentration in a standard solution. As expected, good linear range relationships between the detected current without (Figure 4A) or with (Figure 4B) the PS amplification and the logarithmic concentration of 2,4-D was observed in the range from 1.0 pM-50.0 µM. The fitted linear equations were \( I = 74.20 - 9.68 \log [\text{2,4-D/pM}] \) (\( R = 0.9946 \)) and \( I_{PS} = 949.22 - 122.18 \log [\text{2,4-D/pM}] \) (\( R^2 = 0.985 \)). The limit of detection (LOD) was estimated to be 0.53 pM based on the signal-to-noise (S/N) ratio of 3. It is interesting to see that the LOD obtained with or without PS are the same, which might be mostly attributed to the concomitant amplification of the background. Moreover, it should be observed that the instantaneous amplified current discharged from PS is about 13-fold higher than that without PS charging. The amplified current could be detected through the DMM, causing the complicated electrochemical workstation to be abandoned.

Subsequently, the power density of the assembled glucose/O₂ BFC after incubation with 1.0 µM 2,4-D in 0.10 M PBS (pH 7.0) containing 30 mM glucose was measured. At the
specified conditions, the fabricated BFC exhibited an open-circuit voltage (Voc) of 0.75 V, and the maximum power density ($P_{\text{max}}$) reached 152 mW cm$^{-2}$ at 0.5 V.

![Graphs showing the relationship between current response and 2,4-D concentration, power density on cell voltage, and selectivity investigation.](image)

**Figure 4.** The relationship between the current response and 2,4-D concentration without (A) and with (B) PS amplification; (C) dependence of the power density on the cell voltage in 0.10 M air-saturated PBS with 30 mM glucose; (D) selectivity investigation of the proposed method.

### 3.6. Specificity, Reproducibility, and Stability Investigation

Selectivity is a crucial standard to evaluate the performance of fabricated biosensors. Thus, 2,4,5-trichlorophenoxyacetic acid (2,4,5-T), 4-chlorophenoxyacetic acid (CPA), 2,4-dichlorophenol (2,4-DCP), 4-(2,4-dichlorophenoxy) butyric acid (2,4-DBA), and 4-chloro-o-tolyloxyacetic acid (MCPA) were selected as interferents to access the specificity of our developed BFCPD. As shown in Figure 4D, those interferents could barely influence the signal output and were similar to the blank. Only in the presence of 2,4-D, could a dramatically decreased current be detected. Those results demonstrated that the constructed biosensor possessed satisfied selectivity. Additionally, the reproducibility was investigated through intra- and inter-assays under the same conditions. An identical signal response with the relative standard deviation (RSD) of 3.4% and 5.2% indicated a gratifying reproducibility. Furthermore, the stability of the developed BFCPD was evaluated through measuring the current intensity at intervals of five days. No significant changes of the current output were observed when stored in a refrigerator at 4 °C for four weeks, revealing the good stability of the developed BFCPD.

### 3.7. Practical Application

Finally, the recovery test was conducted to evaluate the feasibility of the fabricated biosensor toward a practical application. The standard addition method was employed by adding different concentrations of 2,4-D into tap water and lake water, respectively. As shown in Table 1, the recovery values of the constructed biosensor were obtained in the range of 98.3-104% for tap water, and 97.5-103.5% for lake water. Together with the RSD
less than 3.2% and 4.3%, this validated the reliability and practicability of our developed BFCPD for 2,4-D determination in real samples.

Table 1. Practical determination of 2,4-D in real samples.

|                      | Tap Water   | Lake Water  |
|----------------------|-------------|-------------|
| Add a                | Detected b  | RSD (%, n = 11) c | Recovery (%) d |
| 1 10 pM              | 9.87 pM     | 2.6         | 98.7          |
| 2 500 pM             | 519.4 pM    | 3.1         | 103.9         |
| 3 100 nM             | 103.2 nM    | 2.8         | 103.2         |
| 4 500 nM             | 491.63 nM   | 2.5         | 98.33         |
| 5 10 µM              | 10.27 µM    | 3.2         | 102.7         |

[a] [Added] means the values that we add into water sample. [b] [Detected] means the amount of 2,4-D got according to the standard curve equations from eleven parallel detections. [c] The relative standard deviation (RSD) of measurements are calculated from eleven independent experiments. [d] Recovery means the ratio of [Detected]/[Added].

4. Conclusions

In this work, a sensitive paper-based self-powered sensing protocol based on a glucose/O2 BFC device integrated with a molecularly imprinted technique for 2,4-D detection was fabricated for the first time. A PS constructed on the fabricated BFCPD and a DMM were used as the current amplifier and the terminal current detector, respectively. Meanwhile, hollow channels were introduced into the BFCPD to transport fluids which could make them suitable for point-of-care testing. The construction of the conductivity of PPy MIPs layers offers a promising approach for 2,4-D-specific recognition. This is based on a competitive reaction between 2,4-D and GOx–2.4-D, and the influence of the catalytic oxidation of glucose at the bioanode to realize quantification. Meanwhile, the introduction of PS could collect and store the generated electrons leading to an amplified current response that can be directly read by the portable DMM. This work provides a novel and efficient approach to develop a multifunctional paper device to implement self-powered sensing, that eliminates the energy source. Such findings also provide an alternative choice for the design and development of POCT devices.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/bios12110974/s1, Figure S1. Schematic representation of this 3D-µ-MOBFCAD. (A) Wax patterns of this 3D-µ-MOBFCAD; (B) the reverse side of (A); (C) Right side of 3D-µ-MOBFCAD the after screen-printing of silver wire and carbon electrode; (D) the reverse side of (C). (a) cathodic tab; (b) anodic tab; (c) hollow-channel tab; (d) hemichannel tab; (e) unprinted paper for paper supercapacitor. (1) cathodic zone; (2) sample inlet; (3) anodic zone; (4) hollow-channel; (5) hemichannel; (6) via hole; (7) silver pad; (8) screen-printed cathodic electrode (SPCE); (9) screen-printed anodic electrode (SPAE); (10) silver wire; (11) paper supercapacitor. (I, II) hollow area; Figure S2. Wax-patterns of 3D-µ-MOBFCAD on a paper sheet (A4) before baking; Figure S3. Right side of the baked wax-patterns of 3D-µ-MOBFCAD on a paper sheet (A4) after dropping CNT/Au on via hole; Figure S4. Reverse side of the baked wax-patterns of 3D-µ-MOBFCAD on a paper sheet (A4) after dropping CNT/Au on via hole; Figure S5. Right side of the 3D-µ-OBFCAD on a paper sheet (A4) after screen-printing of carbon electrodes, silver wire and drawing of graphite electrodes; Figure S6. Reverse side of the 3D-µ-OBFCAD on a paper sheet (A4) after screen-printing of carbon electrodes, silver wire and drawing of graphite electrodes; Figure S7. Schematic Representation, Size, Shape and Folding Procedure of the Wax Patterns on This 3D-µ-OBFCAD; Figure S8. (A) Optimization conditions of blocking time in eluted MIP-grafted Au-PAE; (B) effects of (a) labeling time and (b) competition time in MIP-grafted Au-PAE; Scheme S1. Schematic diagram of the fabrication of
bioanode (A) and biocathode (B), different part in (A) represents PAE (a), Au-PAE (b), MIPs-Au-PAE (c), GOx-2,4-D occupied MIPs-Au-PAE (d), and detection of 2,4-D at MIPs-Au-PAE; different part in (B) represents PCE (a), CNT-PCE (b), and BOD-CNT-PCE (c).

**Author Contributions:** Y.W.: Conceptualization, Methodology, Investigation, Writing–review and editing, Project administration, Funding acquisition. H.S.: Formal analysis, Writing—original draft, Methodology, Investigation. J.S.: Characterizations, Validation, Investigation. J.X.: Characterizations, Validation, Investigation. M.Y.: Writing–review and editing. J.Y.: Writing–review and editing, Supervision, Funding acquisition, Investigation. All authors have read and agreed to the published version of the manuscript.

**Funding:** This work was financially supported by the National Natural Science Foundation of China (22004075), Natural Science Foundation of Shandong Province (ZR2020QB091), Excellent Youth Innovation Team in Universities of Shandong (2019JC016), Case-by-Case Project for Top Outstanding Talents of Jinan, and Science, Education and Industry Integration Innovation Pilot Project from Qilu University of Technology (Shandong Academy of Sciences) (2022JBZ02-04).

**Institutional Review Board Statement:** Not applicable.

**Informed Consent Statement:** Not applicable.

**Data Availability Statement:** Not applicable.

**Conflicts of Interest:** The authors declare that they have no known competing financial interests or personal relationships that could influence the work reported in this article.

**References**

1. Gerbers, R.; Foellacher, W.; Chen, H.; Anagnostopoulo, C.; Faghri, M. A new paper-based platform technology for point-of-care diagnostics. *Lab Chip* **2014**, *14*, 4042–4049. [CrossRef] [PubMed]
2. Yamada, K.; Henares, T.; Suzuki, K.; Citterio, D. Paper-based Inkjet-Printed Microfluidic Analytical Devices. *Angew. Chem. Int. Ed.* **2015**, *54*, 5294–5310. [CrossRef] [PubMed]
3. Wang, Y.; Ge, L.; Wang, P.; Yan, M.; Ge, S.; Li, N.; Yu, J.; Huang, J. Photoelectrochemical lab-on-paper device equipped with a porous Au-paper electrode and fluidic delay-switch for sensitive detection of DNA hybridization. *Lab Chip* **2013**, *13*, 3945–3955. [CrossRef] [PubMed]
4. Yetisen, A.K.; Akram, M.S.; Lowe, C.R. Paper-based microfluidic point-of-care diagnostic devices. *Lab Chip* **2013**, *13*, 2210–2251. [CrossRef]
5. Sun, Y.; Zhao, C.; Niu, J.; Ren, J.; Qu, X. Colorimetric band-aids for point-of-care sensing and treating bacterial infection. *ACS Central Sci.* **2020**, *6*, 207–212. [CrossRef]
6. Hui, C.Y.; Liu, M.; Li, Y.; Brennan, J.D. A Paper Sensor Printed with Multifunctional Bio/Nano Materials. *Angew. Chem. Int. Edit.* **2018**, *57*, 4549–4553. [CrossRef]
7. Liu, H.; Xiang, Y.; Lu, Y.; Crooks, R.M. Aptamer-based origami paper analytical device for electrochemical detection of adenosine. *Angew. Chem. Int. Ed.* **2012**, *51*, 6925–6928. [CrossRef]
8. Zong, C.; Wu, J.; Liu, M.; Yang, L.; Yan, F.; Ju, H. Chemiluminescence Imaging for a Protein Assay via Proximity-Dependent DNAzyme Formation. *Anal. Chem.* **2014**, *86*, 9939–9944. [CrossRef]
9. Huang, Y.; Li, L.; Zhang, Y.; Zhang, L.; Ge, S.; Li, H.; Yu, J. Cerium dioxide-mediated signal “On-Off” by resonance energy transfer on a lab-on-paper device for ultrasensitive detection of lead ions. *ACS Appl. Mater. Interfaces* **2017**, *9*, 32591–32598. [CrossRef]
10. Gao, C.; Yu, H.; Zhang, L.; Zhao, Y.; Xie, J.; Li, C.; Cui, K.; Yu, J. Ultrasensitive paper-based photoelectrochemical sensing platform enabled by the polar charge carriers-created electric field. *Anal. Chem.* **2020**, *92*, 2902–2906. [CrossRef]
11. Wang, Y.; Zhang, L.; Kong, Q.; Ge, S.; Yu, J. Time-resolution addressable photoelectrochemical strategy based on hollow-channel paper analytical devices. *Biosens. Bioelectron.* **2018**, *120*, 64–70. [CrossRef] [PubMed]
12. Glavan, A.C.; Martinez, R.V.; Maxwell, E.J.; Subramaniam, A.B.; Nunes, R.M.D.; Soh, S.; Whitesides, G.M. Rapid fabrication of pressure-driven open-channel microfluidic devices in omniphobic R-F paper. *Lab Chip* **2013**, *13*, 2922–2930. [CrossRef] [PubMed]
13. Li, X.; Scida, K.; Crooks, R.M. Detection of hepatitis B virus DNA with a paper electrochemical sensor. *Anal. Chem.* **2015**, *87*, 9009–9015. [CrossRef] [PubMed]
14. Liu, W.; Cassano, C.L.; Xu, X.; Fan, Z.H. Laminated paper-based analytical devices (LPAD) with origami-enabled chemiluminescence immunosassay for cotinine detection in mouse serum. *Anal. Chem.* **2013**, *85*, 10270–10276. [CrossRef] [PubMed]
15. Renault, C.; Koehne, J.; Ricco, A.J.; Crooks, R.M. Three-dimensional wax patterning of paper fluidic devices. *Langmuir* **2014**, *30*, 7030–7036. [CrossRef]
16. Giokas, D.L.; Tsogas, G.Z.; Vlessidis, A.G. Programming fluid transport in paper-based microfluidic devices using razor-crafted open channels. *Anal. Chem.* **2014**, *86*, 6202–6207. [CrossRef] [PubMed]
17. Wang, Y.; Zang, D.; Ge, S.; Ge, L.; Yu, J.; Yan, M. A novel microfluidic origami photoelectrochemical sensor based on CdTe quantum dots modified molecularly imprinted polymer and its highly selective detection of S-fenvalerate. *Electrochim. Acta* **2013**, *107*, 147–154. [CrossRef]

18. Luo, L.; Ou, Y.; Yang, Y.; Liu, G.; Liang, Q.; Ai, X.; Yang, S.; Nian, Y.; Su, L.; Wang, J. Rational construction of a robust metal-organic framework nanozyme with dual-metal active sites for colorimetric detection of organophosphorus pesticides. *J. Hazard. Mater.* **2022**, *423*, 127253. [CrossRef]

19. Wang, S.; Ge, L.; Li, L.; Yan, M.; Ge, S.; Yu, J. Molecularly imprinted polymer grafted paper-based multi-disk micro-disk plate for chemiluminescence detection of pesticide. * Biosens. Bioelectron.* **2013**, *50*, 262–268. [CrossRef]

20. Cai, Y.; Zhu, H.; Zhou, W.; Qiu, Z.; Chen, C.; Qi, L.; Li, K.; Liu, Y. Capsulation of AuNCs with AIE Effect into Metal-Organic Framework for the Marriage of a Fluorescence and Colorimetric Biosensor to Detect Organophosphorus Pesticides. *Anal. Chem.* **2021**, *93*, 7275–7282. [CrossRef]

21. Ding, L.; Hong, H.; Xiao, L.; Hu, Q.; Zuo, Y.; Hao, N.; Wei, J.; Wang, K. Nanoparticles-doped induced defective ZIF-8 as the novel cathodic luminoaphore for fabricating high-performance electrochemiluminescence aptasensor for detection of omethoate. *Biosens. Bioelectron.* **2021**, *192*, 113492. [CrossRef] [PubMed]

22. Schirhagl, R. Bioapplications for molecularly imprinted polymers. *Anal. Chem.* **2014**, *86*, 250–261. [CrossRef] [PubMed]

23. Yan, K.; Yang, Y.; Zhang, J. A self-powered sensor based on molecularly imprinted polymer-coupled graphic carbon nitride photoanode for selective detection of bisphenol A. *Sensor. Actuat. B-Chem.* **2018**, *259*, 394–401. [CrossRef]

24. Jin, Y.; Luan, Y.; Wu, Z.; Wen, W.; Zhang, X.; Wang, S. Photocatalytic Fuel Cell-Assisted Molecularly Imprinted Self-Powered Sensor: A Flexible and Sensitive Tool for Detecting Aflatoxin B1. *Anal. Chem.* **2021**, *93*, 13204–13211. [CrossRef]

25. Orozo, J.; Cortes, A.; Cheng, G.; Satayasamitsathit, S.; Gao, W.; Feng, X.; Shen, Y.; Wang, J. Molecularly imprinted polymer-based catalytic micromotors for selective protein transport. *J. Am. Chem. Soc.* **2013**, *135*, 5336–5339. [CrossRef]

26. Ouyang, J.; Liu, Z.; Han, Y.; Zeng, K.; Sheng, J.; Deng, L.; Liu, Y.N. Fabrication of Surface Protein-Imprinted Biofuel Cell for Sensitive Self-Powered Glycoprotein Detection. *ACS Appl. Mater. Interfaces* **2016**, *8*, 35004–35011. [CrossRef]

27. Li, G.; Wu, J.; Qi, X.; Wan, X.; Liu, Y.; Chen, Y.; Xu, L. Molecularly imprinted polyppyrole film-coated poly(3,4-ethylenedioxythiophene): Polystyrene sulfonate-functionalized black phosphorene for the selective and robust detection of norfloxacin. *Mater. Today Chem.* **2022**, *26*, 101043. [CrossRef]

28. Li, G.; Qi, X.; Wu, J.; Xu, L.; Wan, X.; Liu, Y.; Chen, Y.; Li, Q. Ultrasensitive, label-free voltammetric determination of norfloxacin based on molecularly imprinted polymers and Au nanoparticle-functionalized black phosphorus nanosheet nanocomposite. *J. Hazard. Mater.* **2022**, *436*, 129107. [CrossRef]

29. Wadie, M.; Marzouk, H.; Rezk, M.; Abdel-Moety, E.; Tantawy, M. A sensing platform of molecularly imprinted polymer-based polyaniline/carbon paste electrodes for simultaneous potentiometric determination of alfuzosin and solifenacin in binary co-formulation and spiked plasma. *Anal. Chim. Acta* **2022**, *1200*, 339599. [CrossRef]

30. Parolo, C.; Merkoci, A. Paper-based nanobiosensors for diagnostics. *Chem. Soc. Rev.* **2013**, *42*, 450–457. [CrossRef]

31. Wang, Y.; Ge, L.; Wang, P.; Yan, M.; Yu, J.; Ge, S. A three-dimensional origami-based immuno-biofuel cell for self-powered, low-cost, and sensitive point-of-care testing. *Chem. Commun.* **2014**, *50*, 1947–1949. [CrossRef] [PubMed]

32. Gu, C.; Gai, P.; Li, F. Construction of biofuel cells-based self-powered biosensors via design of nanocatalytic system. *Nano Energy* **2022**, *93*, 106806. [CrossRef]

33. Guntert, A.T.; Abegg, S.; Konigstein, K.; Gerber, P.A.; Schmidt-Trucksass, A.; Pratsinis, S.E. Breath Sensors for Health Monitoring. *ACS Sens.* **2019**, *4*, 268–280. [CrossRef] [PubMed]

34. Wen, D.; Xu, X.; Dong, S. A single-walled carbon nanohorn-based miniature glucose/air biofuel cell for harvesting energy from soft drinks. *Energy Environ. Sci.* **2011**, *4*, 1358–1363. [CrossRef]

35. Zhang, L.; Zhou, M.; Wen, D.; Bai, L.; Lou, B.; Dong, S. Small-size biofuel cell on paper. *Biosens. Bioelectron.* **2012**, *35*, 155–159. [CrossRef]

36. Wang, Y.; Zhang, L.; Cui, K.; Ge, S.; Zhao, P.; Yu, J. Paper-Supported Self-Powered System Based on a Glucose/O2 Biofuel Cell for Visual MicroRNA-21 Sensing. *ACS Appl. Mater. Interfaces* **2019**, *11*, 5114–5122. [CrossRef]

37. Wang, L.; Shao, H.; Lu, X.; Wang, W.; Zhang, J.R.; Song, R.B.; Zhu, J.J. A glucose/O2 fuel cell-based self-powered biosensor for probing a drug delivery model with self-diagnosis and self-evaluation. *Chem. Sci.* **2018**, *9*, 8482–8491. [CrossRef]

38. Wang, Y.; Zhang, L.; Cui, K.; XU, C.; Li, H.; Liu, H.; Yu, J. Solar driven electrochromic photoelectrochemical fuel cells for simultaneous energy conversion, storage and self-sensing. *Nano尺度* **2018**, *10*, 3421–3428. [CrossRef]

39. Bai, C.; Wang, Z.; Yang, S.; Cui, X.; Li, X.; Yin, Y.; Zhang, M.; Wang, T.; Sang, S.; Zhang, W.; et al. Wearable Electronics Based on the Gel Thermogalvanic Electrolyte for Self-Powered Human Health Monitoring. *ACS Appl. Mater. Interfaces* **2021**, *13*, 37316–37322. [CrossRef]

40. Wang, F.; Xu, J.; Hou, Y.; Huang, K.; Yu, X.; Zhou, X.; Tan, X. Matching Capacitors to Self-Powered Biosensors for Signal Amplification: Toward Ultrasensitive Electrochemical Detection for MicroRNA-21-Triggered Catalytic Hairpin Assembly. *ACS Sustain. Chem. Eng.* **2022**, *10*, 2673–2680. [CrossRef]

41. Wang, Y.; Ge, L.; Ma, C.; Kong, Q.; Yan, M.; Ge, S.; Yu, J. Self-powered and sensitive DNA detection in a three-dimensional origami-based biofuel cell based on a porous Pt-paper cathode. *Chem. Eur. J.* **2014**, *20*, 12453–12462. [CrossRef] [PubMed]

42. Yao, B.; Yuan, L.; Xiao, X.; Zhang, J.; Qi, Y.; Zhou, J.; Zhou, J.; Hu, B.; Chen, W. Paper-based solid-state supercapacitors with pencil-drawing graphite/polyaniline networks hybrid electrodes. *Nano Energy* **2013**, *2*, 1071–1078. [CrossRef]
43. Ge, L.; Wang, P.; Ge, S.; Li, N.; Yu, J.; Yan, M.; Huang, J. Photoelectrochemical lab-on-paper device based on an integrated paper supercapacitor and internal light source. *Anal. Chem.* **2013**, *85*, 3961–3970. [CrossRef] [PubMed]

44. Hou, C.; Fan, S.; Lang, Q.; Liu, A. Biofuel Cell Based Self-Powered Sensing Platform for l-Cysteine Detection. *Anal. Chem.* **2015**, *87*, 3382–3387. [CrossRef] [PubMed]

45. Zhao, M.; Gao, Y.; Sun, J.; Gao, F. Mediatorless glucose biosensor and direct electron transfer type glucose/air biofuel cell enabled with carbon nanodots. *Anal. Chem.* **2015**, *87*, 2615–2622. [CrossRef]