Flexible HIV-1 Biosensor Based on the Au/MoS$_2$ Nanoparticles/Au Nanolayer on the PET Substrate

Minkyu Shin $^{1,†}$, Jinho Yoon $^{1,†}$, Chanyong Yi $^1$, Taek Lee $^2$ and Jeong-Woo Choi $^{1,*}$

$^1$ Department of Chemical & Biomolecular Engineering, Sogang University, 35 Baekbeom-Ro, Mapo-Gu, Seoul 04107, Korea
$^2$ Department of Chemical Engineering, Kwangwoon University, Wolgye-dong, Nowon-gu, Seoul 01899, Korea
$^*$ Correspondence: jwchoi@sogang.ac.kr; Tel.: +82-2-705-8480
$†$ These authors contributed equally to this work.

Received: 9 July 2019; Accepted: 24 July 2019; Published: 26 July 2019

Abstract: An electrochemical flexible biosensor composed of gold (Au), molybdenum disulfide nanoparticles (MoS$_2$ NPs), and Au (Au/MoS$_2$/Au nanolayer) on the polyethylene terephthalate (PET) substrate is developed to detect envelope glycoprotein GP120 (gp120), the surface protein of HIV-1. To fabricate the nanolayer on the PET substrate, Au is sputter coated on the flexible PET substrate and MoS$_2$ NPs are spin coated on Au, which is sputter coated once again with Au. The gp120 antibody is then immobilized on this flexible electrode through cysteamine (Cys) modified on the surface of the Au/MoS$_2$/Au nanolayer. Fabrication of the biosensor is verified by atomic force microscopy, scanning electron microscopy, and cyclic voltammetry. A flexibility test is done using a micro-fatigue tester. Detection of the gp120 is measured by square wave voltammetry. The results indicate that the prepared biosensor detects 0.1 pg/mL of gp120, which is comparable with previously reported gp120 biosensors prepared even without flexibility. Therefore, the proposed biosensor supports the development of a nanomaterial-based flexible sensing platform for highly sensitive biosensors with flexibility for wearable device application.

Keywords: nanolayer; flexible biosensor; HIV-1; MoS$_2$ nanoparticles; PET substrate; gp120

1. Introduction

Flexible biosensors composed of polymer materials have recently attracted significant attention for their application in wearable devices and point-of-care (POC) diagnostic systems. Polymer substrates such as polyethylene terephthalate (PET), polyimide (PI), polycarbonate (PC), and polydimethylsiloxane (PDMS) are widely used as flexible substrates [1,2]. In addition, in order to fabricate flexible biosensors, various types of nanomaterials such as gold nanoparticles (GNPs), carbon nanotubes (CNTs), and graphene oxide (GO) have been introduced for granting the conductivity, enhancing the electron transfer, and biocompatibility [3–8]. Among these nanomaterials, carbon-based nanomaterials have been widely used due to their exceptional properties such as excellent electrical conductivity, high specific activated surface area, and chemical/biological stability [9,10].

Recently, transition metal dichalcogenide (TMD) materials such as tungsten diselenide (WSe$_2$) and molybdenum disulfide (MoS$_2$) have been widely researched for their application in biosensors due to unique properties including electric charge effect and semiconducting property [11–13]. In particular, MoS$_2$ has many advantages for biosensor development due to its electrochemical property and biocompatibility [14,15]. In addition, in order to maximize the benefits of MoS$_2$ on a large activated surface area, MoS$_2$ nanoparticles (MoS$_2$ NPs) have recently been synthesized and applied to develop various sensors [16]. However, most studies related to flexible biosensors composed of nanomaterials have been done through complex manufacturing methods. To replace the complex manufacturing
methods with a simple method for flexible biosensor fabrication, our group reported the flexible biosensor fabricated by a simple manufacturing method using the sputter coating and spin coating on a PI polymer substrate for uniform nanolayer formation. To fabricate an excellent flexible biosensor, PET can be an excellent candidate as the flexible substrate due to its properties including low cost, excellent thermal conductivity, chemical resistance, and dimensionally stability compared with previously used PI substrate [17,18].

Numerous diseases including influenza, severe acute respiratory syndrome (SARS), and acquired immunodeficiency syndrome (AIDS) are caused by small infectious agents called viruses [19]. Among the various lethal viruses, human immunodeficiency virus (HIV) has received massive attention since it causes AIDS which aggravates the human immune system, eventually resulting in death. Since HIV has a long incubation period and can be transmitted during this incubation period, its early and accurate diagnosis is a highly critical issue in the biomedical field [20,21]. The surface of HIV is composed of an envelope glycoprotein made of Envelope glycoprotein GP120 (gp120) and glycoprotein GP41 (gp41) which are connected non-covalently. The gp120 binds with the cluster of differentiation 4 (CD4) receptors and thus, plays an important role in HIV infection [22,23]. Until now, to accurately detect HIV, various methods such as enzyme-linked immunosorbent assay (ELISA) and polymerase chain reaction (PCR) have been used [24,25]. However, these methods have limitations such as complex sample treatment process, time consumption, and low sensitivity [26]. To overcome these limitations, electrochemical immunosensors have been developed for high sensitivity and real-time detection of HIV.

From these points of view, MoS$_2$ nanoparticles (NPs), PET and gp120 can be the core components to develop the HIV biosensor with flexibility and high sensitivity. In this study, an electrochemical flexible biosensor based on gold (Au), MoS$_2$ NPs, and Au (Au/MoS$_2$/Au nanolayer) on a PET substrate was fabricated for the first time to detect the HIV surface protein gp120 using poly(vinylpyrrolidone) (PVP)-modified MoS$_2$ NPs on the PET substrate. PVP-modified MoS$_2$ NPs were synthesized to prevent aggregation of MoS$_2$ NPs, and to facilitate their effective dispersion on the PET substrate. Synthesis of PVP-modified MoS$_2$ NPs was verified by transmission electron microscopy (TEM), energy-dispersive X-ray spectroscopy (EDS), and X-ray diffraction (XRD), while the fabrication of the Au/MoS$_2$/Au nanolayer on the PET substrate was confirmed by field emission scanning electron microscopy (FE-SEM), EDS, and atomic force microscopy (AFM). A gp120 antibody (Ab) was then immobilized on the substrate through the 1-Ethyl-3-(3-dimethylaminopropyl) carbodiimide (EDC)/N-hydroxysuccinimide (NHS) reaction in order to prepare the biosensor (Ab/Cys/Au/MoS$_2$/Au nanolayer). Immobilization of the gp120 antibody was confirmed by cyclic voltammetry (CV). In addition, the electrochemical property for gp120 detection was confirmed by CV and square wave voltammetry (SWV). A micro-fatigue tester (E3000LT, Instron, Buckinghamshire, UK.) was used to evaluate the flexibility of the fabricated Au/MoS$_2$/Au nanolayer on the PET substrate.

2. Materials and Methods

2.1. Materials

Ammonium molybdate tetrahydrate (99.98%, Sigma-Aldrich, Burlington, MA, USA), thiourea (AMRESCO, Solon, OH, USA), and polyvinylpyrrolidone (Sigma-Aldrich, Burlington, MA, USA) were used to synthesize the MoS$_2$ NPs. Potassium hexacyanoferrate (III) ($K_3[Fe(CN)_6]$) (approx. 99.0%, Sigma-Aldrich, Burlington, MA, USA) and potassium hexacyanoferrate (II) trihydrate ($K_4[Fe(CN)_6]$) (≥99.0%, Sigma-Aldrich, Burlington, MA, USA) solution in phosphate-buffered saline (PBS) (Sigma-Aldrich, Burlington, MA, USA) was used as the electrolyte in this study. EDC and NHS were purchased from Thermo Scientific (≥99.0%, Waltham, MA, USA). Human serum (Sigma-Aldrich, Burlington, MA, USA), cysteamine (Cys) (≥98.0%, Sigma-Aldrich, Burlington, MA, USA), gp120 antibody (Sino biological, Wayne, PA, USA), and gp120 antigen (ACRO Biosystem, Newark, DE, USA) were used to fabricate the biosensor. All aqueous solutions were prepared using deionized (DI)
water from a Millipore Milli-Q water purifier operating at a resistance of 18 MΩ·cm. Myoglobin (Mb) (≥90.0%, Sigma-Aldrich, Burlington, MA, USA), hemoglobin (Hb) (Sigma-Aldrich, Burlington, MA, USA), thioredoxin (Trx) (Sino biological, Wayne, PA, USA), and prostate-specific antigen (PSA) (Abcam, Cambridge, UK) were used to investigate the selectivity of the fabricated biosensor.

2.2. MoS$_2$ NPs Synthesis

Ammonium molybdate tetrahydrate (0.35 g) and 0.76 g of thiourea were mixed with 50 mL of DI water. Then 0.25 g of PVP was added to the mixture. The resulting solution was reacted at 800 rpm for 1 h at 60 °C. The solution was transferred into a Teflon-lined stainless-steel autoclave. The autoclave was set at 200 °C for 24 h. After the reaction was complete, the resulting black precipitate was cooled down to room temperature and washed with DI water and ethanol using centrifugal filtration with 8000 rpm for 30 min. The final product was air-dried at 60 °C for 24 h. The PVP-modified MoS$_2$ NPs (concentration: 1 mg/mL) were characterized by high-resolution TEM using a JEOL JEM-3010 operated at 300 kV and by XRD (Rigaku, Tokyo, Japan).

2.3. Fabrication of the Au/MoS$_2$/Au Nanolayer on the PET Substrate, and Immobilization of the gp120 Antibody

The fabrication of a flexible biosensor composed of Ab/Cys/Au/MoS$_2$/Au nanolayer on PET substrate is shown in Figure 1. To fabricate the Au/MoS$_2$/Au nanolayer on the PET substrate, the rectangular PET substrate (size 1 mm × 2 mm) was cleaned in a sonication bath for 30 min using acetone and DI water, and then completely dried with N$_2$ gas. After cleaning the PET, the Au/MoS$_2$/Au nanolayer on the PET substrate was prepared by gold sputter and spin coater. The Au layer was formed by sputter coating on the PET substrate. Then, 200 µL of the synthesized MoS$_2$ NPs dissolved in DI water (concentration: 5 mg/mL) was dropped onto the Au layer and spin coated (Au/MoS$_2$) on the PET substrate at 2000 rpm for 30 s; this process was done twice. After the spin coating process, the Au layer was sputter coated once more on the MoS$_2$ layer. Fabrication of the Au/MoS$_2$/Au nanolayer on the PET substrate was verified by FE-SEM, EDS, and AFM. To fabricate the Ab/Cys/Au/MoS$_2$/Au nanolayer on the PET substrate, the gp120 antibody was immobilized on the Au/MoS$_2$/Au nanolayer on the PET substrate. The fabricated electrode was washed with ethanol and DI water in preparation for the gp120 antibody immobilization. Cys (77 mg) dissolved in 10 mL of DI water (10 mM concentration) was then immobilized on the electrode by self-assembly for 3 h at room temperature. The Cys-immobilized electrode was washed with DI water to remove any unbound Cys. Then, the gp120 antibody was attached to Cys using the EDC/NHS reaction. To achieve this, 100 µL of the antibody solution (concentration: 5 µg/mL) was mixed with 100 µL of EDC (concentration: 4 mg/mL) and 100 µL of NHS (concentration: 6 mg/mL) for 1 h at room temperature. Then, the modified gp120 antibody with EDC/NHS was dropped on the Cys-immobilized electrode for 2 h at room temperature. In addition, the gp120 antibody-immobilized electrode was washed with DI water to remove any unbound gp120 antibody and dried with N$_2$ gas.
19.63% of MoS$_2$ were as follows: amplitude of 20 mV, frequency of 10 Hz, and 1
The measured DLS data indicated that 80.37% of synthesized MoS$_2$ and Figure S1. EDS and EDS mapping images showed that the ratio of molybdenum (Mo) and sulfur (S) was approximately 1:2, which matched the theoretical composition of MoS$_2$. Thus, these experiments verified the synthesis of MoS$_2$.

3. Results and Discussion

3.1. Confirmation of MoS$_2$ NPs Synthesis

The conformation and composition of the synthesized MoS$_2$ NPs were investigated by TEM and EDS analysis. As shown in Figure 2a, the TEM images indicated that the synthesized MoS$_2$ NPs were structured as well-dispersed nanospheres. The EDS and EDS mapping results are shown in Figure 2b and Figure S1. The XRD pattern of the synthesized MoS$_2$ NPs showed peaks at 13°, 17°, 24°, 33°, 43°, and 58° for the hexagonal phase of MoS$_2$ NPs shown in Figure 2d. The electrochemical properties of the fabricated biosensor were analyzed by electrochemical analyzer CHI-660E (CH Instruments, Inc., Austin, TX, USA). CV and SWV were conducted with a three-electrode system composed of a fabricated working electrode, a platinum (Pt) wire counter electrode, and a silver/silver chloride (Ag/AgCl) reference electrode. The parameters for the CV experiment were as follows: amplitude of 20 mV, frequency of 10 Hz, and 1 x 10$^{-3}$ (A/V) sensitivity. The applied voltage range was from 500 mV to −200 mV. The parameters for the SWV experiment were as follows: amplitude of 20 mV, frequency of 10 Hz, and 1 x 10$^{-3}$ (A/V) sensitivity. The applied voltage range was from 500 mV to −200 mV. The electrochemical experiments were conducted with PBS solution (pH = 7.4) containing 5 mM K$_3$Fe(CN)$_6$ and 5 mM K$_4$Fe(CN)$_6$ which were used as the redox generator for electrochemical investigation.

2.5. Flexibility Test of Fabricated Biosensor

The flexibility of the fabricated biosensor was investigated using a micro-fatigue tester. To estimate the flexibility, the electrode was fixed to the tester and the tip of the tester was moved down to apply force on the electrode. The electrode was bent by the force applied to the center of the electrode. The elongation of the electrode by the applied force was measured to investigate the flexibility of the electrode. The fabricated biosensor was compared with a widely used rigid gold coated silicon electrode.
Electron transfer by MoS\(_2\) compared to conventional bare gold electrode was due to the large surface area and the efficient electron transfer by MoS\(_2\) NPs.

3.2. Verification of the Au/MoS\(_2\)/Au Nanolayer on the PET Substrate

Fabrication of the Au/MoS\(_2\)/Au nanolayer on the PET substrate was confirmed by AFM and FE-SEM. Figure 3a–c shows AFM images of the PET substrate, Au sputter coated PET substrate, and Au/MoS\(_2\) on the PET substrate, respectively. In Figure 3a, the AFM results of the PET substrate indicated a height of 1.296 nm. When Au was sputter coated on the PET substrate, the height of the substrate increased to 7.856 nm due to the deposited Au as shown in Figure 3b. The MoS\(_2\) NPs immobilized on the Au coated PET substrate by spin coating was shown in Figure 3c, the height of the immobilized MoS\(_2\) NPs was about 155.81 nm. In addition, Figure 3d–f shows FE-SEM images of the fabricated Au/MoS\(_2\)/Au nanolayer on the PET substrate. Compared to the results in Figure 3a, the existence of the sputter coated Au particles on MoS\(_2\) NPs was confirmed in Figure 3f. EDS analysis of all the acquired SEM images is shown in Figure S2. The amount of sputter coated gold in the Au/MoS\(_2\)/Au nanolayer on the PET substrate was doubled to the Au/MoS\(_2\) on the PET substrate due to the twice gold deposition. In addition, the EDS mapping result of the Au/MoS\(_2\)/Au nanolayer on the PET substrate is shown in Figure S3. Also, the CV result is shown in Figure 4a for confirmation of the fabrication of the Au/MoS\(_2\)/Au nanolayer on the PET substrate. In Figure 4a, the CV results of the PET substrate showed no electrochemical signals, but signals were observed after Au sputter coating on the PET substrate. Also, the fabricated Au/MoS\(_2\)/Au nanolayer on the PET substrate had higher electrochemical signals than the conventional bare gold electrode composed of gold (50 nm)/Cr (2 nm)/SiO\(_2\) (Silicon dioxide). The current increase of the Au/MoS\(_2\)/Au nanolayer on the PET substrate compared to conventional bare gold electrode was due to the large surface area and the efficient electron transfer by MoS\(_2\) NPs.
10 mV, 1.38 mA, and 1.30 mA. In Figure 4c, the CV results were obtained by increasing the scan rate from with a \([\text{Fe(CN)}_6]^{3-/4-}\) reduction and oxidation peak currents of the measured CV showed a linear response to the increase in the scan rate. Figure 4d showed that the plotted reduction and oxidation peak currents of the measured CV showed a linear response to the increase in the scan rate.

### 3.3. Investigation of the Electrochemical Properties of the Fabricated Biosensor

CV and SWV were performed to confirm the electrochemical properties of the fabricated biosensor with a \([\text{Fe(CN)}_6]^{3-/4-}\) redox probe. Figure 4b shows the CV result of the Au/MoS\(_2\)/Au nanolayer on the PET substrate, the Cys/Au/MoS\(_2\)/Au nanolayer on the PET substrate, the Ab/Cys/Au/MoS\(_2\)/Au nanolayer on the PET substrate. The reduction and oxidation peak currents of the Au/MoS\(_2\)/Au nanolayer on the PET substrate was found to be 1.13 mA and -1.17 mA, respectively. After Cys immobilization, the reduction and oxidation peak potentials were changed from 0.05 V and 0.35 V to 0.12 V and 0.28 V, and the reduction and oxidation peak potentials were also increased to 1.36 mA and -1.37 mA, respectively. This increase was due to the electrostatic interaction between the negatively charged \([\text{Fe(CN)}_6]^{3-/4-}\) and the positively charged amine group of the Cys immobilized on the Au/MoS\(_2\)/Au nanolayer on the PET substrate. However, when the gp120 antibody was immobilized on the Cys, the efficiency of electron transfer between the multilayer and the redox probe was reduced by the gp120 antibody, and the reduction and oxidation peak currents were lowered to 1.30 mA and -1.34 mA, respectively. In addition, to confirm the reproducibility of fabricated biosensor, the average reduction peak current was investigated with four different measurements. As shown in Figure S4, the average reduction peaks of Au/MoS\(_2\)/Au nanolayer on the PET substrate, Cys/Au/MoS\(_2\)/Au nanolayer on the PET substrate and Ab/Cys/Au/MoS\(_2\)/Au nanolayer on the PET substrate were 1.12 mA, 1.38 mA, and 1.30 mA. In Figure 4c, the CV results were obtained by increasing the scan rate from 10 mV/s to 200 mV/s to verify the relationship between the current peak and the scan rate of the fabricated the Au/MoS\(_2\)/Au nanolayer on the PET substrate. Figure 4d showed that the plotted reduction and oxidation peak currents of the measured CV showed a linear response to the increase in the scan rate.

**Figure 3.** Atomic force microscopy (AFM) images of (a) the PET substrate, (b) Au sputter coated PET substrate, and (c) the Au/MoS\(_2\) on the PET substrate, and SEM images of (d) Au sputter coated PET substrate, (e) the Au/MoS\(_2\) on the PET substrate, and (f) the Au/MoS\(_2\)/Au nanolayer on the PET substrate.
3.4. Detection of the gp120 Antigen and Selectivity

The SWV technique was performed to investigate the electrochemical detection performance and selectivity of the fabricated biosensor. To confirm its detection, the gp120 antigen dissolved in PBS solution was immobilized on the prepared Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate for 1 h at room temperature. The detection of the gp120 antigen at a concentration of 0.1 pg/mL to 10 ng/mL was confirmed by SWV. As shown in Figure 5a, the electron transfer reaction between the redox probe and the biosensor surface was blocked by the antigen–antibody binding on the surface of the fabricated biosensor, and the current value was decreased when the concentration of the gp120 antibody was increased. In addition, the linearity of the current value for the gp120 antigen concentration was confirmed through the results obtained from SWV. As shown in Figure 5b, the gp120 antigen ranged from 0.1 pg/mL to 10 ng/mL and showed excellent linearity with 0.973 of the coefficient of determination value ($R^2$). In general, the concentration of gp120 in HIV-infected patients is approximately 200 pg/mL to 2000 pg/mL, therefore the fabricated biosensor showed the possibility for gp120 detection in practice. The mean value and the error bars were plotted with the standard deviation (SD) from four different measurements. The detection limit of an electrochemical biosensor for gp120 detection in practice is approximately 200 pg/mL to 2000 pg/mL, therefore the fabricated biosensor showed the possibility for gp120 detection in practice. The mean value and the error bars were plotted with the standard deviation (SD) from four different measurements. To confirm its detection, the gp120 antigen dissolved in PBS solution was immobilized on the prepared Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate, and the Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate against scan rates. (c) Cyclic voltammograms of the Au/MoS2/Au nanolayer on the PET substrate at scan rates increasing from 10 to 200 mV/s. (d) Linear-response plot of redox current peaks of the Au/MoS2/Au nanolayer on the PET substrate against scan rates.
concentrations from 1 pg/mL to 10 ng/mL was mixed with serum. As shown in Figure 5c, in the case of the gp120 antigen prepared in serum, the current value decreased with increase in concentration, the same as the gp120 antigen dissolved in PBS. The selectivity of the Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate-based biosensor was measured by the addition of various types of antigens and proteins including Hb, Mb, PSA, and Trx prepared in PBS solution. The concentrations of the gp120 antigen, Hb, Mb, PSA, and Trx were fixed at 100 ng/mL. As shown in Figure 5d, the gp120 antigen had a low current value of 0.925 mA, while Hb, Mb, PSA, and Trx had high current values of 1.26 mA, 1.24 mA, 1.30 mA, and 1.26 mA, respectively. In addition, the rate of change of current value of gp120 antigen, Hb, Mb, PSA, and Trx were indicated 29.9%, 4.42%, 6.43%, 1.45%, and 4.60%. The mean value and the error bars were obtained as the standard deviation (SD) of five measurements. As shown in the results of the selectivity test, the measurement of gp120 was confirmed since the current value of the gp120 antigen, a surface protein of HIV was smaller than that of other antigens and proteins, and the electrochemical signal decreased due to selective immobilization of the gp120.

Figure 5. (a) Square wave voltammetry (SWV) results and (b) linear-response plot of the current peak values for the concentration of gp120 from 0.1 pg/mL to 10 ng/mL in phosphate-buffered saline (PBS) solution. (c) SWV results for the concentration of gp120 from 1 pg/mL to 10 ng/mL in serum. (d) Selectivity test of the Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate to various types of antigens and proteins including Hb, Mb, PSA, and Trx prepared in PBS solution.
Table 1. Electrochemical biosensors for human immunodeficiency virus (HIV) detection.

| Electrode                                      | Technique | Target | Detection Limit | Linear Range       | Reference |
|------------------------------------------------|-----------|--------|-----------------|--------------------|-----------|
| GCE/CNF-Bi/MIP/NBD-556@ gp120                 | DPV       | gp120  | 0.3 pg/mL       | 0.002–200 ng/mL    | [27]      |
| FP-50 fusion peptide                          | Dot blot  | gp120  | -               | 0–100 pg/mL        | [28]      |
| Poly(propylene imine) Dendrimer-Streptavidin   | SWV       | gp120  | 4.12 pg/ml      | 12.04 ng/mL–0.19 μg/mL | [29] |
| Au nonodot/ITO                                 | CV        | HIV-1  | -               | 0.6–375 pg/mL      | [30]      |
| Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate | SWV     | gp120  | 0.066 pg/mL     | 0.1 pg/mL–10 ng/mL| This research |

3.5. Investigation of the Flexibility of the Au/MoS2/Au Nanolayer on the PET Substrate

The flexibility of the fabricated biosensor was confirmed by a micro-fatigue tester. Figure 6a,b show flexibility results of a conventional bare gold electrode, Au sputter coated PET substrate, and the Au/MoS2/Au nanolayer on the PET substrate. In Figure 6a, the flexure extension results of the conventional bare gold electrode were 0.076 mm due to the hardness of the electrode. SiO2-based rigid substrates were hard to bend using applied forces due to their lack of flexibility. However, Au sputter coated PET and the Au/MoS2/Au nanolayer on the PET substrate showed excellent flexibility and flexure extension with 1.60 mm and 1.54 mm, respectively. These results were significantly higher than those of the conventional electrode because these electrodes were easily bent by the applied force due to their excellent flexibility. In addition, the Au sputter coated PET and Au/MoS2/Au nanolayer on the PET substrate had flexure strength of 92.9 MPa and 99.2 MPa, respectively, which were lower than that of the conventional gold electrode due to the excellent flexibility. As shown in Figure 6b, the conventional gold electrode was found to be rapidly damaged by strong forces due to the hardness of the electrode. However, the fabricated biosensor had a high flexure extension compared with the conventional gold electrode because of the small force applied to the substrate due to the characteristics of the flexible substrate. In addition, SWV was performed to investigate electrochemical detection performance of the bent biosensor. To confirm the detection performance, 10 ng/mL of gp120 antigen was immobilized on the Ab/Cys/Au/MoS2/Au nanolayer on the PET substrate. As shown in Figure S5, the current value of fabricated biosensor before bent was indicated 1.06 mA, and after the fabricated biosensor was bent, the current value was maintained to 1.03 mA.

![Figure 6](image-url)
4. Conclusions

In this study, the flexible biosensor based on a Au/MoS$_2$/Au nanolayer on a PET substrate was developed to detect gp120 with high sensitivity. To develop the flexible biosensor, the Au/MoS$_2$/Au nanolayer was fabricated by Au sputter coating and MoS$_2$ NPs spin coating onto a flexible PET substrate. The fabricated Au/MoS$_2$/Au nanolayer on the PET substrate showed the well-oriented NPs and uniform nanolayer formation on the PET substrate. The reduction and oxidation peak currents of the Au/MoS$_2$/Au nanolayer on the PET substrate derived from the redox generator were 1.13 mA and −1.17 mA, respectively, which were much higher than those peaks of the bare gold electrodes with 0.96 mA and −1.01 mA due to the large surface area and effective electron transfer of the synthesized MoS$_2$ NPs. The fabricated biosensor showed highly sensitive detection of gp120 with a detection limit of 0.066 pg/mL, which was more sensitive than previously reported electrochemical HIV biosensors. This biosensor showed a selective detection of gp120 added with various antigens and proteins such as Hb, Mb, PSA, and Trx. In addition, this biosensor showed excellent flexibility with flexure extension of 1.54 mm compared to SiO$_2$-based conventional gold electrodes, and the fabricated biosensor maintained the detection performance after bending. In conclusion, the proposed flexible biosensor based on a Au/MoS$_2$/Au nanolayer on a PET substrate can suggest the milestone for nanomaterial-based flexible sensing platform to develop the highly sensitive biosensors with flexibility for a wearable device application. In addition, because the gp120 concentration range of the HIV infected patient can be measured, it can be used in the commercial field.

Supplementary Materials: The following are available online at http://www.mdpi.com/2079-4991/9/8/1076/s1, Figure S1: TEM and EDS analysis of MoS$_2$ NPs. Figure S2: SEM and EDS analysis of (a) Au sputter coated PET substrate, (b) the Au/MoS$_2$ on the PET substrate and (c) the Au/MoS$_2$/Au nanolayer on the PET substrate. Figure S3: EDS mapping results of the Au/MoS$_2$/Au nanolayer on the PET substrate. Figure S4: The reproducibility of reduction peaks of the Au/MoS$_2$/Au nanolayer on the PET substrate, Cys/Au/MoS$_2$/Au nanolayer on the PET substrate and Ab/Cys/Au/MoS$_2$/Au nanolayer on the PET substrate. Error bars indicate the standard deviations of four different measurements. Figure S5: Reduction peak currents of the 10 ng/mL of gp120 antigen on the Ab/Cys/Au/MoS$_2$/Au nanolayer on the PET substrate before bending and after bending. Error bars indicate the standard deviations of four different measurements.

Author Contributions: Conceptualization, M.S., J.Y., C.Y., T.L., and J.-W.C.; investigation, M.S., J.Y., and C.Y.; supervision, J.-W.C.; project administration, J.-W.C.; writing—original draft preparation, M.S. and J.Y.; writing—review and editing, J.Y., T.L., and J.-W.C.

Funding: This research was supported by the National Research Foundation of Korea (NRF) grant funded by the Korea government (MSIT) (No. 2019R1A2C3002300) and Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education (No. 2016R1A6A1A03012845).

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

References

1. Yang, Y.; Luo, C.; Jia, J.; Sun, Y.; Fu, Q.; Pan, C. A Wrinkled Ag/CNTs-PDMS Composite Film for a High-Performance Flexible Sensor and Its Applications in Human-Body Single Monitoring. Nanomaterials 2019, 9, 850. [CrossRef] [PubMed]
2. Wang, X.; Tao, L.; Hao, Y.; Liu, Z.; Chou, H.; Khomlanov, I.; Chen, S.; Tan, C.; Jayant, N.; Yu, Q.; et al. Direct delamination of graphene for high-performance plastic electronics. Small 2014, 10, 694–698. [CrossRef] [PubMed]
3. Jo, J.; Yoon, J.; Lee, T.; Cho, H.-Y.; Lee, J.-Y.; Choi, J.-W. H$_2$O$_2$ biosensor consisted of hemoglobin-DNA conjugate on nanoporous gold thin film electrode with electrochemical signal enhancement. Nano Converg. 2019, 6, 1. [CrossRef] [PubMed]
4. Venditti, I. Engineered Gold-Based Nanomaterials: Morphologies and Functionalities in Biomedical Applications. A Mini Review. Bioengineering 2019, 6, 53. [CrossRef] [PubMed]
5. Kurdekar, A.D.; Avinash Chunduri, L.A.; Manohar, C.S.; Haleyurgrisetty, M.K.; Hewlett, I.K.; Venkataramaniah, K. Streptavidin-conjugated gold nanoclusters as ultrasensitive fluorescent sensors for early diagnosis of HIV infection. Sci. Adv. 2018, 4, eaar6280. [CrossRef]
6. Hsu, M.-S.; Chen, Y.-L.; Lee, C.-Y.; Chiu, H.-T. Gold Nanostructures on Flexible Substrates as Electrochemical Dopamine Sensors. *ACS Appl. Mater. Interfaces* 2012, 4, 5570–5575. [CrossRef]

7. Loghin, F.; Rivadeneyra, A.; Becherer, M.; Lugli, P.; Bobinger, M. A Facile and Efficient Protocol for Preparing Residual-Free Single-Walled Carbon Nanotube Films for Stable Sensing Applications. *Nanomaterials* 2019, 9, 471. [CrossRef]

8. Lin, S.; Feng, W.; Miao, X.; Zhang, X.; Chen, S.; Chen, Y.; Wang, W.; Zhang, Y. A flexible and highly sensitive nonenzymatic glucose sensor based on DVD-laser scribed graphene substrate. *Biosens. Bioelectron.* 2018, 110, 89–96. [CrossRef]

9. El-Said, W.A.; Yoon, J.; Choi, J.-W. Nanostructured surfaces for analysis of anticancer drug and cell diagnosis based on electrochemical and SERS tools. *Nano Converg.* 2018, 5, 11. [CrossRef]

10. Mohapatra, J.; Ananthoju, B.; Nair, V.; Mitra, A.; Bahadur, D.; Medhekar, N.V.; Aslam, M. Enzymatic and non-enzymatic electrochemical glucose sensor based on carbon nano-onions. *Appl. Surf. Sci.* 2018, 442, 332–341. [CrossRef]

11. Wu, S.; Liu, G.; Li, P.; Liu, H.; Xu, H. A high-sensitive and fast-fabricated glucose biosensor based on Prussian blue/topological insulator Bi₂Se₃ hybrid film. *Biosens. Bioelectron.* 2012, 38, 289–294. [CrossRef] [PubMed]

12. Mazánek, V.; Mayorga-Martinez, C.C.; Bouša, D.; Sofer, Z.; Pumera, M. WSe₂ nanoparticles with enhanced hydrogen evolution reaction prepared by bipolar electrochemistry: Application in competitive magneto-immunoassay. *Nanoscale* 2018, 10, 23149–23156. [CrossRef] [PubMed]

13. Wang, T.; Zhu, H.; Zhuo, J.; Zhu, Z.; Papakonstantinou, P.; Lubarsky, G.; Lin, J.; Li, M. Biosensor Based on Ultrasmall MoS₂ Nanoparticles for Electrochemical Detection of H₂O₂ Released by Cells at the Nanomolar Level. *Anal. Chem.* 2013, 85, 10289–10295. [CrossRef] [PubMed]

14. Yoon, J.; Lee, S.N.; Shin, M.K.; Kim, H.-W.; Choi, H.K.; Lee, T.; Choi, J.-W. Flexible electrochemical glucose biosensor based on GOx/gold/MoS₂/gold nanofilm on the polymer electrode. *Biosens. Bioelectron.* 2019, 140, 111343. [CrossRef] [PubMed]

15. Su, S.; Zou, M.; Zhao, H.; Yuan, C.; Xu, Y.; Zhang, C.; Wang, L.; Fan, C.; Wang, L. Shape-controlled gold nanoparticles supported on MoS₂ nanosheets: Synergistic effect of thionine and MoS₂ and their application for electrochemical label-free immunosensing. *Nanoscale* 2015, 7, 19129–19135. [CrossRef]

16. Kim, S.J.; Mondal, S.; Min, B.K.; Choi, C.-G. Highly Sensitive and Flexible Strain–Pressure Sensors with Cracked Paddy-Shaped MoS₂/Graphene Foam/Ecoflex Hybrid Nanostructures. *ACS Appl. Mater. Interfaces* 2018, 10, 36377–36384. [CrossRef] [PubMed]

17. Yaqoob, U.; Phan, D.-T.; Uddin, A.S.M.I.; Chung, G.-S. Highly flexible room temperature NO₂ sensor based on MWCNTs-WO₃ nanoparticles hybrid on a PET substrate. *Sens. Actuators B Chem.* 2015, 221, 760–768. [CrossRef]

18. König, T.; Münstedt, H. Coatings of indium tin oxide nanoparticles on various flexible polymer substrates: Influence of surface topography and oscillatory bending on electrical properties. *J. Soc. Inf. Disp.* 2008, 16, 559. [CrossRef]

19. Saxena, S.K.; Tiwari, S.; Nair, M.P.N. A Global Perspective on HIV/AIDS. *Science* 2012, 337, 798. [CrossRef]

20. Embretson, J.; Zupanic, M.; Ribas, J.L.; Burke, A.; Racz, P.; Tenner-Racz, K.; Haase, A.T. Massive covert infection of helper T lymphocytes and macrophages by HIV during the incubation period of AIDS. *Nature* 1993, 362, 359–362. [CrossRef]

21. Pilcher, C.D.; Tien, H.C.; Eron, J.J.; Vernazza, P.L.; Leu, S.-Y.; Stewart, P.W.; Goh, L.-E.; Cohen, M.S. Brief but Efficient: Acute HIV Infection and the Sexual Transmission of HIV. *J. Infect. Dis.* 2004, 189, 1785–1792. [CrossRef] [PubMed]

22. Kim, Y.-G.; Moon, S.; Kuritzkes, D.R.; Demirci, U. Quantum dot-based HIV capture and imaging in a microfluidic channel. *Biosens. Bioelectron.* 2009, 25, 253–258. [CrossRef] [PubMed]

23. Zhu, P.; Liu, J.; Bess, J.; Chertova, E.; Lifson, J.D.; Grisé, H.; Ofek, G.A.; Taylor, K.A.; Roux, K.H. Distribution and three-dimensional structure of AIDS virus envelope spikes. *Nature* 2006, 441, 847–852. [CrossRef] [PubMed]

24. De La Rica, R.; Stevens, M.M. Plasmonic ELISA for the ultrasensitive detection of disease biomarkers with the naked eye. *Nat. Nanotechnol.* 2012, 7, 821–824. [CrossRef] [PubMed]

25. McFall, S.M.; Wagner, R.L.; Jangam, S.R.; Yamada, D.H.; Hardie, D.; Kelso, D.M. A simple and rapid DNA extraction method from whole blood for highly sensitive detection and quantitation of HIV-1 proviral DNA by real-time PCR. *J. Virol. Methods* 2015, 214, 37–42. [CrossRef]
26. Chen, C.-C.; Lai, Z.-L.; Wang, G.-J.; Wu, C.-Y. Polymerase chain reaction-free detection of hepatitis B virus DNA using a nanostructured impedance biosensor. *Biosens. Bioelectron.* 2016, 77, 603-608. [CrossRef]

27. Ma, Y.; Liu, C.; Wang, M.; Wang, L.-S. Sensitive electrochemical detection of gp120 based on the combination of NBD-556 and gp120. *Talanta* 2019, 196, 486-492. [CrossRef]

28. Lines, J.A.; Yu, Z.; Dedkova, L.M.; Chen, S. Design and expression of a short peptide as an HIV detection probe. *Biochem. Biophys. Res. Commun.* 2014, 443, 308-312. [CrossRef]

29. John, S.V.; Rotherham, L.S.; Khati, M.; Mamba, B.B.; Arotiba, O.A. Towards HIV Detection: Novel Poly(propylene imine) Dendrimer-Streptavidin Platform for Electrochemical DNA and gp120 Aptamer Biosensors. *Int. J. Electrochem. Sci.* 2014, 9, 13.

30. Lee, J.-H.; Oh, B.-K.; Choi, J.-W. Electrochemical sensor based on direct electron transfer of HIV-1 virus at Au nanoparticle modified ITO electrode. *Biosens. Bioelectron.* 2013, 49, 531-535. [CrossRef]

© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).