Multimode optoelectrochemical detection of cysteine based on an electrochromic Prussian blue electrode

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Abstract

In the present work, multiple detection modes of an electrochromic (EC) sensor were studied and compared. We utilized a Prussian blue (PB) thin film modified F-doped tin oxide (FTO) electrode to detect l-cysteine (Cys), a naturally occurring amino acid bearing a thiol group. The sensing was triggered by a cyclic voltammetric (CV) scan and functioned upon the catalytic oxidation of Cys at the surface of a PB/FTO electrode that had been oxidized to the Berlin green (BG) state. With the EC property of PB and the use of transparent FTO electrode, the Cys concentration, denoted as [Cys], could be detected both electrochemically and optically. As a result, the following four types of calibration curves were obtained simultaneously from the same \textit{in situ} optoelectrochemical measurement: amperometry (current versus [Cys]), coulometry (charge versus [Cys]), potentiometry (potential versus [Cys]), and absorptometry (optical density change versus [Cys]). This proof-of-concept study suggests that the EC detection can provide more informative results than pure electrochemical or optical sensing approach does.

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1. Introduction

Cysteine (C\textsubscript{3}H\textsubscript{7}NO\textsubscript{2}S) is a naturally occurring, sulfur-containing amino acid that plays many important roles in biological systems, such as detoxification, metabolism, a critical substrate for protein synthesis and folding (formation of disulfide bonds), a precursor for antioxidant glutathione synthesis, and a key extracellular reducing agent [1]. For food industry, cysteine has been used as the raw materials for the production of various flavors [2] and has been served as a processing aid for baking [3]. In clinics, it was reported that altered levels of the plasma Cys could link in many pathological conditions, including Alzheimer’s and Parkinson’s diseases [4]. For pharmaceutical application, the well-known N-acetyl cysteine (NAC), a Cys derivative, is often used as a cough medicine and has been investigated for other new treatments recently [5]. In some biotechnology research, cysteine contains a highly nucleophilic thiol group (∼SH, also called sulfhydryl group) and is thus a very popular target for site-directed labeling bioassays [6] that investigate biomolecular structure and dynamics. As a consequence, cysteine detection is of great importance for industrial, health care, and fundamental research applications. To date, many analytical approaches have been developed for Cys detection, and they can be categorized into the following two types, electrical detection and optical detection. Electrical Cys detection can be carried out with amperometry [7], potentiometry [8], cyclic voltammetry (CV) [9], and differential pulse voltammetry (DPV) [9]. Optical Cys detection can be done with spectrofluorimetry [10] and colorimetry [11]. Both electrical and optical detection schemes are well-suited for flow injection analysis (FIA) [12] or high-performance liquid chromatography (HPLC) [13,14] that helps in the precise determination of Cys level in body fluids and other biological samples. In order to develop a Cys sensor that generates both electrical and optical signals simultaneously for more informative detection, an electrochromic Prussian blue (PB) thin film electrode was used as the transducer for Cys sensing in this work.

An electrochromic (EC) material changes color when its redox state is altered electrochemically. Based on this property,
a variety of EC thin films have been synthesized and deposited onto optically transparent electrodes (OTEs), such as F-doped tin oxide (FTO) and indium tin oxide (ITO) glasses, for assembling into a unique class of electro-optical devices, namely electrochromic devices (ECDs) [15]. For instance, Prussian blue (PB, KFeIII[FeII(CN)6]), a classical dark blue pigment, is one of the frequently used EC materials [16]. When deposited on an OTE, Prussian blue is able to exhibit a reversible transparent-blue-green multiple color change corresponding to the following redox states: Prussian white (PW, K2FeIII[FeII(CN)6]), PB, and Berlin green (BG, KFeIII[FeII(CN)6]1/3, FeIII[FeII(CN)6]2/3). Hence, Prussian blue-based ECDs have been investigated for the applications in commercial displays, smart daylighting-control windows, anti-glare rearview mirrors, and sunglasses [17]. Here we demonstrate another promising application of EC thin films – a novel biosensing approach with real-time multiple detection modes – by the PB-modified FTO electrode (denoted as PB/FTO).

In fact, most of the EC materials also can serve as solid-state redox mediators or recognition elements for their superior electrocatalytic activity to certain biomolecules and high ion selectivity. Hence, a variety of electrode materials, such as glassy carbon [18], graphite [19], carbon paste [20], Pt [21], gold [22], ITO [23], and FTO [24], have been modified with PB thin films for electrochemical sensor research and development. Some of the PB electrochemical sensors are based on the function of high selectivity of counter ions [25,26]. Other systems employ direct electrocatalysis strategy, such as detection of hydrogen peroxide [18,21,24], cysteine [13], morphine [23], and so on. The others take effect via coupling with redox enzymes [19,20] like glucose oxidase [27,28], alcohol oxidase [28], and cholesterol oxidase [29], just to name a few. The comprehensive discussions over PB-based biosensors can be found in the recent review articles [30,31]. In addition to electrochemical detection, many efforts had been made to develop optical sensors based on the photosynthesized PB/polymer composite film [32–36] or PB-immobilized beads [37]. In general, such a PB-based optical biosensor worked through the absorptometric detection of analyte-induced PB/PW interchange. For example, the PB film chemically deposited on a non-conductive, transparent substrate could be reduced by a reductant like ascorbic acid [32–34,37] and thus exhibited a blue-to-colorless optical change; on the other hand, the colorless PW film could be oxidized to PB (blue state) by an oxidant like hydrogen peroxide [33,34]. This approach could be used to fabricate an optical glucose [35] and urea [36] biosensor for flow injection analysis, when enzymes were entrapped into the PB films. But, different from a typical amperometric sensor, the optical sensors utilized chemical method to regenerate the PB mediator in stead of applying a dc bias and might therefore elongate the recovery time before the next sensing experiment. In contrast, what we developed in this work is a new biosensing approach, featuring both electrochemical and optical detections by taking advantages of the EC property of PB thin film. To validate the EC sensing method, cysteine (Cys) was chosen as a model analyte.

In the following sections, we will report an in situ optoelectrochemical Cys measurement at a PB/FTO electrode that electrocatalyzes Cys in a buffered solution by a voltammetric scan. Through simultaneous analysis of voltammetric and voltabsorptometric responses [38], our new detection scheme is able to generate amperometric, coulometric, potentiometric, and absorptometric data in a single sensing experiment. It will be shown that EC sensing can provide more informative data than pure amperometric or optical detection does. In addition, the mechanism of EC sensing will be discussed.

2. Experimental

2.1. Materials and instrumentation

The main chemicals used in this work were FeCl3, K3Fe(CN)6, KCl, HCl, H3PO4, KH2PO4, and L-cysteine. All of them were ACS reagent grade and not further purified. Deionized water (DIW) was used throughout. F-doped SnO2 (FTO)-coated glass substrates (Rsh = 20Ω□ and 2 mm in thickness) were obtained from a local supplier (Sinonor Corporation, Hsinchu, Taiwan). Before using, FTO glass substrates were washed ultrasonically with 0.1N HCl for 5 min and with DIW for another 5 min. After an extra DIW rinse, the substrates were dried in air. When preparing a FTO electrode, a piece of cooper tape (3 M Company), serving as the bus bar, was applied to the top edge of the FTO-coated surface, and then an insulating tape was applied to the same surface of glass substrate to define an active electrode area of 3.0 cm × 1.5 cm.

All of the electrochemical experiments were performed in a static, three-electrode fashion. A homemade Ag/AgCl/sat’d KCl reference electrode (−94.7 mV versus commercial SCE) and a Pt coil counter electrode were used. All electrochemical experiments were controlled and monitored using a potentiostat/galvanostat (Autolab, model PGSTAT30), including electrodeposition of PB thin films and in situ optoelectrochemical detection of Cys. A UV–vis spectrophotometer (Shimadzu, model UV-1601PC) was used to collect absorptometric data. All of the experiments were done at room temperature.

2.2. Electrodeposition of PB thin films on FTO glass substrates

Electrodeposition of PB thin films on FTO glass substrates (active area = 3.0 cm × 1.5 cm) was done by applying a cathodic current density of 20 μA/cm2 for 900 s, i.e., total charge capacity input for the deposition was equal to 18.0 mC/cm2. The deposition bath was composed of 10 mM K3Fe(CN)6, 10 mM FeCl3, 0.1 M KCl, and 1.0 M HCl. Presumably, the PB thin film deposited in a KCl-containing solution was the soluble form (KFeIII[FeII(CN)6]) [16,30,31]. The as-prepared PB modified FTO electrodes (PB/FTO) were washed with DIW and were then dried in air for at least 24 h prior to use. A typical charge capacity of the PB thin film was ca. 14.0 mC/cm2 that was estimated from the voltammogram of PB/PW redox reaction in a 0.1 M potassium phosphate buffer, pH 2.5 (scan range = 0.6 to −0.2 V versus Ag/AgCl/sat’d KCl; scan rate, v = 5 mV/s). This sug-
Fig. 1. Schematic apparatus for the in situ spectroelectrochemical Cys detection with a PB/FTO working electrode (RE & CE: reference & counter electrodes).

gest a deposition efficiency of 77.8% (= redox capacity/current density input while deposition = 14.0/18.0).

2.3. In situ optoelectrochemical cysteine detection at a PB/FTO electrode

The schematic apparatus for the in situ optoelectrochemical Cys detection is drawn in Fig. 1, which is very similar to that of our previous work [38]. The EC sensing took place in a UV cell (4 cm × 4 cm × 1 cm) positioned in the UV–vis spectrometer. The UV cell not only filled with an analyte solution but also equipped with a wired PB/FTO working electrode, a homemade reference electrode (Ag/AgCl/sat’d KCl), and a Pt coil counter electrode. The analyte solution was buffered with 0.1 M H₃PO₄ and 0.1 M KH₂PO₄ (pH 2.5) and contained 0, 0.08, 0.04, 0.2, 1, 5, or 25 mM cysteine. The acidity (pH 2.5) was required for stabilizing the deposited PB thin film without the aid of a polymer matrix, carbon paste, or other binder material [16,30,31], as such an additive might interfere with the proof-of-concept EC sensing. Then the EC detection was triggered by cyclic voltammetry (CV) with the potentiostat, scanning back and forth between 0.6 V (PB) and −0.2 V (PW) and/or +0.6 V (PB) and +1.2 V (BG) (versus Ag/AgCl/sat’d KCl) at a scan rate of 5 mV/s. Finally, voltammograms and voltabsorptometric responses at 690 nm [38] obtained at different Cys concentrations were collected simultaneously and analyzed.

3. Results and discussions

3.1. Detection of cysteine at an electrochromic PB/FTO electrode

The oxidized state of PB thin film, Berlin green (BG), can be viewed as an artificial oxidase for some reductant-like biomolecules such as morphine [23] and cysteine [13]. Especially, it was reported that Cys (denoted as RSH) could be oxidized to disulfide cystine (denoted as RSSR) by BG that was then reduced to PB via acquiring both electrons and potassium ions (counter ions), and PB was regenerated under a positive bias, sufficiently larger than the standard potential of the BG/PB redox system (ca. 0.9 V versus Ag/AgCl), for continuous interrogation of Cys [13]. According to the above fact and the electrochemistry of “soluble-form” PB [16,30,31], we propose a working principle for the EC detection of Cys at a PB/FTO electrode, as shown in Fig. 2. The mechanism also can be illustrated by the following equations:

\[ 3\text{PB} \leftrightarrow 3\text{BG} + 2K^+ + 2e^- \]  
\[ \text{(electro-oxidation of PB to BG)} \]  
\[ 3\text{BG} + 2\text{RSH} + 2K^+ \rightarrow 3\text{PB} + \text{RSSR} + 2H^+ \]  
\[ \text{(chemical oxidation of Cys)} \]  

The mechanism is a reversible electrochemical as well as EC reaction followed by an irreversible chemical oxidation. Therefore, we considered that not only electrochemistry but also spectroscopic behavior of PB would be affected by the BG-induced Cys oxidation and thus influenced by the Cys concentration (denoted as [Cys]).

In this study, we triggered the EC detection by a cyclic voltammetric scan (ν = 5 mV/s), and we followed Fig. 2 to simultaneously monitor voltammograms and voltabsorptometric responses at 690 nm, a characteristic wavelength of PB thin film [38], of PB/FTO in buffered Cys solutions. Before that, we had assured cycling stability of PB/FTO in 0.1 M H₃PO₄/KH₂PO₄ (pH 2.5) (both PB/PW and BG/PB redox systems) and observed a decrease in optical density (OD) at 690 nm for both PB-to-PW (0.6 V → −0.2 V) and PB-to-BG (0.6 V → 1.2 V) electrochromism. The data are shown in Fig. 3. The results of different Cys detection methods (voltammetry, voltabsorptometry, and differential voltabsorptometry) degenerated from the same in situ optoelectrochemical experiment are presented and discussed as follows.

3.2. Voltammetric detection of cysteine at PB/FTO

Basically, the cyclic voltammograms (CVs) responded to the entire process of electrocatalytic Cys oxidation at a PB/FTO electrode, i.e., both Eqs. (1) and (2). But, we found that the
CV behavior of PB/FTO was not altered significantly when [Cys] was lower than 40 μM (data not shown). Fig. 4 shows the CVs of PB/FTO measured in the phosphate-buffered solutions containing 0.2, 1, 5, and 25 mM Cys. It can be seen that the PB/PW redox wave (ranged between −0.2 and 0.6 V) is not changed with [Cys], except for the case of [Cys] = 25 mM. Presumably, cysteine (pI = 5.0 [1]) must be positively charged at pH 2.5 and might then compete with reversible, potassium counter ions [25,26] (from 0.1 M KH₂PO₄) for the PB-to-PW redox reaction at such a concentration level ([Cys] = 25 mM). By contrast, the BG/PB redox wave (ranged between 0.6 and 1.2 V) becomes more and more asymmetric and distorted to the anodic side when [Cys] increases. This corresponds to the mechanism mentioned in Section 3.1.

Since Fig. 4 is a typical electrocatalytic data, both amperometric and coulometric detections of Cys can be achieved by plotting calibration curves from the CV responses. Fig. 5 compares two amperometric calibration curves obtained from anodic peak current density and sampling current density at 1.2 V (versus Ag/AgCl/sat’d KCl). It can be found that the sampling current method provides both larger dynamic range and sensitivity than the peak current picking does. This is because more BG forms at 1.2 V than at the peak potential of BG/PB redox wave, although Cys addition tends to shift the peak potential anodically. A more reliable electrochemical calibration curve is to plot the total reacted charge against analyte concentration, as shown in Fig. 6. The coulometric detection is done by mathematical integration of anodic current density ranged between 0.6 to 1.2 V for each CV data obtained at different Cys concentrations. The method can provide accurate concentration calibration as every electron involved in the electrocatalytic process has been counted. The limit of detection (LOD) for both electrochemical methods is ca. 40 μM and corresponds to the value reported in literature [13].

3.3. Voltabsorptometric detection of cysteine at PB/FTO

Since both Cys (RSH) and disulfide cystine (RSSR) are colorless within the visible spectrum, the optical density (OD) change at 690 nm responds to the EC reaction at the PB/FTO electrode. It has been shown in Fig. 3 that the conversion of BG from PB causes OD attenuation at 690 nm. By real-time absorbance measurement during the CV scan, we observed that increasing Cys in the analyte solution would reduce and finally eliminate the OD attenuation. The cyclic voltabsorptometric (CVA, absorbance versus time) data are given in Fig. 7. (Note: The data for PB/PW interchange and for [Cys] < 0.2 mM are not shown.)
Although the initial absorbance shifts at 0.6 V (the PB state) might be due to operating factors like changing analyte solutions, a conclusive finding in Fig. 7 revealed that high [Cys] (>1 mM) interferes the formation of BG from PB. Especially for the case of [Cys] = 25 mM, Berlin green seems never being formed or not detectable throughout the potential-scanning range (0.6–1.2 V). Such a phenomenon would not be discovered using an ordinary amperometric or voltammetric detection and under a facile redox mediator assumption. However, Fig. 7 serves as an evidence to our proposed model in Fig. 2 that the BG/PB EC reaction, Eq. (1), would be affected by the BG-induced Cys oxidation, as seen in Eq. (2).

Therefore, when the rate of Cys-induced BG reduction (an alternative explanation of Eq. (2)) is faster than the electrochemical oxidation of PB, the OD change (ΔOD) at 690 nm can become a measure of [Cys], similar to those reported in PB-based optical sensors [32–37]. Fig. 8 plots ΔOD at 690 nm against [Cys] and can thus play the role of a calibration curve for the optical or absorptometric detection. Different from Figs. 5 and 6, the optical calibration is rather non-linear, and the sensitivity is significantly enlarged when [Cys] > 1 mM.

Besides, it contains both upper and lower detection limits. As the lower limit is ca. 40 μM, the upper limit is 25 mM in the present case attributed to total elimination of BG formation. We believe that the thickness as well as the charge capacity of PB thin film may be another key factor for the dynamic range of the absorptometric detection.

3.4. Differential voltabsorptometric detection of cysteine at PB/FTO

To further investigate how the Cys-induced BG reduction affects the PB-to-BG electrochromism, we differentiate cyclic voltabsorptometric (CVA) responses shown in Fig. 7. Fig. 9 presents the differential cyclic voltabsorptometric (DCVA, dA/dt versus time) data obtained in 0.2, 1, 5, and 25 mM Cys solutions. (Note: no significant difference between DCVA responses for Cys concentration < 0.2 mM is found.) Since the DCVA data can be regarded as optical CV curves [38] and do not respond to thiols, the BG/PB redox mechanism in the presence of Cys can be determined more precisely than that of real, electrochemical CV curves. From Fig. 9, it can be inferred that the PB-to-BG conversion is retarded by the Cys-induced BG reduction, and thus more positive potential is required to regenerate an observable amount of BG. Because of the retardation, the extension of PB-to-BG EC reaction is reduced. For the case of 25 mM Cys, the DCVA response is not observable. And this implies that the seriously distorted anodic wave (0.6 V → 1.2 V) for [Cys] = 25 mM in Fig. 4 responds to Cys oxidation only.

On the basis of the peak shift in the BG/PB redox wave, the potentiometric detection of Cys from the same in situ opto-electrochemical measurement is possible. Fig. 10 presents the relationship between both anodic peak potential values obtained from CV and DCVA data and [Cys]. The trends for both peak potentials are similar, and the values determined from DCVA curves (Fig. 9) are slightly higher than those from CV data (Fig. 4). However, the dynamic range and limit of detection (ca. 200 μM) of the potentiometric calibration curve are obviously worse than those of amperometric, coulometric, and absorptometric detection modes.
4. Conclusions

We have demonstrated the new application of EC thin film for biosensors through the in situ optoelectrochemical Cys detection at a PB/FTO electrode. It has been shown that at least four types of calibration curves can be obtained simultaneously from the EC detection triggered by a single CV scan: they are amperometry (current versus [Cys]), coulometry (charge versus [Cys]), potentiometry (potential versus [Cys]), and absorbometry (absorbance versus [Cys]). This suggests that the EC detection can provide more informative results than sole electrochemical or optical sensing approach does. Table 1 summarizes the estimated limits of detection for each sensing scheme in the present work. Since the normal concentration of plasma Cys in a healthy person is in the range between 152.8 and 378.0 μM, our EC detection is applicable for clinical purposes. Moreover, such a sensing method can be coupled with liquid chromatography [13,14,39] or other separation techniques to detect other organic thiols, Cys-rich proteins, or thiol-labeled biomolecules for many purposes. Besides, there exist many other EC thin film materials, like transition metal oxides and conducting polymers [15,17], to provide a variety of choices of scanning potential range and characteristic wavelength for different EC biosensing applications against different biomolecular targets.

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Table 1
Detection limits estimated for each sensing scheme in this work

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<thead>
<tr>
<th>Method</th>
<th>Lower detection limit (μM)</th>
<th>Upper detection limit</th>
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<tbody>
<tr>
<td>Amperometric detection</td>
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<td>Not observed</td>
</tr>
<tr>
<td>Coulometric detection</td>
<td>40</td>
<td>Not observed</td>
</tr>
<tr>
<td>Potentiometric detection</td>
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<td>within 5–25 mM</td>
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<tr>
<td>Absorptometric detection</td>
<td>40</td>
<td>25 mM</td>
</tr>
</tbody>
</table>

References


Biographies

Lin-Chi Chen received his BS degree in chemical engineering from National Taiwan University, Taipei, Taiwan in 1997. He received PhD degree in chemical engineering from National Taiwan University in 2001. He was a postdoctoral research fellow in the Institute of Biomedical Sciences at Academia Sinica, Taipei, Taiwan, from 2002 until 2005. Currently, he is an assistant professor jointly appointed by the Department of Bio-industrial Mechatronics Engineering and Bioenergy Research Center at National Taiwan University. His research interest includes electrochromic devices, biosensors, biochips and antibody mimics.

Kuo-Chuan Ho received BS and MS degrees in chemical engineering from National Cheng Kung University, Tainan, Taiwan, in 1978 and 1980, respectively. In 1986, he received the PhD degree in chemical engineering at the University of Rochester. The same year he joined PPG Industries, Inc., first as a senior research engineer and then, from 1990 until 1993, as a research project engineer. He has worked on the electrochemical properties of various electrode materials, with emphasis on improving the performances of sensor devices. Following a 6-year industrial career at PPG Industries, Inc., he joined his alma mater at National Cheng Kung University in 1993 as an associate professor in the Chemical Engineering Department. In 1994, he moved to the Department of Chemical Engineering at National Taiwan University. Currently, he is a professor jointly appointed by the Department of Chemical Engineering and Institute of Polymer Science and Engineering at National Taiwan University.