Cyclic Voltammetric and Electrochemical Simulation Studies on the Electro-Oxidation of Catechol in the Presence of 4, 4-bipyridine

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To cite this article:

Tekalign Kasa, Theodros Solomon. Cyclic Voltammetric and Electrochemical Simulation Studies on the Electro-Oxidation of Catechol in the Presence of 4, 4-bipyridine. *American Journal of Physical Chemistry*. Vol. 5, No. 3, 2016, pp. 45-55. doi: 10.11648/j.ajpc.20160503.11

Received: February 21, 2016; **Accepted:** March 9, 2016; **Published:** April 16, 2016

Abstract: The studies were made using cyclic voltammetry on a glassy carbon electrode in aqueous solution containing phosphate buffer solution as supporting electrolyte. The purpose of the investigation was to carry out a quantitative detailed study of the electrochemical oxidation of catechol in the presence of 4, 4-bipyridine in aqueous solution. The electrooxidation of catechol produces a very reactive intermediate, *o*-benzoquinone, which subsequently reacts with 4, 4-bipyridine in 1, 4-Michael addition reaction to form the corresponding catechol derivative product. The kinetic data were extracted from cyclic voltammograms with the help of digital simulation. The results of the study show that catechol is oxidized in aqueous phosphate buffer solution to a very reactive intermediate *o*-benzoquinone.

Keywords: Cyclic Voltammetry, EC Mechanism, Catechol, Benzoquinone, 4, 4-bipyridine

1. Introduction

Electrochemistry provides very interesting and versatile means for the study of chemical reactions. The majority of organic electrode reactions are characterized by the generation of a reactive intermediate at the electrode by electron transfer and subsequent reactions typical for that species. The main goal of the electrochemical studies is the elucidation of the sequence of electron transfer and chemical reactions that occur near the electrode surface and its applications to electro-synthesis of organic compounds. Among the organic compounds, catechol can be easily oxidized to the corresponding reactive *o*-benzoquinone mainly due to its low oxidation potential [1]. Catechol has the molecular formula of C₆H₆O₂ and structure which is shown in the scheme 1 below.

Catechol is well known in biological systems often as a reactive center of electron transfer in the structure of many natural compounds and biologically reactive molecules capable of exhibiting both anti- and pro-oxidant behavior [2]. Catechol has a great importance in both biological and environmental analysis because of its excellent

electrochemical activity and it can be used for the characterization of different analytical methods. It is used in a variety of applications including photography, dyeing, agrochemicals, rubber, plastic production antiseptic, dyestuffs, electroplating, specialty inks, antioxidants and light stabilizers, in organic synthesis, pharmaceutical industry, antibiotics, and flavor compounds such as vanillin, precursors are essential; they are the starting material for the production of the desired compound [1, 3-6].

Catechol

Scheme 1. Structure of catechol.

The electrochemical oxidation of catechol in aqueous solutions and in the presence of nucleophiles shows that catechol undergoes 1, 4-Michael addition reactions according to the so-called EC mechanism with the consumption of 2 electrons per molecule of catechol, and a conversion to catechol derivatives [7]. The purpose of the investigation was to carry out a quantitative detailed study of the electrochemical oxidation of catechol in the presence of 4, 4-bipyridine in aqueous solution.

2. Experimental Parts

2.1. Chemicals

The chemicals used were catechol (BDH Poole, England), 4, 4-bipyridine (Aldrich-chemie, Germany), HCl (Riedel-deHaen, Germany), NaOH (BDH Poole, England), sodium dihydrogen phosphate ($\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$, Riedel-deHaen, Germany) and disodium hydrogen phosphate ($\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$, Techno Pharmchem, India).

2.2. Instrumentation

The cyclic voltammetry apparatus used was BAS 100A electrochemical analyzer [Bio-analytical systems (BAS), USA], coupled to a Dell computer (Pentium 4). A JENWAY 3510 pH Meter (Barloworld Scientific Ltd, Dunmow, Essex, U.K.) and Electronic Balance (Model: LA 204) were used. The three electrodes used in the voltammetry experiments were the glassy carbon electrode (3mm diameter) as working electrode, platinum wire as an auxiliary electrode and Ag/AgCl (3MKCl) as reference electrode, all from BAS. The working electrode (glassy carbon electrode) (GCE) was polished in each set of experiments with aluminum oxide powder ($0.3\mu\text{m}$) on a polishing cloth and followed by distilled water and were carried out at room temperature.

2.3. Electrochemical Digital Simulation

Digital simulation is a useful method in evaluation of complicated electrode reactions and used to visualize and verify electrochemical mechanisms and also used to determine the normalized current-over voltage curves for reversible as well as for quasi reversible reactions [8, 9]. By the development of simulation software cyclic voltammetry has become a very powerful technique [10]. The experimental parameters entered for digital simulation consisted of the following; $E_{\text{start}} = -0.2\text{ V}$, $E_{\text{switch}} = 0.8\text{ V}$ and $T = 298\text{ K}$. The concentration of catechol and 4, 4-bipyridine were 1mM. The transfer coefficient (α) was assumed to be 0.5. The homogeneous rate constant (k_f) of the chemical reaction of o-benzoquinone with 4, 4-bipyridine was estimated based on an EC mechanism by fitting the simulation data with experimental cyclic voltammogram data at various scan rates and pHs, while the heterogeneous rate constant (k^0) for the electrochemical oxidation of catechol to o-benzoquinone was estimated based on an E- mechanism of

the simulation [11]. The best simulation fits were obtained for the E_rC_i mechanism.

3. Result and Discussion

3.1. Cyclic Voltammetry of Catechol

The number of electrons and protons involved in the oxidation of catechol can be determined from the separation peak potentials and from plots of E^o vs pH and E_{pa} vs pH respectively. The formal potential (E^o), which is approximated by the midpoint potential (E_{mid}) between the anodic and cathodic peaks is given by;

$$E^o = E_{pH_0}^o - \left(\frac{2.303mRT}{nF} \right) pH$$

Where $E_{pH_0}^o$ is the formal potential at pH_0 , and R , T , and F have their usual meanings and n and m are number of electron and proton respectively. The values of E^o evaluated from the midpoint potential between the anodic and cathodic peaks, (E_{mid}).

Both E^o and E_{pa} were shifted to negative potentials with the slope of 62 mV/pH with the correlation function R ($R = -0.999601$) and 63.3 mV/pH with the correlation function R ($R = -0.999004$) respectively. In all cases, the slopes are in good agreement with the theoretical slope ($2.303mRT/nF$) of 59 mV/pH with $m \approx 2$ and $n = 2$, where n and m are number of electron and proton respectively [12].

3.2. Effect of PH

Figure 1. Typical cyclic voltammograms for 1 mM catechol at various pHs a) 4, b) 5, c) 6, d) 7 and e) 8 at a scan rate of 50 mV/s.

On increasing the pH of supporting electrolyte both the anodic and cathodic peak potentials shifted towards more negative potential [13] as shown in the Figure 1. This is expected because of the participation of proton in the oxidation reactions and these shifts of potential indicate that the electron transfer process from catechol to o-benzoquinone occurs at glassy carbon electrode easily at higher pH than at lower pH. The electrooxidation of catechol was found to be pH dependent. In acidic and neutral media catechol showed a well developed quasi-reversible wave as indicated in Figure 1

below. Under these conditions, a peak current ratio $|i_{pc}/i_{pa}|$ of nearly unity can be considered as a criterion for the stability of o-benzoquinone produced at the surface of the electrode under the experimental conditions. In basic solutions, the peak current ratio is less than unity and decreases with increasing pH. These shifts can be associated to the coupling of the anionic or dianionic forms of catechol with o-benzoquinones (dimerization reactions) [2].

Figure 2 below shows that the anodic peak current increases as the pH increases and reaches the highest peak at (pH = 7) and then decreases. The highest anodic peak current (at pH = 7) was taken as the optimum pH for the electrooxidation of catechol.

Figure 2. i_{pa} as a function of pH for 1 mM catechol at a scan rate of 50 mV/s.

3.3. Effect of Scan Rate

As shown in Figure 3, when the scan rate increases, both the anodic and cathodic peaks increase; the anodic peak shifts towards less positive potentials while the cathodic peak shifts

towards more positive potentials. Overall, the current ratios are almost unity and the peak separation between the anodic and cathodic peak potential at scan rate of 50 mV/s is 69 mV/n which is greater than 59 mV/n and it is one of the indications that the system is a quasi-reversible process.

A peak current ratio $|i_{pc}/i_{pa}|$ of nearly unity, particularly during the repetitive recycling of potential, can be considered as criteria for the stability of o-benzoquinone produced at the surface of electrode under the experimental conditions. It means that any hydroxylation or dimerization reactions are very slow which was observed in the time-scale of cyclic voltammetry [2, 14, 15].

Figure 3. Cyclic voltammograms of 1mM catechol at scan rates of; 30, 50, 100, 150, 200, 250, 300 and 400 mV/s at phosphate buffer solution of pH=7.

DETERMINATION OF HETEROGENEOUS RATE CONSTANT (k^0) AT VARIOUS SCAN RATES

Figure 4. Typical cyclic voltammograms of 1mM catechol at various scan rates at pHs 7. (a) Experimental; (b) simulated.

DETERMINATION OF HETEROGENEOUS RATE CONSTANT (k^0) AT VARIOUS pHs

Figure 5. Typical cyclic voltammograms of 1mM catechol at various pHs at a scan rate of 50mV/s. (a) experimental; (b) simulated.

Figure 6. Variation of experimental (a) and simulated (b) heterogeneous rate constant (k°) as the function scan rate at pH 7.

Figure 7. Variation of experimental (a) and simulated (b) heterogeneous rate constant (k^o) as the function pH at 50 mV/s.

As can be seen from the data in Figure 6 and 7, there is a good correlation between the heterogeneous rate constant (k^o) calculated using the Nicholson and Shain method and the heterogeneous rate constant (k^o) obtained using digital simulation at various scan rates and pHs.

3.4. Cyclic Voltammetry of Catechol in the Presence of 4, 4-Bipyridine

The cyclic voltammogram of catechol in aqueous solution of 0.2 M phosphate buffer at pH = 7 shows one anodic and the corresponding cathodic peak. As 1mM 4, 4-bipyridine is added to 1mM catechol, the height of the cathodic and anodic peak currents decreased, while the potential of anodic peak to some extent shifted towards a less positive potential as shown in Figure 8 below. The shift of the anodic peak potential in the presence of 4, 4-bipyridine may due to the coupled homogeneous chemical reaction product formed at the surface of the electrode.

Figure 8. Cyclic voltammograms of 1mM catechol in the absence (a) and in the presence of 1mM 4, 4-bipyridine (b), and that of a 1 mM 4, 4-bipyridine alone (c) at the glassy carbon electrode in 0.2 M phosphate buffer at different pH at a scan rate of 50 mV/s.

3.5. Effect of pH

According to figure 9 the cyclic voltammogram of 1mM catechol in the presence of 4, 4-bipyridine shows that the anodic and cathodic peak potentials shifted towards less positive values as the pH increased. As shown in Figure 10 below also the peak current ratio $|i_{pc}/i_{pa}|$ of catechol in the presence of 4, 4-bipyridine increased with increasing pH until it reached a maximum at pH 6 and then decreased after this pH. This can be related to protonation of the nucleophile 4, 4-bipyridine at lower pHs and its deprotonation at higher pHs. In other words, in acidic and neutral medium, the peak current ratio $|i_{pc}/i_{pa}|$ of nearly unity can be considered as criteria for the stability of o-benzoquinone produced at the surface of the electrode.

However, in basic solutions, the peak current ratio is less than unity and decreases with increasing pH. These changes can be related to the coupling of the anionic or dianionic forms of catechols with o-quinones (dimerization reactions) [16, 17].

Figure 9. Cyclic voltammograms of catechol in the presence of 4, 4-bipyridine at glassy carbon electrode at a scan rate of 50mV/s and various pHs: a) 4, b) 5, c) 6, d) 7 and e) 8.

Figure 10. Current ratio $|i_{pc}/i_{pa}|$ versus pH for 1mM catechol in the presence of 1mM 4, 4-bipyridine.

Figure 11 below shows that the anodic peak current decreases as pH increases and attain its highest value at (pH = 6) and then slightly decreases with pH. The highest anodic peak current occurred at pH = 6 and this is the optimum pH for electrooxidation of catechol in the presence of 4, 4-bipyridine.

Figure 11. Anodic peak current i_{pa} as a function of pH for catechol in the presence of 4, 4-bipyridine at a scan rate of 50 mV/s.

3.6. Effect of Scan Rate

Figure 12. Cyclic voltammograms of 1mM catechol in the presence of 4, 4-bipyridine at scan rates of (10-300) mV/s at phosphate buffer solution of pH=6.

The effect of the scan rates (10-400 mV/s) on the electrooxidation of catechol in the presence of 4, 4-bipyridine was studied. Figure 12 show that both the magnitude of anodic and cathodic peak potentials increased with increasing scan rate. The peak current ratio $|i_{pc}/i_{pa}|$ increased as the scan rate increased and this indicates that a chemical reaction occurred between 4, 4-bipyridine and o-benzoquinone [18].

Figure 13. Current ratio $|i_{pc}/i_{pa}|$ as function of scan rate for 1mM catechol in the presence of 1mM 4, 4-bipyridine.

Figure 14. Current ratio $|i_{pa}/i_{pc}|$ versus scan rate for catechol in the presence of 4, 4-bipyridine.

Figure 15. Current function $i_{pa}/v^{1/2}$ at different scan rate (v) for catechol in the presence of 4, 4-bipyridine at pH = 6.

The cyclic voltammogram of catechol in the presence of the nucleophile at different scan rates shows an increasing of peak current ratio $|i_{pc}/i_{pa}|$, decreasing of $|i_{pa}/i_{pc}|$ and the decreasing of current function $i_{pa}/v^{1/2}$ with increasing scan rate. As shown in Figures 13, 14 and 15; it is a clear

indication of coupled chemical reaction following an E_rC_i mechanism [19] and also the reactivity of deprotonated 4, 4-bipyridine towards o-benzoquinone.

Determination of homogeneous rate constant (k_f) by using electrochemical simulation method at various scan rates.

Figure 16. Separation of peak potentials (ΔE_p) as a function of scan rate (v) for 1 mM catechol in the presence of 1 mM 4, 4-bipyridine at (pH = 6).

Figure 17. Cyclic voltammograms for 1 mM catechol in the presence of 4, 4'-bipyridine at pH= 6 at different scan rates. (a) Experimental; (b) simulated.

As shown in Figure 18 below the k_f values from Nicholson's working curve were in good agreement with the k_f values of the electrochemical simulation results.

Figure 18. Variation of experimental (a) and simulated (b) homogeneous rate constant (k_f) as a function scan rate at pH of 6.

Determination of homogeneous rate constant (k_f) by using electrochemical simulation method at various PHs

Figure 19. Typical cyclic voltammograms of 1mM catechol in the presence of 1mM 4, 4'-bipyridine at various pHs at a scan rate of 50mV/s. (a) experimental; (b) simulated.

Figure 20. Calculated (a) and simulated (b) homogeneous rate constant (k_f) as a function pH at 50 mV/s.

As shown in the Figure 20 above the experimental values of k_f at various pHs at the scan rate of 50 mV/s were in good agreement with the k_f values of simulation values.

Scheme 2. Proposed mechanism for electrooxidation of catechol in the presence of 4, 4'-bipyridine.

4. Conclusion

The results of the study show that catechol is oxidized in aqueous phosphate buffer solution to a very reactive intermediate *o*-benzoquinone. The oxidation of catechol in the absence of 4, 4-bipyridine showed properties of a quasi-reversible two-electron two-proton process, whereas, the oxidation of catechol in the presence of 4, 4-bipyridine follows an E_rC_i mechanism with an irreversible chemical reaction. The *o*-benzoquinone is attacked in 1, 4-Michael addition reaction by 4, 4-bipyridine to form catechol derivative. The kinetic parameters for the electrooxidation of catechol in the absence and presence of 4, 4-bipyridine at different scan rates and pHs were studied using cyclic voltammetry and electrochemical digital simulation.

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