

Electrochemical Oxidation of Caffeic and Ferulic Acid Derivatives in Aprotic Medium

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O comportamento eletroquímico em função da estrutura de uma série de derivados dos ácidos cafeico e ferúlico, assim como de seus precursores catecol e guaiacol, eletroquimicamente ativos, foi avaliado por voltametria cíclica. Os resultados revelaram que as condições experimentais são a chave para as mudanças no mecanismo de oxidação do guaiacol e ácido ferúlico. A oxidação eletroquímica dos derivados amida do ácido ferúlico revelou que o átomo de nitrogênio desempenha um papel importante na derivatização da superfície do eletrodo. Além disso, a atividade sequestradora de radicais dos compostos, avaliada através da porcentagem de inibição do radical 2,2'-difênil-1-picril-hidrazila, mostrou uma boa correlação com os potenciais de oxidação.

We studied the electrochemical behaviour as a function of the structure of a series of caffeic and ferulic acids derivatives as well as their corresponding redox moieties catechol and guaiacol by cyclic voltammetry. Results revealed that the medium is key for changes in the oxidation mechanism of guaiacol and ferulic acid. Electrochemical oxidation of the ferulic acid amide derivatives revealed that the nitrogen atom plays an important role in the derivatization of the electrode surface. In addition, radical scavenging activity of the compounds evaluated through the percentage of inhibition of the 2,2'-diphenyl-1-picrylhydrazyl radical showed a good relationship with the oxidation potentials.

Keywords: cyclic voltammetry, hydroxycinnamic acids, caffeic acid phenethyl ester, antioxidant activity, DPPH radical

Introduction

Caffeic (CA) and ferulic (FA) acids and their corresponding esters are widely distributed in plants and also in propolis from diverse geographic zones of the world.^{1,2} During the last years, these compounds have received the attention of numerous research groups since it has been suggested that some of them have important biological properties.³⁻¹⁸ Special interest has received caffeic acid phenethyl ester (CAPE), which is one of the most active components isolated from honeybee propolis.

It has been demonstrated that CAPE is an efficient free radical scavenger and an inhibitor of some enzymes as cyclooxygenase,¹⁷ lipoxygenase, and the HIV-1 integrase.^{5,18} Additionally, it possesses antiinflammatory and antitumor activities.¹⁴⁻¹⁶ Particularly, the antitumor activity of CAPE has been related to its capacity to induce apoptosis and also to modulate the cellular redox states.^{19,20}

Since the biological activity of CA and FA is related to the electron transfer processes, several mechanistic studies on the electrochemical oxidation of these acids have been documented in the literature.²¹⁻²⁵ However, to the best of our knowledge, no study has been focused to understand the electrochemical behavior of its corresponding esters

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or amides, nevertheless some compounds such as CAPE and caffeic acid phenethyl amide (CAPEA) have shown even higher *in vitro* antioxidant activity respect to that of CA.^{9,10,26}

In the present work the anodic oxidation of a series of caffeic and ferulic acids derivatives (Figure 1) in aprotic medium is reported, as well as for their corresponding redox moieties catechol (C) and guaiacol (G). Some mechanistic details of the redox behavior of the complete series were investigated and are also discussed here, since they may be related to the known antioxidant properties of this kind of compounds in biological systems. The radical scavenging activity of the ten compounds was also evaluated using the DPPH assay, in order to analyze if there is a relationship between this property and the corresponding potential oxidation value.

Experimental

Chemicals

Tetrabutylammonium hexafluorophosphate 98% (n-Bu₄NPF₆), catechol (C) (99%), guaiacol (G) (> 98%),

ferulic acid (FA) (99%), dihydrocaffeic acid (DHCA) (99%), caffeic acid (CA) (99%), and 2,2'-diphenyl-1-picrylhydrazyl radical (DPPH) were purchased from Sigma-Aldrich. Acetonitrile (AN) (content of H₂O < 0.1%), dimethylsulfoxide (DMSO) (content of H₂O < 0.1%) spectra grade (Merck Uvasol) and ethanol chromasolv® were used as solvents. Caffeic acid phenethyl ester (CAPE) was prepared according to the procedure reported by Grunberger.²⁷ All of the amides were synthesized from the free acid and the corresponding amine using (benzotriazol-1-yloxy)tris(dimethylamino)phosphonium hexafluorophosphate (BOP) as coupling reagent, as has been described before.⁹

Free radical scavenging activity (DPPH assay)

The free radical scavenging activity (FRS) was measured according to the following procedure: 500 μL of a 2,2'-diphenyl-1-picrylhydrazyl radical (DPPH) ethanolic solution (300 μmol L⁻¹) was mixed with the same volume of the test compound solution (17.5 μmol L⁻¹). The reaction mixtures were vigorously vortexed for 10 s. Absorbance of the resulting solutions was read at

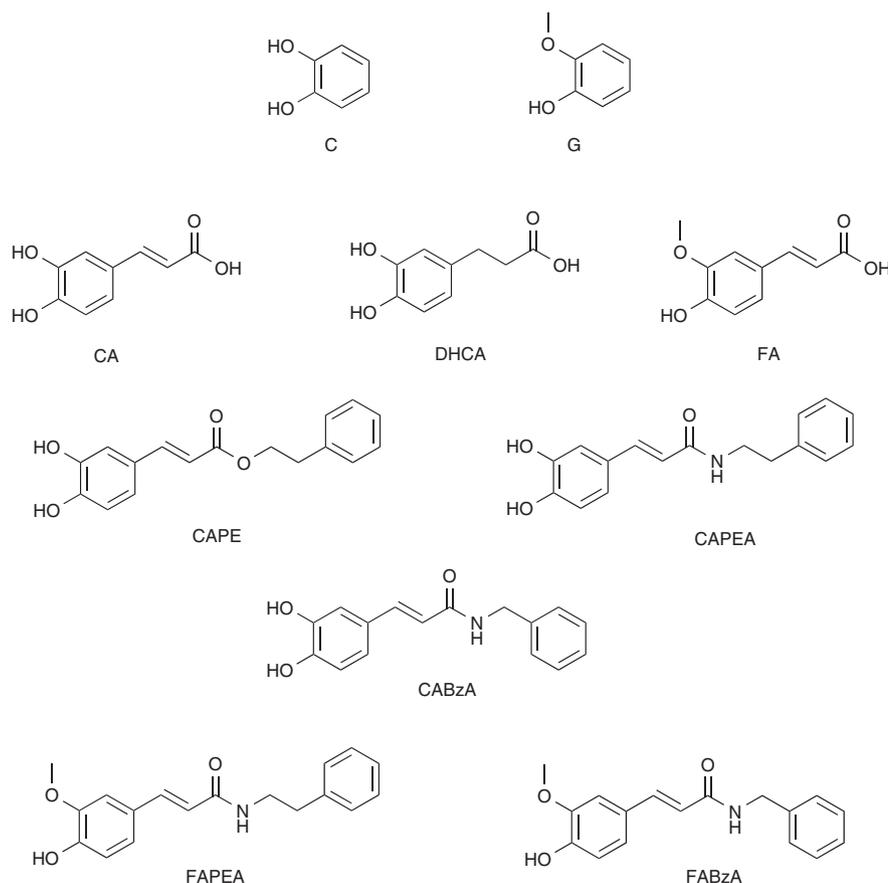


Figure 1. Compounds studied in the present work.

517 nm in a spectrophotometer (Varian, CARY 100), after 30 min of incubation at room temperature in darkness. The scavenging activity was determined by comparing the absorbance with that of the blank containing only 2,2'-diphenyl-1-picrylhydrazyl radical (DPPH) and ethanol. Results were expressed as inhibition percentage.³

Instrumentation and electrodes

Cyclic voltammetry and coulometry experiments were performed in a radiometer potentiostat 1MT102-DEA332 with positive feedback resistance compensation. A three-electrode cell was used in the electrochemical experiments. The working electrode was a 3 mm diameter glassy carbon disk (Sigradur G from HTW, Germany). This electrode was polished with 1 μm alumina powder and rinsed in ultrasound bath with distilled water and ethanol before each run. Platinum mesh was used as counter electrode and saturated calomel electrode (SCE) as reference. The reference electrode was connected to the cell through a salt bridge containing the same supporting electrolyte concentration as the working solution. All electrochemical experiments were performed at 25 $^{\circ}\text{C}$ under an argon atmosphere. Coulometry was carried out with a glassy carbon rotating rod electrode under constant rotation (1000 rpm) and controlled potential.

Results and Discussion

Electrochemical study of the oxidation process of C, G, CA, DHCA, CAPE and FA

The electrochemical behaviour in acetonitrile of caffeic acid (CA), dihydrocaffeic acid (DHCA), caffeic acid phenethyl ester (CAPE), guaiacol (G), and ferulic acid (FA) showed a typical chemically irreversible oxidation wave (I) and the corresponding broadened reduction wave (II) in the reverse scan. The voltammogram of CAPE is shown in Figure 2a as a representative example of this behaviour. The bielectronic nature of the oxidative process of CAPE was confirmed by coulometry ($n_{\text{app}} = 2.02$). Since the current peak intensity (i_p) for the rest of the systems was similar to the current peak intensity recorded for CAPE at the same scan rate and concentration, it was deduced that the electron stoichiometry was also two in all the cases.

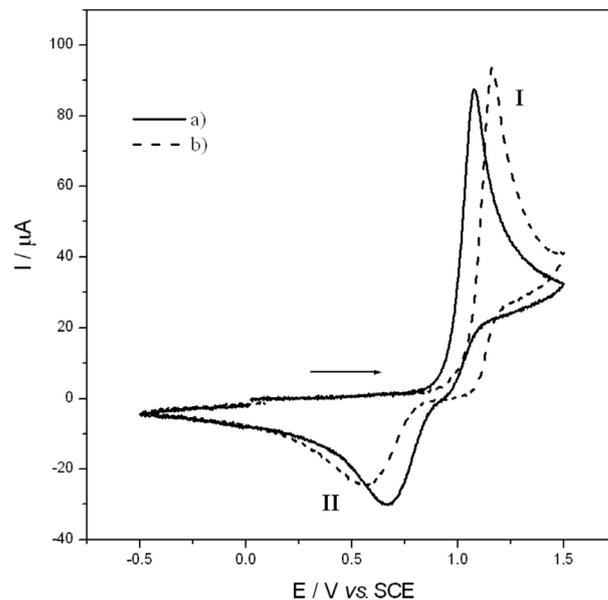
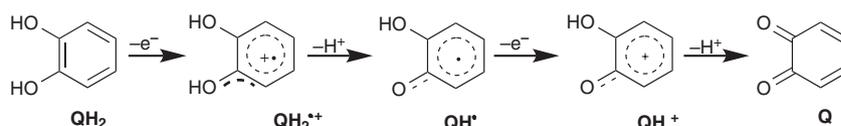


Figure 2. Cyclic voltammetry in acetonitrile + 0.2 mol L⁻¹ *n*-Bu₄NPF₆, on glassy carbon electrode (3 mm ϕ) at 2 mmol L⁻¹ and 0.1 V s⁻¹ of a) CAPE and b) FA.

The electrochemical responses of CA, DHCA, and CAPE under these experimental conditions are similar to that found for catechol, and are in good agreement with the anodic oxidation of hydroquinone (an isomer of catechol) reported before,²⁸⁻³² which follows an ECE mechanism that involves the loss of one electron in the system, followed by a fast deprotonation reaction and, finally, the loss of a second electron. Scheme 1 shows the complete mechanism, which is the starting point for the following discussion. A further deprotonation reaction is also included as the last step, since additional spectroscopic studies have suggested that the *o*-quinone (Q) is obtained as the final product.²⁸

In Figure 2b, it can also be observed that the voltammogram of FA did not show significant differences with respect to the electrochemical responses of catechol and caffeic acid derivatives, such as CAPE (Figure 2a). Consequently, a two-electron mechanism is also involved in the oxidation of G and FA in acetonitrile. Despite the fact that the electrochemical oxidation of both compounds is still unclear, it is known that the mechanism involves the *o*-quinone form, which has been identified as the main product obtained during the oxidation of FA.^{25,28}

When the scan rate is increased in the case of CA, FA, and CAPE, the peak potential I is anodically shifted,



Scheme 1.

showing a linear variation with the logarithm of the scan rate of 31, 30, and 42 mV dec⁻¹, respectively (Table 1). The analysis of a transition of the mechanism controlled by the follow-up chemical reaction to a mechanism where the electron transfer is rate determining, has been well characterized.³³ Thus, at 25 °C, this transition involves a variation of $\partial E_p/\partial \log \nu$ from 29.6 to 59.2 mV dec⁻¹. The results presented here suggest that in the case of CA and FA the initial electron and proton transfer reactions are consecutive, and they occur under a pure kinetic control of the radical cation deprotonation. In contrast, the substitution of the acidic proton of the caffeic acid by the phenethyl group induces an increase in the rate of the deprotonation chemical reaction in the case of CAPE and, as a consequence, the global kinetic occurs under mixed control.

Table 1. Variation of oxidation potentials vs. the logarithm of the scan rate

Compound	$\partial E_p/\partial \log \nu$
C	38
CA	31
DHCA	39
CAPE	42
G	40
FA	30

On the other hand, $\partial E_p/\partial \log \nu$ was analyzed for C, DHCA and G (Table 1), in order to know the effect that the conjugation between the aromatic ring and the double bond exerts on the global kinetic oxidation mechanisms of CA and FA systems. The variation of $\partial E_p/\partial \log \nu$ was 38, 39, and 40 mV dec⁻¹ for C, DHCA, and G, respectively. This observation suggests that the oxidation mechanism for these compounds proceeds under mixed kinetic control.³³

In addition, we found that the difference between the oxidation wave I and the reduction wave II for the conjugated systems CA, CAPE, and FA is smaller (326, 355, and 447 mV, respectively) than in the case of the non-conjugated compounds C, DHCA, and G (530, 552, and 677 mV, respectively), since there is a higher stabilization of the radical cations coming from the oxidation of the

catechol and guaiacol moieties due to the presence of the conjugated double bond (Scheme 2).

On the other hand, Astudillo *et al.*³² reported that hydrogen bonding acceptor solvents play an important role on the anodic oxidation of hydroquinone. They proposed that DMSO interacts with hydroquinone through hydrogen bonding associations, causing the shift of the oxidation peaks toward less positive potentials. Taking into account the structural similarity between hydroquinone and the catechol moiety in C, CA, DHCA, and CAPE, we also analyzed the electrochemical behavior of these compounds in DMSO (Figure 3).

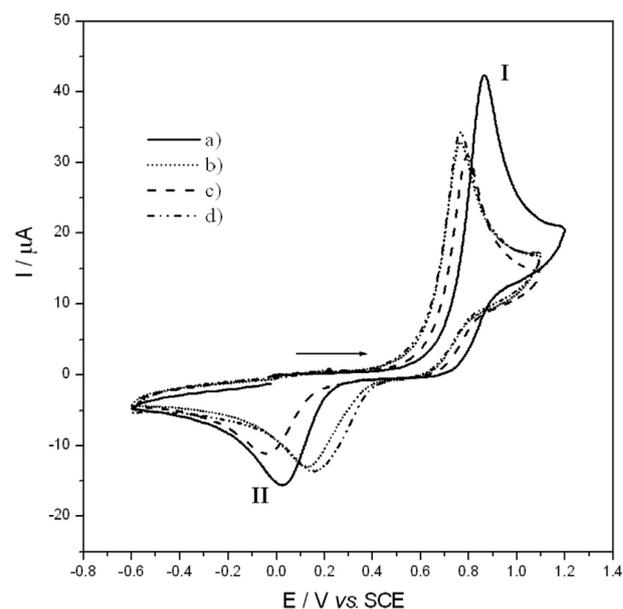
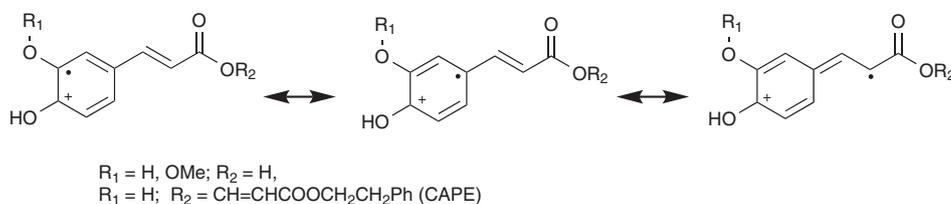


Figure 3. Cyclic voltammetry in DMSO + 0.2 mol L⁻¹ *n*-Bu₄NPF₆, on glassy carbon electrode (3 mm ϕ) at 2 mmol L⁻¹ and 0.1 V s⁻¹ of a) C, b) CA, c) DHCA and d) CAPE.

As in the case of hydroquinone, the peak potentials of C, CA, DHCA, and CAPE presented a displacement of >250 mV towards less anodic potentials in DMSO, in comparison with the measurements obtained in AN (Table 2).

The behaviour suggests that under these conditions the interaction between the catechol moiety (QH₂, scheme 1) of the four compounds with DMSO decreases the dissociation energy of the O–H bonds, making the proton



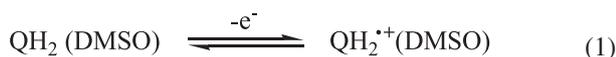
Scheme 2.

Table 2. Oxidation potentials of C, CA, DHCA, and CAPE in AN and DMSO^a

Compound	E_p (AN) / V vs. SCE	E_p (DMSO) / V vs. SCE
C	1.11	0.86
CA	1.08	0.78
DHCA	1.07	0.79
CAPE	1.01	0.73

^aAll oxidation potentials were obtained at 2 mmol L⁻¹ and at 0.1 V s⁻¹.

transfer reaction more favorable in the second step of the oxidation mechanism (reactions 1-3). This observation can be explained considering the fact that the proton affinity is higher in DMSO than in AN, and that the proton transfer in pure DMSO is carried out into a hydrogen bonding complex formed between the hydroxyl groups of catechol and the DMSO molecules.^{32,34}



Additionally, the variation of E_p vs. $\log \nu$ for all the compounds (38.4–41.9 mV dec⁻¹) indicates a mixed kinetic control of steps 1 and 2 on the global reaction, which is not affected by the structural differences among the four compounds (C, CA, CAPE, and DHCA). This is a consequence of the competition between deprotonation and radical cation delocalization, where the most favored process is the first one.

Since hydrogen bonding acceptor solvents play an important role on the oxidation mechanisms of catechol and caffeic acid derivatives, we also carried out the cyclic voltammetry of G and FA in DMSO (Figure 4). A two-electron chemically irreversible oxidation wave I was observed in both cases, however the voltammogram of FA shows additional oxidation and reduction waves, which

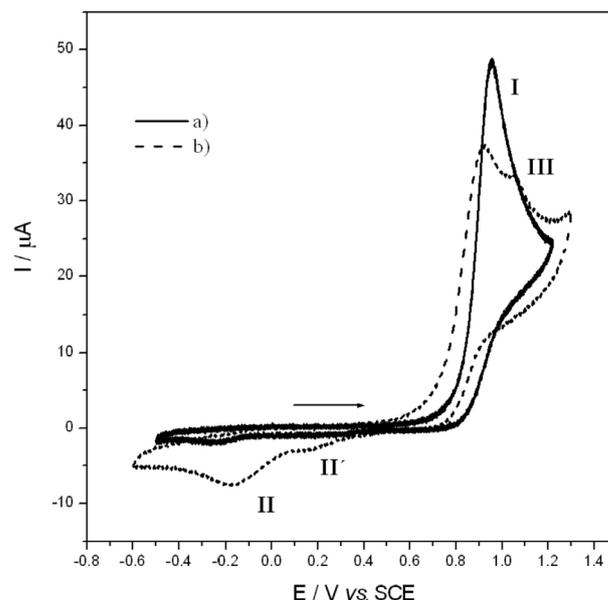
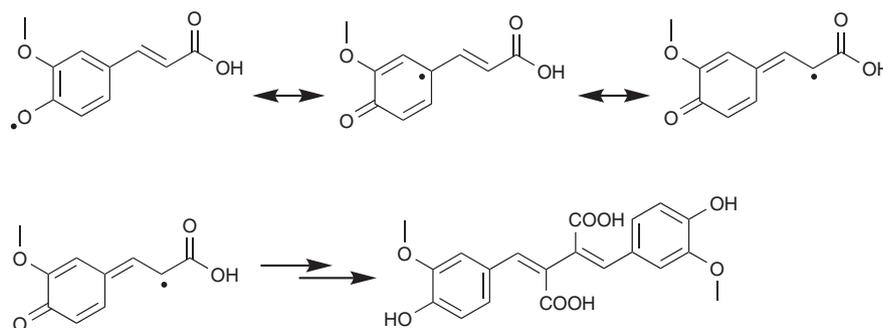


Figure 4. Cyclic voltammetry in DMSO + 0.2 mol L⁻¹ *n*-Bu₄NPF₆, on glassy carbon electrode (3 mm ϕ) at 2 mmol L⁻¹ and 0.1 V s⁻¹ of a) G and b) FA.

were labeled as III and II' respectively in the Figure 4b. In addition, the reduction wave (II) intensity of G was smaller than the recorded for FA.

On the other hand, the analysis of the variation of E_p vs. $\log \nu$ on the wave I from the voltammogram of FA showed a slope of 23 mV dec⁻¹. This value is consistent with a radical-radical dimerization mechanism, since it has been previously reported that dimeric species can be formed during the chemical and electrochemical oxidation of neutral and dianionic hydroxycinnamic acid, respectively.^{28,33,35} Taking into account this result, it was proposed the dimerization mechanism of FA in DMSO as follow (Scheme 3).

Thus, the wave III observed in the cyclic voltammetry of FA corresponds to the oxidation of the guaiacol moieties present in the resulting dimer (Scheme 3). The absence of a second oxidation wave in the voltammogram of G and a slope of 36 mV dec⁻¹ calculated from the E_p vs. $\log \nu$



Scheme 3.

variation from the corresponding wave I confirm this proposal. In addition, the reduction waves II observed in the voltammograms of both G and FA in DMSO (Figure 4), showed lower intensities respect to the data obtained in AN, which suggest that *o*-quinone is obtained in small quantities during their oxidation processes under these experimental conditions. However, the potential scan of the electrochemical responses of G and FA from -2 to 1.5 V (Figures 5 and 6 respectively), showed an additional

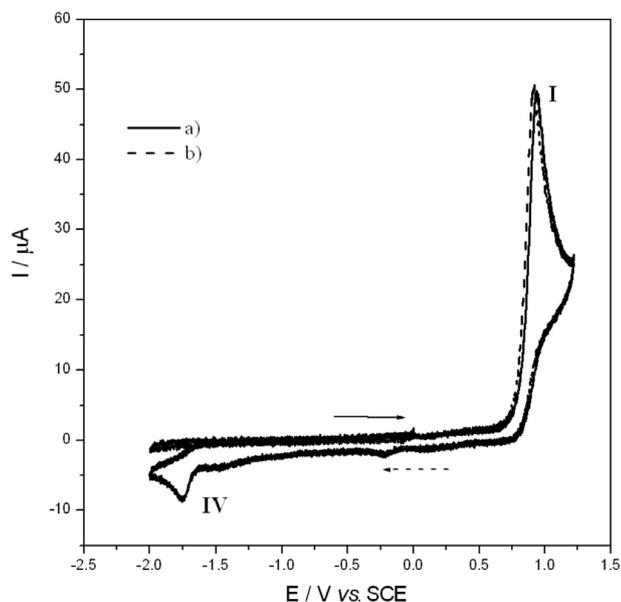


Figure 5. Cyclic voltammetry of G at 2 mmol L^{-1} in DMSO + 0.2 mol L^{-1} $n\text{-Bu}_4\text{NPF}_6$, on glassy carbon electrode ($3 \text{ mm } \phi$). Scan from 1.5 to -2.0 V. a) anodic direction and b) cathodic direction.

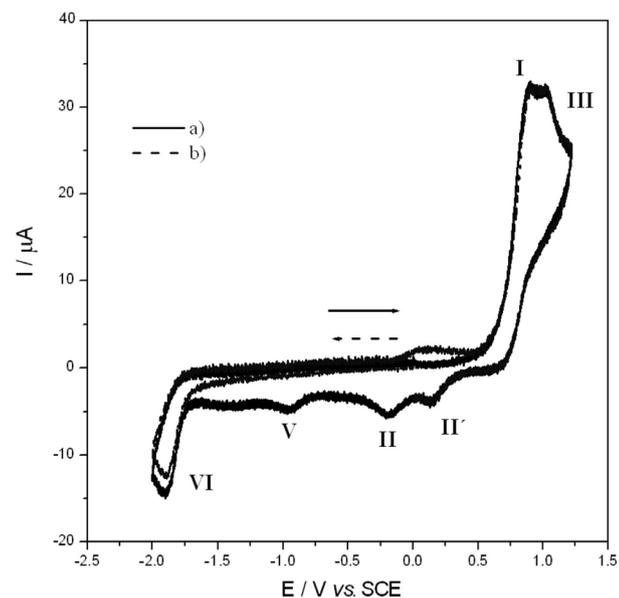


Figure 6. Cyclic voltammetry of FA 2 mmol L^{-1} in DMSO + 0.2 mol L^{-1} $n\text{-Bu}_4\text{NPF}_6$, on glassy carbon electrode ($3 \text{ mm } \phi$). Scan from 1.5 to -2.0 V. a) anodic direction and b) cathodic direction.

reduction wave (IV) in -1.75 V for G and two waves more (V and VI) for FA. It should be mentioned, however that the wave VI is also observed in the cyclic voltammetry of FA in AN.

From these results, it can be inferred that the products obtained during the oxidation processes of both compounds in DMSO, correspond to *o*-quinone or *p*-quinone derivatives, which are substituted with the methoxy group, and emerge from a nucleophilic attack between the water traces present in DMSO and a phenoxy cation generated in a second oxidation step (favored by the electron-releasing power of the methoxy group).

The presence of a methoxy group on the quinone type products is in agreement with the fact that the reduction waves found for G and FA (IV and V, respectively) are more negative than the wave II.³⁶

Electrochemical behaviour of caffeic and ferulic acids amide derivatives in acetonitrile

The electrochemical behaviours of the caffeic (CABZA and CAPEA) and ferulic acid (FABZA and FAPEA) amide derivatives, in AN, are shown in Figures 7 and 8, respectively. In all the cases, it was observed the bielectronic oxidation peak I and its corresponding reduction peak II. Additionally, a small reduction wave (III) to less positive potential values is also observed for all the amides, and an oxidation wave I' is recorded only in the voltammograms of FABZA and FAPEA.

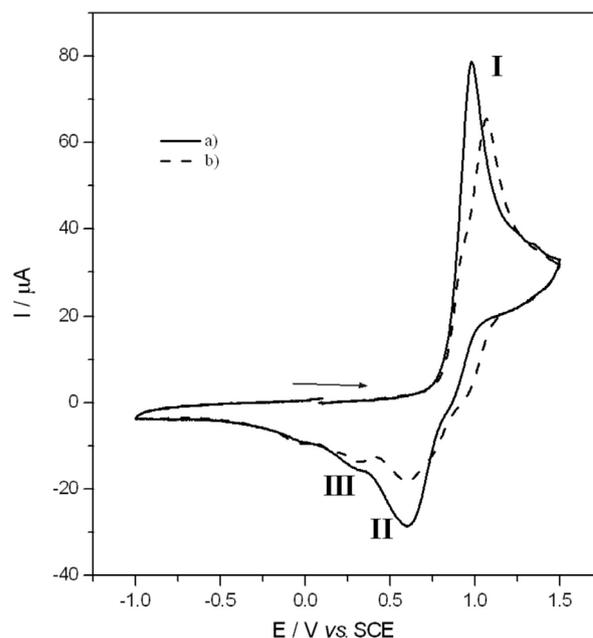


Figure 7. Cyclic voltammetry in acetonitrile + 0.2 mol L^{-1} $n\text{-Bu}_4\text{NPF}_6$, on glassy carbon electrode ($3 \text{ mm } \phi$) at 2 mmol L^{-1} and 0.1 V s^{-1} of a) CABZA and b) CAPEA.

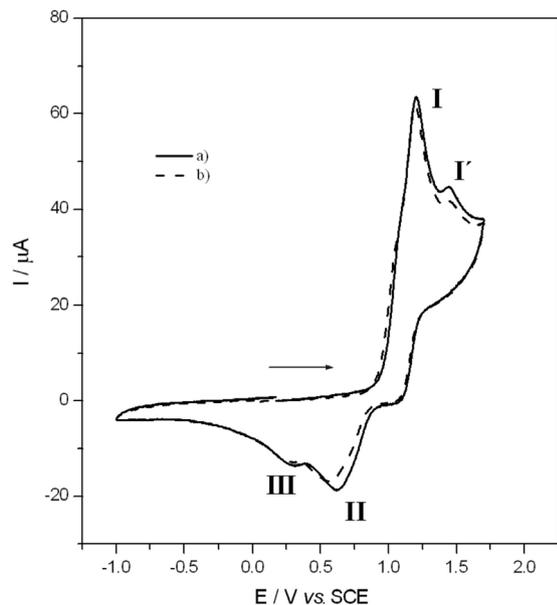


Figure 8. Cyclic voltammetry in acetonitrile + 0.2 mol L⁻¹ *n*-Bu₄NPF₆, on glassy carbon electrode (3 mm ϕ) at 2 mmol L⁻¹ and 0.1 V s⁻¹ of a) FABzA and b) FAPEA.

In the case of CABzA, when the scan rate is increased a new peak (I') appears toward more positive potentials respect to the wave I (Figure 9), which indicates that two oxidative processes must be occurring on this compound. On the other hand, the peak II and the post wave observed in both, CABzA and CAPEA, become a new broad reduction wave at high scan rates.

In addition, the corresponding anodic peak of FABzA and FAPEA decrease steadily with the number of cycles until they disappears completely (see supplementary information).

This behaviour could be related to the derivatization of the electrode surface phenomenon, since the voltammetric behaviour of FABzA is recovered if the electrode surface is polished. From these results, it is evident that the reactivity of the nitrogen atom must be considered also as a part of

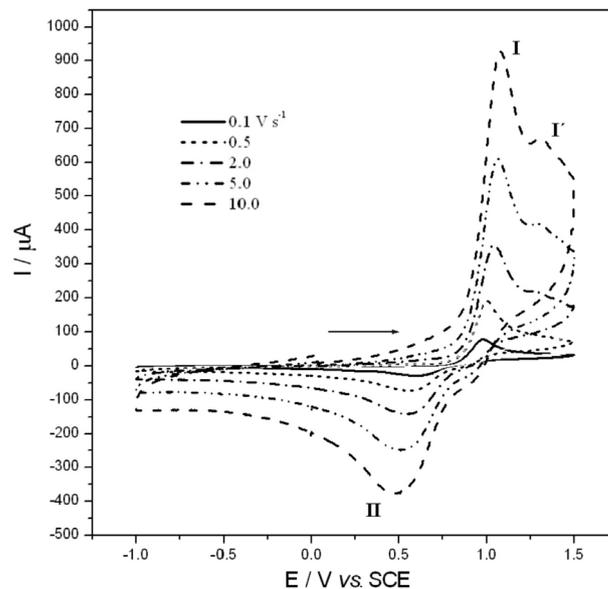
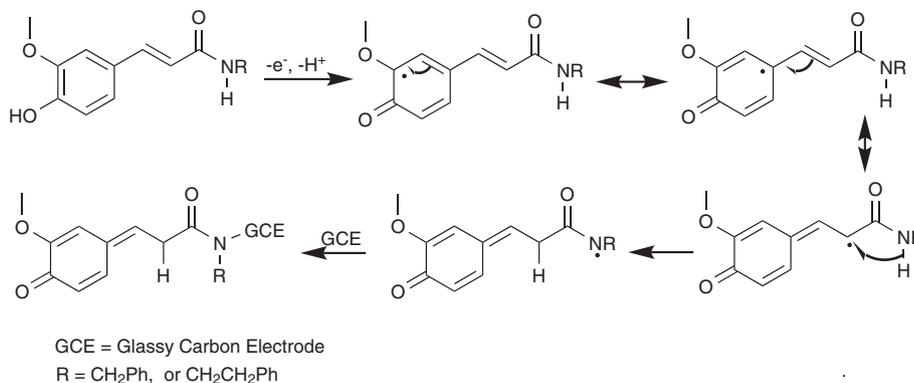


Figure 9. Cyclic voltammetry in DMSO + 0.2 mol L⁻¹ *n*-Bu₄NPF₆, on glassy carbon electrode (3 mm ϕ) of CABzA at 2 mmol L⁻¹ and different scan rate.

the global oxidation mechanism of both compounds^{37,38} as follow (Scheme 4).

Oxidation potentials vs radical scavenging activity

The electrochemical parameters such as the redox potential, the number of electrons, and the electron transfer rate have been used to evaluate the oxidant capabilities of polyphenols.³⁹⁻⁴² The redox potential could be a key factor that governs the antioxidant activity of this type of systems. Therefore, the oxidation potentials of polyphenols are often measured and compared to their antioxidant activities, including the radical scavenging activity. The DPPH assay has been extensively used to evaluate this property of the phenolic compounds, hydroxycinnamic acids and related systems.^{8-13,43,44} In order to see if the electrochemical



Scheme 4.

oxidation capability of all the compounds is linked in some way with their free radical scavenging activity, we carried out DPPH assays of the complete series.

The oxidation potential of the ten compounds reported in this work is shown in Table 3. As it can be observed, the highest values correspond to the guaiacol and ferulic systems, from 1.20 to 1.16 V vs. SCE, whereas the catechol and caffeic acid derivatives exhibit oxidation potentials between 1.11 and 0.98 V vs. SCE.

Table 3. Oxidation potential of C, G, CA and FA derivatives and their corresponding inhibition percentage of 2,2'-diphenyl-1-picrylhydrazyl radical (DPPH)

Compound	E_p / V vs. SCE ^a	Inhibition / (%) (\pm SD) DPPH ^b
C	1.11	31.77 \pm 1.06
CA	1.08	30.56 \pm 1.52
DHCA	1.07	40.52 \pm 0.45
CAPE	1.01	43.48 \pm 0.54
CABzA	0.98	50.38 \pm 0.49
CAPEA	1.07	36.86 \pm 1.49
G	1.18	10.86 \pm 0.50
FA	1.16	19.33 \pm 2.05
FABzA	1.20	8.03 \pm 1.58
FAPEA	1.19	10.81 \pm 0.47

^aOxidation potential of caffeic and ferulic acid derivatives were obtained at 2 mmol L⁻¹ and at 0.1 V s⁻¹ in acetonitrile. ^bThe values represent means \pm S.D. from three experiments and indicate the percentage of inhibition at 17 μ mol L⁻¹ of C, G, CA and FA derivatives.

In agreement with the electrochemical results, the catechol and caffeic systems showed a major inhibition percentage of the 2,2'-diphenyl-1-picrylhydrazyl radical (DPPH) between 30 to 50%, whereas the guaiacol and ferulic acid derivatives showed a lower inhibition percentage between 10 and 20% (Table 3). In addition, the lower oxidation potentials values of the series corresponded to caffeic acid phenethyl ester (CAPE) and one of the caffeic acid amides (CABzA) (1.01 and 0.98 V, respectively). Both compounds exhibited also the higher scavenging radical activity (43.48 and 50.38% of inhibition respectively). A linear regression of the E_p for the ten compounds against their inhibition percentage of a 2,2'-diphenyl-1-picrylhydrazyl radical (DPPH) showed a good agreement between the two sets of data (correlation coefficient of 0.978).

Conclusions

The electrochemical oxidation of the compounds studied in this work, catechol (C), guaiacol (G), caffeic acid (CA), dihydrocaffeic acid (DHCA), ferulic acid (FA), and caffeic acid phenethyl ester (CAPE), in acetonitrile follows an ECE

mechanism. The analyses of the oxidation peak variation with the scan rate of the six compounds, demonstrated that in the case of CA and FA, the initial electron and proton transfer reactions are consecutive and they occur under kinetic control of the deprotonation chemical reaction, since the conjugation of the aromatic ring with the double bond exerts a major stabilization of the radical cation QH₂⁺ generated from the two compounds. In contrast, the oxidations of C, DHCA, G, and CAPE occur under kinetic mixed control. In addition, the substitution of the acidic proton of the caffeic acid by the phenethyl group in the case of CAPE, induces an increase in the rate of the chemical deprotonation.

On the other hand, the formation of hydrogen bond association complexes between DMSO and the catechol moiety of caffeic acid derivatives induce a displacement of the oxidation peaks toward less anodic values respect to the oxidation potentials obtained in AN. In addition, several methoxy quinone type compounds must be generated during the oxidation of G and FA. In the case of anodic oxidation of FA, the analysis of variation of E_p vs. scan rate revealed the establishment of a radical-radical dimerization mechanism in DMSO.

The voltammograms of the four amides analyzed here showed additional waves in AN, respect to the rest of the series. Adsorption processes were even observed during the oxidation of FABzA and FAPEA, which suggests that in addition to the catechol and guaiacol moieties, the nitrogen atom plays also an important role in the global oxidation mechanism of this type of compounds.

Finally, the analysis of the oxidation potential of the ten compounds in AN demonstrated that catechol and caffeic acid derivatives have a major oxidation capacity respect to the guaiacol and ferulic analogues. This is in agreement with the radical scavenging activity evaluated through a DPPH assay. The results showed that CAPE and CABzA exhibited the lower oxidation potentials, as well as the highest reactivity with 2,2'-diphenyl-1-picrylhydrazyl radical. These results corroborate the observation made by other authors, since the reducing power of an antioxidant could be a key factor that governs its antioxidant activity.^{39,42}

Supplementary Information

Supplementary data are available free of charge at <http://jbcs.sbq.org.br>, as PDF file.

Acknowledgments

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Supplementary Information

Electrochemical Oxidation of Caffeic and Ferulic Acid Derivatives in Aprotic Medium

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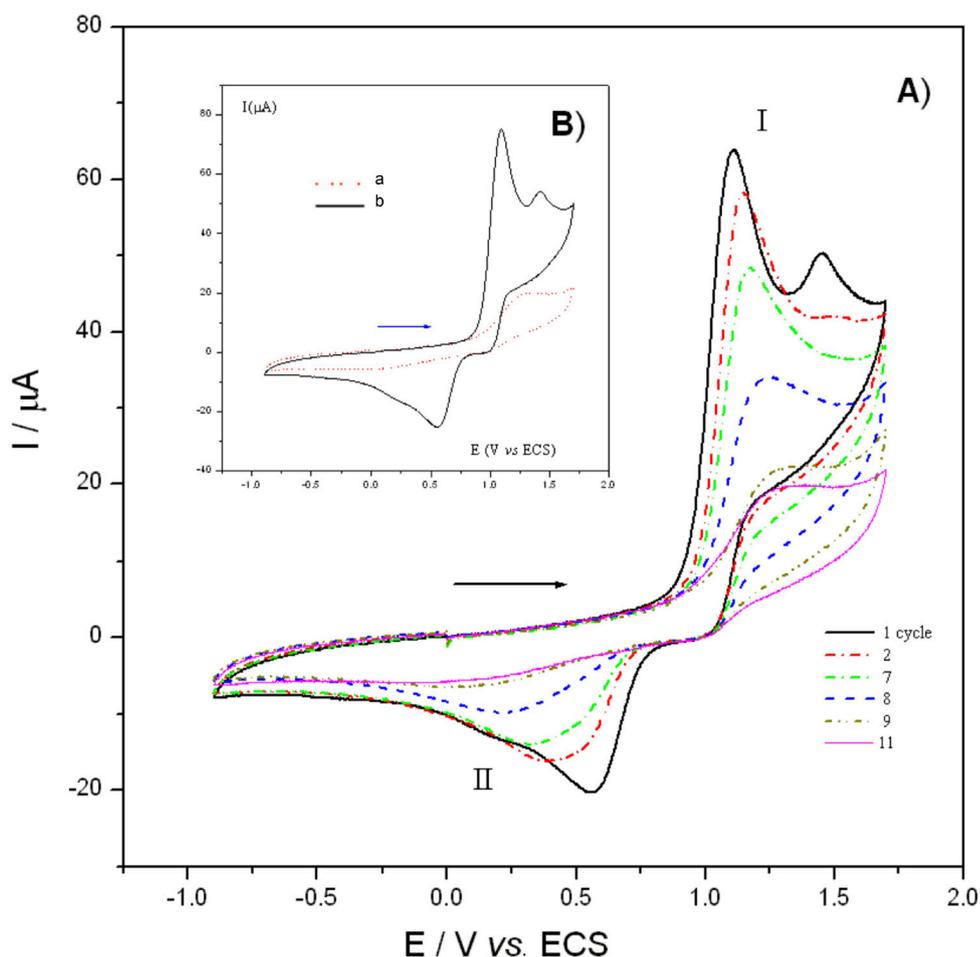


Figure S1. Cyclic voltammeteries of FABZA in acetonitrile + 0.2 mol L⁻¹ n-Bu₄NPF₆, on glassy carbon electrode (3 mm ϕ) at 0.1 V s⁻¹ and 2 mmol L⁻¹. A) During eleven continuous cycles and B) (a) Cyclic voltammetry of FABZA on 11th cycle; (b) after polishing electrode surface.

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