Inexpensive Bismuth-Film Electrode Supported on Pencil-Lead Graphite for Determination of Pb(II) and Cd(II) Ions by Anodic Stripping Voltammetry

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The present work reports the development and application of bismuth-film electrode (BiFE), obtained by in situ method on the pencil-lead graphite surface, for simultaneous Cd(II) and Pb(II) determination at trace levels, as alternative to replace the mercury-film electrodes. Experimental factors, deposition time ($t_d$), deposition potential ($E_d$), and Bi(III) concentration ($C_{Bi}$), were investigated by applying a $2^3$ factorial design using 0.10 mol/L acetate buffer solution (pH 4.5) as supporting electrolyte. The analysis conditions of the differential pulse technique were $t_d = 250$ s, $E_d = -1.40$ V, and $C_{Bi} = 250$ mg L$^{-1}$. The validation of the method employing BiFE was accomplished by determination of merit figures. The detection limits were of 11.0 $\mu$g L$^{-1}$ for Cd(II) and 11.5 $\mu$g L$^{-1}$ for Pb(II), confirming that proposed method is attractive and suitable for heavy metals determination. Additionally, the BiFE developed was successfully applied for the Cd(II) and Pb(II) determination in wastewater sample of battery industry.

1. Introduction

Trace elements, such as Cu, Mo, Mn, and Zn, are considered essential for human health, while others as Hg, Pb, and Cd may accumulate in body tissues causing problems due to their toxicity [1, 2]. This accumulation is due to anthropogenic activities, associated with disposal of solid and liquid waste without proper treatment [3]. Considering the increasing industrial use of these heavy metals and their serious environmental and toxicological impacts, it is necessary to develop new analytical methods to determine them at trace levels [4, 5].

The analytical methods used for heavy metals determination include spectrometry and electroanalytical techniques. Atomic absorption spectrometry (AAS) is the most applied technique for metals determination, since it provides satisfactory sensitivity, high selectivity, and relatively low cost equipment. However, this technique has the disadvantage of not allowing simultaneous determination of chemical elements [6]. The inductively coupled plasma optical emission spectrometry (ICP-OES) and inductively coupled plasma mass spectrometry (ICP-MS) are techniques which allows multielement analysis; however, the high cost of installation and maintenance has restricted their use in research and routine analysis [7–10].

Anodic stripping voltammetry (ASV) is a versatile electroanalytical technique for the trace metals determination in various environmental, clinical and industrial samples [6]. Stripping methods are important in trace analysis because the electrodeposition step concentrates the analyte on the electrode surface, enabling its determination even in extremely low quantities with reasonable accuracy [11, 12].

During the past five decades, mercury-based electrodes were often applied in ASV for presenting an excellent analytical performance [13, 14]. However, due to the high toxicity of mercury and its compounds, the use of this metal in electrodes was restricted. Consequently, an intense research for less toxic and environmentally friendly materials than mercury has been promoted [15–17].

Bismuth-film electrodes were introduced in the last decade to replace the mercury-based electrodes used for the determination of trace heavy metals and organic compounds,
due to very similar electrochemical properties of Bi and Hg, including alloying heavy metals with bismuth as an analogy to amalgamation at mercury [10, 18]. The bismuth-film electrode consists of a bismuth film deposited on a substrate, which can be Au [19], Pt [20], carbon paste [21], glassy carbon [22], carbon fiber [23], pyrolytic graphite [24], and oxides [3, 25, 26]. Moreover, different combinations also are possible as Bi$_2$Te$_3$-graphene oxide hybrid film [27] and bismuth-dispersed xerogel-based composite film [28], besides new methods for Bi$_2$O$_3$-electrode modification as spark discharge [29], proving that there are still issues to be explored in the development of bismuth-based electrodes. Performance of bismuth-film electrodes is similar to traditional mercury electrodes, in addition to the advantage of having negligible toxicity compared to them [30]. The bismuth-film electrode has been applied in voltammetric studies from the ex situ or in situ method, at potential range of -1.2 to 0 V. Positives potentials are not applied, since bismuth is completely removed from the substrate surface by oxidation under these conditions [14].

Carbon electrodes are used extensively as voltammetric sensors in various applications [31]. Mechanical pencils are inexpensive and alternative material to produce carbon electrodes, since they are commercially available with different diameters and hardness. This type of graphite has the advantage of not being fragile as the pyrolytic or paste carbon electrodes but is not hard as the glassy carbon electrode. In addition, graphite has good characteristics such as high electrical conductivity, quick and easy pretreatment, low cost, wide availability, minimum trace metals residue in its composition, and low background current [18].

Bond et al. [12] reported the use of a graphite electrode, which was optimized to anodic stripping analysis using mercury thin-film for Cd and Pb determination and the results were consistent with those obtained by analysis with glassy carbon electrode. The use of bismuth-film electrode supported on pencil graphite (BiFE) has been reported in the literature [3, 15, 32, 33]; however, investigation of manufacturing settings can provide better responses for analytical determination of metal ions.

The present work aimed to prepare a BiFE, investigating the experimental parameters by $2^3$ fractional factorial design, for the simultaneous determination of Pb(II) and Cd(II) ions by differential pulse anodic stripping voltammetry (DPASV). Additionally, the applicability of BiFE in DPASV was evaluated by the analytical figures of merit.

2. Materials and Methods

2.1. Reagents, Chemicals, and Samples. All chemicals used were of analytical grade and milli-Q water was used in the preparation of solutions. The Bi(III), Pb(II), and Cd(II) stock solutions (1000 mg L$^{-1}$) were prepared from Bi(NO$_3$)$_3$·4H$_2$O, Pb(NO$_3$)$_2$ and Cd(NO$_3$)$_2$·4H$_2$O acquired from Sigma-Aldrich. The powder graphite, H$_2$SO$_4$, HNO$_3$, K$_3$Fe(CN)$_6$, and CH$_3$COOH were purchased from Merck. A 0.20 mol L$^{-1}$ H$_2$SO$_4$ solution and a 0.10 mol L$^{-1}$ acetate buffer (pH 4.50) were prepared as substrates electrolytes for the cyclic voltammetry studies of the prepared electrode and for Cd(II) and Pb(II) determination from DPASV, respectively. To evaluate the applicability of the electrode, wastewater samples of battery industry without any previous preparation were analyzed.

2.2. Apparatus. Voltammetric measurements were performed using a potentiostat (Autolab potentiostat/galvanostat GPES IME 663, PGSTAT302n 247V 50/60 Hz). The electrochemical cell coupled to potentiostat was composed of pencil-lead graphite working electrode (GE) or bismuth-film electrode supported on pencil-lead graphite (BiFE), Ag/AgCl as reference electrode and counter electrode of Pt wire.

2.3. Preparation of BiFE. Pencil-lead rods (Pentel Super, 2B, 0.7 mm in diameter) were used to prepare the GE and its assembly scheme is shown in Figure 1. Pencil-lead pieces of 2.5 cm were fitted to the micropipette tips, which were filled with carbon paste prepared by mixing graphite powder and mineral oil (Nujol) to promote the electric contact between the GE and copper wire. This wire was fixed in a glass tube that was connected to the micropipette tip, forming the electrode body. All connections were isolated using paste (cyanoacrylate), Teflon tape, and nonconductive epoxy resin. Due to this assembly, only the lower extremity of the pencil-lead rod (approximately 3.0 mm) is out of the micropipette tip and then available to act as the contact surface for film formation and determination of cations of interest. Since this surface was completely isolated with cyanoacrylate, a polishing step was necessary prior the experiments. Then, the polishing and the surface renewal of the electrodes were done by polishing them on a silk paper, until the base of the electrodes remained with metallic appearance.

The BiFE was prepared from the Bi(III) film formation on the GE surface by the cathode potential application (versus Ag/AgCl sat) by the in situ method. In this, the bismuth deposition occurred simultaneously with the electrochemical deposition of analytes. Moreover, the in situ procedure was chosen because it allows better adhesion of the Bi film to the GE surface, besides the obtaining of higher and better resolved peaks in the voltammograms, along with the higher sensitivity to Cd(II) and Pb(II) when compared to the ex situ method [34, 35].

2.4. Procedure. The cyclic voltammetry analysis was performed as a performance test of the GE prepared as working electrode, using 2.0 mol L$^{-1}$ H$_2$SO$_4$ as the supporting electrolyte. Differential pulse anodic stripping voltammetry (DPASV) was carried out using standard Bi(III) solution (150 to 250 mg L$^{-1}$) in the electrochemical cell to form the Bi film. The deposition time, deposition potential and Bi(III) concentration were investigated using acetate buffer solution (0.1 mol L$^{-1}$) as supporting electrolyte and potential step of 1.95 mV, while the solution was stirred. Before each cycle a 30 s conditioning step at 300 mV (under stirring) was used to remove the bismuth excess and/or the target metals on the electrode surface. The DPASV measurements were performed from equilibrium time of 15 s, time modulation of 0.05 s, pulse interval time of 0.25 s, amplitude modulation of
Table 1: Factors and respective lower and superior levels values.

<table>
<thead>
<tr>
<th>Factors</th>
<th>Lower level (-)</th>
<th>Superior level (+)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bi(III) concentration (mg L(^{-1}))</td>
<td>150</td>
<td>250</td>
</tr>
<tr>
<td>Deposition potential (V)</td>
<td>-1.50</td>
<td>-1.10</td>
</tr>
<tr>
<td>Deposition time (s)</td>
<td>100</td>
<td>250</td>
</tr>
</tbody>
</table>

25 mV, scan rate of 10 mV s\(^{-1}\), and being with different Cd(II) and Pb(II) concentrations.

2.5. Experimental Design. The 2\(^3\) factorial design was applied to investigate the factors: deposition time (t\(_d\)), Bi(III) solution concentration (C\(_{Bi}\)), and deposition potential (E\(_d\)), represented in Table 1. The responses to the proposed design were the current peak areas of the ASV for Cd(II) and Pb(II) at concentrations of 48.3 to 233 \(\mu\)g L\(^{-1}\). The selection of the experimental domain for each factor was determined by previous experiences and literature. Experimental design and data processing were performed using Design Expert 7.1.3 software.

3. Results and Discussion

3.1. Cyclic Voltammetry. The GE electrochemical performance was investigated from cyclic voltammetry measurements of the potassium ferroferricyanide (K\(_4\)[Fe(CN)\(_6\)] / K\(_3\)[Fe(CN)\(_6\)]) redox process. Figure 2 shows the cyclic voltammograms of different K\(_4\)[Fe(CN)\(_6\)] concentrations. It can be seen that cathodic and anodic peaks occur in a short potential interval, indicating the reversibility of redox system.

Current values for anodic and cathodic peaks were proportional to the increase of the K\(_4\)[Fe(CN)\(_6\)] concentration. From the voltammograms of the cyclic voltammetry, it can be seen that the performance of the developed GE is comparable with the glassy carbon [12]. Additionally, a linear relationship between the peak current intensity and K\(_4\)[Fe(CN)\(_6\)] concentration was obtained and showed a linear regression equation of I\(_d\) = 3.16x10\(^{-7}\) + 4.18x10\(^{-9}\) C\(_{K_4[Fe(CN)6]}\) and determination coefficient (R\(^2\)) of 0.9992.

3.2. Anodic Stripping Voltammetry (ASV) with BiFE. During the electrodeposition step in ASV method with BiFE, the codeposition of metallic ions present in the solution along with Bi\(^0\) on the graphite surface is possible [36]. Therefore, the film formation on the GE surface contributes to the improvement of the analytical signal corresponding to such ions, as shown in the Figure 3. It can be seen that the measurement performed with GE showed low current signals at -0.77 and -0.57 V, which are corresponding to Cd(II) and Pb(II), respectively. Applying BiFE, an increase in the analytic signal for the metal species present in solution can be observed, indicating a high sensitivity in the analysis. The highest current peak observed at -0.10 V corresponds to oxidation of bismuth film. According to the results, it can be inferred that other species with characteristic potential values higher than -0.10 V can be analyzed by the method.

3.3. Effect of the Factors Time (t\(_d\)), Deposition Potential (E\(_d\)), and Bi(III) Concentration (C\(_{Bi}\)) on ASV Analyses. In order to investigate the effects of the factors t\(_d\), E\(_d\), and C\(_{Bi}\) on current measured for Cd(II) and Pb(II) determination using BiFE, a 2\(^3\) factorial design was carried out. Table 2 shows the lower and superior levels values of the factors and its responses (current) obtained from the 8 ASV experiments.

Current measurements were made from the voltammograms obtained in potential values of -0.70 and -0.49 V for Cd(II) and Pb(II), respectively. As can be seen in Table 2, the lowest current values were observed for Cd(II), which ranged from 18.4 to 1.79 nA. The current peaks for Pb(II) determination ranged from 3.23 to 29.3 nA. Effects analyses were investigated by Pareto chart (Figure 4), showing the statistic t-test for each effect, where each bar represents the standard effect, i.e., the estimated effect divided by its standard error and the t-critical value [37].

The factors represented by the bars that extend beyond the line are considered significant. According to the results, the t\(_d\) factor was significant for both metals causing a positive effect on the response. The other factors, E\(_d\) and C\(_{Bi}\), as well as the interaction effects between the factors, were not significant. This indicates that metal and film saturation on the electrode surface is not achieved at low time values. Therefore, t\(_d\) is the most important factor for the sensitivity of the technique.
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**Figure 2:** Cyclic voltammograms of potassium ferroferricyanide in 2.0 mol L\(^{-1}\) \(\text{H}_2\text{SO}_4\) using GE (scan rate: 100 mV s\(^{-1}\)).

**Figure 3:** Differential pulse anodic stripping voltammograms obtained for Cd(II) and Pb(II) in 0.10 mol L\(^{-1}\) acetate buffer solution (pH 4.5) as supporting electrolyte; (a) 250 mg L\(^{-1}\) of Bi(III) dissolved in supporting electrolyte, (b) without Bi(III), determined with GE containing 142 \(\mu\)g L\(^{-1}\) of Cd(II) and Pb(II) (\(E_{dp} = -1.40\) V; \(t_{dp} = 250\) s; potential step =1.95 mV; amplitude = 25 mV).

**Figure 4:** Pareto chart to evaluate the peaks area for Cd(II) (a) and Pb(II) (b), where \(C_{\text{Bi}}\) (A) and \(t_{d}\) (C).

using BiFE, which is in agreement with other works that reported the development of bismuth electrodes [18, 37, 38]. Thus, the optimized conditions selected for subsequent studies were those frequently reported in the literature: 250 s deposition time, Bi(III) concentration of 250 mg L\(^{-1}\), and -1.40 V deposition potential [3, 26, 38–40].

3.4. **Analytical Method Development.** Analytical figures of merit were determined using the selected conditions \((t_{dp}, E_{dp}, \text{ and } C_{\text{Bi}})\) to evaluate the applicability of BiFE (film obtained by the in situ method) on the Cd(II) and Pb(II) determination. Figure 5 shows differential pulse anodic stripping voltammograms for simultaneous determination of these ions using the
Table 2: Experiments of $2^3$ factorial design and response values.

<table>
<thead>
<tr>
<th>Order</th>
<th>$c_B$ (mg L$^{-1}$)</th>
<th>$E_d$ (V)</th>
<th>$t_d$ (s)</th>
<th>$I_{Cd(II)}$ (nA)</th>
<th>$I_{Pb(II)}$ (nA)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>150</td>
<td>-1.50</td>
<td>100</td>
<td>5.31</td>
<td>7.92</td>
</tr>
<tr>
<td>2</td>
<td>150</td>
<td>-1.50</td>
<td>250</td>
<td>16.8</td>
<td>29.3</td>
</tr>
<tr>
<td>3</td>
<td>250</td>
<td>-1.50</td>
<td>100</td>
<td>4.71</td>
<td>9.26</td>
</tr>
<tr>
<td>4</td>
<td>150</td>
<td>-1.10</td>
<td>250</td>
<td>18.4</td>
<td>29.3</td>
</tr>
<tr>
<td>5</td>
<td>250</td>
<td>-1.50</td>
<td>250</td>
<td>7.02</td>
<td>14.9</td>
</tr>
<tr>
<td>6</td>
<td>250</td>
<td>-1.10</td>
<td>250</td>
<td>6.97</td>
<td>14.9</td>
</tr>
<tr>
<td>7</td>
<td>150</td>
<td>-1.10</td>
<td>100</td>
<td>2.75</td>
<td>3.23</td>
</tr>
<tr>
<td>8</td>
<td>250</td>
<td>-1.10</td>
<td>100</td>
<td>1.79</td>
<td>6.37</td>
</tr>
</tbody>
</table>

![Figure 5: Differential pulse anodic stripping voltammograms for increasing concentrations ranging from 48.3 to 233 $\mu$gL$^{-1}$ of Cd(II) and Pb(II) in 0.10 mol L$^{-1}$ acetate buffer solution (pH 4.5) as supporting electrolyte using BiFE.](image)

According to the figure, the two peaks corresponding to metal ions are well resolved and increase with increasing concentrations.

From the calibration curve of each chemical species, it can be demonstrated linearly between the current values and concentrations within the range of 48.3 to 233 $\mu$gL$^{-1}$ by the linear regression equation and determination coefficient of experimental data: $I_d = 2.44 \times 10^{-3} C_{Cd} + 4.29 \times 10^{-3}$ and $R^2 = 0.9964$ for Cd(II) and $I_d = 8.35 \times 10^{-4} C_{Pb} - 5.02 \times 10^{-3}$ and $R^2 = 0.9954$ for Pb(II).

3.4.1. Detection and Quantification Limits. To evaluate the limit of detection (LOD) and limit of quantification (LOQ), five different blank solutions were used, which were analyzed in triplicate and the standard deviation ($S_p$) of the means was calculated. The measurements were performed in 5.0 mL of blank solutions with 5.0 mL of acetate buffer solution, under the optimized conditions. The LOD and LOQ values were determined from the $3S_p/m$ and $10S_p/m$ relations, respectively, where $m$ is the slope of the calibration curve of each chemical species. For Cd(II), the calculated values of LOD and LOQ were of 11.0 $\mu$gL$^{-1}$ and 36.8 $\mu$gL$^{-1}$, respectively. For Pb(II), the LOD was 11.5 $\mu$gL$^{-1}$ and LOQ was 38.2 $\mu$gL$^{-1}$. The LOD values obtained at levels of $\mu$gL$^{-1}$ are comparable with other studies involving modified electrodes. Serrano et al. [41] prepared modified graphite-epoxy composite electrodes which showed LOD of 2.40 and 4.70 $\mu$gL$^{-1}$ for Cd(II) and of 1.50 and 3.30 $\mu$gL$^{-1}$ for Pb(II), simultaneously determined. Kadara and Toffill [42] found LOD of 8.00 and 16.0 $\mu$gL$^{-1}$ for Pb(II) and Cd(II), respectively, using a screen-printed Bi$_2$O$_3$-modified electrode. Løche et al. [43] employed screen-printed electrodes modified with five bismuth precursor compounds in Pb(II) and Cd(II) determination, obtaining LOD values of 0.90-1.40 and 1.10-3.20 $\mu$gL$^{-1}$, respectively. Therefore, low LOD values could be achieved by increasing the deposition time during the Bi film formation in the proposed method, once the deposition time showed a positive effect on the response by increasing the current intensity, as demonstrated in Table 2. Some studies in the literature corroborate this fact, as an example: Zhang et al. [35] employed a deposition time of 300 s during the in situ Bi deposition on working electrode and LOD of 0.02 $\mu$gL$^{-1}$ for Pb(II) and 0.01 $\mu$gL$^{-1}$ for Cd(II) were obtained; Demetriades et al. [18] prepared a bismuth-film electrode supported on pencil graphite with Bi deposition time of 600 s and LOD of 0.30 $\mu$gL$^{-1}$ for Cd(II) and 0.40 $\mu$gL$^{-1}$ for Pb(II) were determined.
3.4.2. Repeatability. The repeatability (intraday) was evaluated from five measurements (in triplicate) of solutions containing 150 \( \mu \text{g L}^{-1} \) Cd(II) or 150 \( \mu \text{g L}^{-1} \) Pb(II) with 10.0 mL of acetate buffer solution, with just one BiFE. The mean of triplicates and the corresponding relative standard deviation (RSD) were calculated to obtain the analytical parameter. The repeatability values obtained, expressed as % RSD, were of 9.07% for Cd(II) and 11.6% for Pb(II), which are acceptable with respect to trace level determination of heavy metals.

3.4.3. Reproducibility. The reproducibility was determined from measurements (in triplicate) of five different BiFEs for solutions of 150 \( \mu \text{g L}^{-1} \) Cd(II) or 150 \( \mu \text{g L}^{-1} \) Pb(II) with 10.0 mL of acetate buffer solution. This analytical parameter was calculated from RSD obtained from the mean value of measurements. The reproducibility values (expressed as the % RSD) achieved in this work were of 9.72% for Cd(II) and 7.87% for Pb(II) determination.

3.4.4. Standard Addition Method. The BiFE was applied in the determination of Cd(II) and Pb(II) in a wastewater sample from a battery industry. The sample was analyzed without any previous preparation and ions determination was performed by DPASV under optimized conditions. Figure 6 shows the voltammograms obtained by the standard addition method for Cd(II) and Pb(II) determination in the sample. The curve was constructed using concentrations of Cd(II) and Pb(II) ranging from 48.3 to 233 \( \mu \text{g L}^{-1} \).

The calibration curves for the standard addition method of each chemical species are shown in Figure 7. According to the figure, a good linearity between current signals and analytes concentrations can be observed. The linear regression equation and determination coefficient obtained were:

\[
I_d = 1.36 \times 10^{-3} \ C_{\text{Cd}} - 1.05 \times 10^{-2}
\]

and

\[
I_d = 1.35 \times 10^{-3} \ C_{\text{Pb}} + 1.79 \times 10^{-2}
\]

and \( R^2 = 0.9873 \) for Cd(II) and \( R^2 = 0.9706 \) for Pb(II). The Cd(II) was not detected in the sample, while the Pb(II) determined concentration was of 48.8 \( \mu \text{g L}^{-1} \). To
Table 3: Recovery of Cd(II) and Pb(II) in battery industry wastewater sample fortified in two concentration levels: (I) 190 μg L$^{-1}$ and (II) 235 μg L$^{-1}$.

<table>
<thead>
<tr>
<th></th>
<th>Cd(II) (μg L$^{-1}$)</th>
<th>Recovery (%)</th>
<th>Pb(II) (μg L$^{-1}$)</th>
<th>Recovery (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blank</td>
<td>(I)</td>
<td>(II)</td>
<td>(I)</td>
<td>(II)</td>
</tr>
<tr>
<td>ND</td>
<td>205 ± 7.4</td>
<td>228 ± 8.5</td>
<td>108 ± 3.9</td>
<td>96.9 ± 3.6</td>
</tr>
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<td></td>
<td></td>
<td></td>
<td>15.1</td>
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</tr>
</tbody>
</table>

3.4.5. Recovery Test. The recovery method was used in the battery industry wastewater sample, which was fortified with 190 μg L$^{-1}$ Cd(II) and 235 μg L$^{-1}$ Pb(II) solutions. Table 3 shows the results of the recovery study in wastewater sample using BiFE and optimized experimental conditions. From the table, it can be noticed that the recovery for both metals was close to 100%, ranging from 96.9 to 108% for the addition levels studied. This indicates adequate precision and accuracy of the method employing BiFE for determination of heavy metals at trace levels.

4. Conclusions

The BiFE developed by in situ method for Pb(II) and Cd(II) simultaneous determination at trace levels showed an efficient analytical performance. Among the parameters investigated for differential pulse anodic stripping voltammetry, the deposition time was the most important factor for the technique sensitivity. From the optimized conditions, the BiFE exhibited a linear response in the range between 48.3 and 233 μg L$^{-1}$ with a detection limit of 11.0 μg L$^{-1}$ for Cd(II) and 11.5 μg L$^{-1}$ for Pb(II). Repeatability and reproducibility showed acceptable values for trace level determination of heavy metals. A wastewater sample of battery industry was used in the recovery method to evaluate the analytical application of BiFE on real samples. The recovery for both metals was close to 100%, demonstrating the precision and accuracy of the electrode developed. In this way, the results imply that BiFE can be useful in trace level determination of Cd(II) and Pb(II) in heavily polluted samples (industrial waste waters, aqueous sources from mining areas, etc.), thanks to its analytical performance, low cost, wide availability, easy preparation, and less time consuming.

Data Availability

The authors confirm that all data created during this research, which are required to prove the results, are included within the article. In addition, questions about the data used to support the findings of this study are available from the corresponding author (vcalmeida@uem.br) upon request.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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