Application of CPE-FAAS Methodology for the Analysis of Trace Heavy Metals in Real Samples using Phenanthraquinone Monophenyl Thiosemicarbazone and Triton X-114

Magda Akl A*, Magdy Bekheit M and Ibraheim Helmy
Faculty of Science, Mansoura University, Mansoura, Egypt

Abstract

In the present study, a cloud point extraction method was used for the preconcentration and extraction of cadmium (II) and zinc (II) ions in different environmental samples. The zinc and cadmium ions formed hydrophobic complexes with phenanthraquinone monophenyl thiosemicarbazone (PPT). These complexes were extracted using Triton X-114 nonionic surfactant. The surfactant-rich phase was diluted with acidified methanol. Then, the concentrations of the metal ions were determined by FAAS. The experimental factors controlling the process of separation were investigated e.g., pH, complexing agent concentration, surfactant's concentration, temperature, and incubation time. The present CPE-FAAS procedure has been used to preconcentrate and determine Cd(II) and Zn(II) metal ions in natural water samples, drug samples and certified reference materials. The LODs for cadmium(II) and zinc(II) ions were 0.38 and 1.85 μg/L, respectively with a preconcentration factor of 100. The recovery % of the extracted Cd(II) and Zn(II), is greater than 90% and the relative standard deviation (RSD, %) is less than 5%.

Keywords: CPE; Zinc(II); Cadmium(II); Surfactants; FAAS

Introduction

Heavy metals in the environment are really increasing due to human activity, particularly mining process and heavy industry in the developing world. This has caused great accumulation of high concentrations of toxic heavy metals in natural waters [1].

Monitoring the presence of toxic trace metals in different matrices is of great importance in order to evaluate the exposure to these heavy metals in the environment. In this sense, cadmium is considered as one of the most toxic elements; it accumulates in humans body mostly in the kidneys and liver and is classified as a rampant toxic element with biochemical functions in all living organisms [3]. Deficiency of zinc can lead to many disorders such as growth retardation, diarrhea, inefficient immunological defense, eye and skin lesions, delaying of wound healing and other skin diseases [4]. Excess amounts of zinc are harmful and can be toxic when exposures exceed the physiological needs.

Monitoring trace element levels in environmental samples might be considered a difficult analytical task, mostly because of the complexity of the matrix and the very low concentrations of these elements, which requires a prior preconcentration step followed by sensitive instrumental techniques [5]. Many separation/preconcentration techniques have been developed for trace metal analysis such as solid phase extraction [6], liquid-liquid extraction [7], co-precipitation [8], column extraction [9], ion-selective electrode [10] and cloud point extraction (CPE) [11-16].

Cloud point extraction (CPE) has attracted a great attention because it complies with the “Green Chemistry” principle [17]. CPE is simple, highly efficient, cheap, rapid and of lower toxicity than those procedures using organic solvents.

Cloud point extraction (CPE) is based on the phase behavior of non-ionic surfactants in aqueous solutions. Non-ionic surfactants undergo phase separation upon raising the temperature or the addition of a salting-out agent [18,19]. This procedure has been successfully employed to extract and preconcentrate different trace metals from different samples [20-23].

In recent years, considerable attention has been paid to the use of chelating agents, containing sulphur and nitrogen in analytical chemistry, in separation, purification and estimation of metal ions [24]. The extensive application and rapid growth in the popularity of sulphur ligands is because of their outstanding property as potential donors to form stable and characterized complexes. In addition, the presence of nitrogen along with sulphur tends to reduce the solubility of the complexes, making the isolation of these complexes easier.

Among various organic chelating reagents containing S and N, thiosemicarbazones and their aromatic derivatives occupy a unique place. Thiosemicarbazones are a group of organic compounds prepared by condensing thiosemicarbazide with carbonyl compounds in the presence of few drops of glacial acetic acid. These organic reagents function as good complexing agents and form complexes with various metal ions through thionate Sulphur atom and hydrazine nitrogen atom binding sites.

Phenanthraquinone monononyphenyl thiosemicarbazone (PPT) reacts with different metal ions such as Ni(II), Zn(II), Cu(II), Cd(II) and Pb(II) to form hydrophobic colored complexes. These metal ions were further determined in media of diverse origin by FAAS and/or ultraviolet-visible (UV-Vis) spectrophotometry [25-28].

*Corresponding author: Magda Akl A, Faculty of Science, Mansoura University, Mansoura, Egypt, E-mail: magdaakl@yahoo.com

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In the present work, CPE method has been used for the preconcentration of zinc and cadmium after the formation of hydrophobic complexes with phenanthraquinone monophenyl thiosemicarbazone (PPT). These complexes were simply extracted using Triton-X114 nonionic surfactant followed by FAAS determination. The analytical conditions for the preconcentration, extraction and FAAS determination of the analytes were investigated. The proposed combined CPE-FAAS methodology was successfully used for the analysis of zinc(II) and cadmium(II) in water, drug and certified samples.

Experimental

Instruments

A Perkin-Elmer 2380 air-acetylene atomic absorption spectrometer was used for the determination of Cd(II) and Zn(II) concentrations. To measure the pH of sample solutions, a digital pH meter was used.

Chemicals

All the chemicals used in the present study are of analytical-reagent grade. Double distilled water (DDW) was used through. Zinc stock solution (1000 mg L⁻¹) was prepared by dissolving 2.089 g ZnCl₂ in 100 ml of water with the addition of 2 ml of concentrated HCl and dilution to 1 L with bidistilled water. Cd stock solution (1000 mg L⁻¹) was prepared by 2.03 g of CdCl₂.5 H₂O in 1 L DDW. Triton X-114 was obtained from Sigma. To prepare a 1% (v/v) Triton X-114 stock solution, 1 mL of Triton X-114 was dispersed in 5 mL ethyl alcohol and completed with 100 ml DDW.

Phenanthraquinone monophenyl thiosemicarbazone, PPT was synthesized as previously reported [26]. A 10⁻³ mol. L⁻¹ solution of PPT was prepared by dissolving 0.357 gm of phenanthraquinone monophenyl thiosemicarbazone (PPT) in 100 ml of acetone.

Analytical procedure

An aliquot of 20 ml of aqueous solution that contains 3 μg mL⁻¹ of metal ions, 2 × 10⁻⁴ mol. L⁻¹ of PPT and 0.05% (v/v) of Triton X-114 at pH 6 was prepared (Scheme 1). This mixture was shaken for 1 minute and left to stand for 20 minutes in water bath at 60°C. Centrifugation at 3500 rpm for 10 minutes was performed to achieve separation of the two phases. Upon cooling in an ice bath for 15 minutes, the surfactant-rich phase became viscous and the mother aqueous phase was smoothly separated. The micellar phase is dissolved in 0.2 mL of acidified methanol and aspirated directly to the flame to determine the concentration of the investigated metal ions.

Calibration curves were constructed in the concentration ranges 0.40-2.0 mg/l and the concentrations of the investigated metal ions were determined by FAAS at specified wave lengths for each analyte.

For determination of zinc(II) in pharmaceutical samples, two commercial drug samples were selected viz., Vitamax plus and Totavit tablets. The samples were brought into solution using the procedure previously described in the literature [28].

Results and Discussion

Cloud point extraction (CPE)

Effect of pH: The pH is a very important parameter for both the coacervation of the micelles and the complexation of the ligand with the metal ions. The influence of pH on analytes extraction was investigated by performing a number of CPE experiments and varying the pH of sample solution over a wide range (2-9) using 3 μg mL⁻¹ of Zn(II) and 3 μg mL⁻¹ of Cd(II) in the presence of 2 × 10⁻⁴ mol L⁻¹ PPT and 0.05% (v/v) Triton X-114. As it can be seen from Figure 1, the maximum recovery was achieved at pH 6.

Effect of PPT concentration: The CPE efficiency depends on the hydrophobicity of the ligand and the complex formed. So, in order to evaluate the role of PPT in CPE of Zn(II) and Cd(II), similar CPE experiments were performed by adding different concentrations of PPT to a suitable concentrated, 3 μg mL⁻¹ of Zn(II), 3 μg mL⁻¹ of Cd(II) and 0.05% (v/v) Triton X-114 at pH 6. Figure 2 shows that, at low PPT concentration, the recovery of the analytes is low because the amount of ligand required for complexation of all the amount of the analytes is insufficient. Then, the recovery significantly increases by increasing PPT concentration till maximum recovery was attained at 1.5 × 10⁻⁴ mol. L⁻¹ PPT. The recovery of the analytes remains constant upon the use of higher concentrations of the chelating agent. So, 2 × 10⁻³ molL⁻¹ PPT was selected for the subsequent work.

Effect of Triton X-114 concentration: The effect of Triton X-114 concentration on the extraction efficiency of 3 μg mL⁻¹ of Zn(II) and 3 μg mL⁻¹ of Cd(II) in the presence of 2 × 10⁻⁴ mol L⁻¹ PPT at pH 6 was studied in the concentration range of 0.02-0.25% (v/v). In Figure 3, the results showed that, at low surfactant concentration the recovery of the analytes is poor because there are few surfactant molecules to entrap the ligand-metal complexes quantitatively [29]. Then, the recovery sharply increases by increasing the concentration of Triton X-114. The highest recoveries were obtained with 0.05% (v/v) Triton X-114. By increasing the surfactant concentration above 0.1% the recovery starts to decrease gradually.

Effect of temperature and incubation time: Like the other parameters, temperature and duration of the CPE procedure seem to play significant roles. Raising the incubation temperature above the cloud point temperature is an important factor for obtaining satisfactory extraction and enhancing CPE efficiency in a reduced incubation time. A series of experiments was performed to investigate the effect of temperature on the CPE method under the previous optimum condition of pH, PPT concentration and Triton X-114 concentration over a wide range of temperatures (20-80°C) through 20 minutes’ incubation time. The data in Figure 4 shows that maximum recovery was obtained at range (50-80°C). The enhanced recovery at high temperature is related to the dehydration between hydrogen bonds that occur by increasing temperature and decrease amount of water in surfactant rich phase hence the volume of the phase decrease leading to enhancing the recovery. The effect of the incubation time, on the other hand, was investigated within the ranges 5-60 min. The results, Figure 5, illustrate that an equilibration time of 20 min was quite adequate to achieve quantitative extraction of cadmium(II) and zinc(II) ions [30].

Scheme 1: Phenanthraquinone monophenyl thiosemicarbazone (PPT).
Effect of centrifugation time and rate: For the best efficiency of the method, it is required to preconcentrate trace amount of metal ions with high sensitivity and in a short time. To test the influence of centrifugation time on the method sensitivity, CPE has been carried out for a series of experiments, in which an aliquot of 10 ml of aqueous solution containing 3 μg mL⁻¹ of Zn(II) and 3 μg mL⁻¹ of Cd(II) in the presence of 2 × 10⁻⁴ mol L⁻¹ PPT and 0.05% (v/v) of Triton X-114 at pH 6 has been incubated at 60°C for 20 minutes and centrifuged at various rates (1500-4500 rpm) and for different time periods (5-20 min). The best conditions for sample centrifugation were 10 min at 3500 rpm.

Effect of foreign ions: Because of the high selectivity attributed to flame atomic absorption spectrometry, the only source of low sample recovery must be the preconcentration step. This problem may be attributed to the fact that cations may react with ligand and anions may form stable complex with metal ions and resulting in a decrease in extraction efficiency (Table 1). Therefore studying the effect of foreign ions was carried out using 3 μg mL⁻¹ of Zn(II) and 3 μg mL⁻¹ of Cd(II) in the presence of an excess amount of PPT (5 × 10⁻⁴ mol L⁻¹). The effect of almost foreign ions, even those which formed colored complexes with PPT such as Cu²⁺, Ni²⁺ and Pb²⁺, interference could be eliminated.

Analytical characteristics

To evaluate the reproducibility of the method, 20 mL of model solution containing metal ions (n=10) was used. The relative standard deviation (RSD, %) as precision of the method (RSD, %) was 1.86 and 3.06%, for Cd(II) and Zn(II), respectively (Table 2). The LODs based on three times the SD of the blank for cadmium(II) and zinc(II) ions were 0.38
Table 1: Effect of interfering ion on the recovery of Zn(II) and Cd(II) in the presence of 2 × 10⁻⁴ PPT and 0.05% (v/v) Triton X-114 (n=5).

| Foreign ion | Concentration (mg, L⁻¹) | Recovery % |
|-------------|-------------------------|------------|
|             |                         | Zn(II)     | Cd(II)    |
| Hg²⁺        | 20                      | 100        | 96.9      |
| Ni²⁺        | 10                      | 100        | 97.7      |
| Co³⁺        | 20                      | 97.9       | 98.4      |
| Cr³⁺        | 5                       | 98.9       | 99.3      |
| Fe³⁺        | 10                      | 99.8       | 98.5      |
| Cu²⁺        | 10                      | 98.2       | 97.6      |
| Pb²⁺        | 10                      | 99.7       | 99.2      |
| Ba²⁺        | 20                      | 99.8       | 98.4      |
| Mn²⁺        | 20                      | 100        | 96.8      |
| Br⁻         | 20                      | 100        | 97.3      |
| Al³⁺        | 20                      | 100        | 98.1      |
| Na⁺         | 230                     | 98.8       | 97.9      |
| SO₄²⁻       | 480                     | 99.5       | 99.8      |
| Cl⁻         | 177.5                   | 97.9       | 99.6      |
| NO₃⁻        | 230                     | 98.7       | 97.8      |
| CO₃²⁻       | 310                     | 98.1       | 100       |
| C₂O₄²⁻      | 300                     | 97.2       | 96.8      |
| CH₃COO⁻     | 440                     | 97.9       | 100       |
| µS          | 295                     | 98.8       | 98.5      |

Linear range, mg/L

Analyte Correlation coefficient Linear range, mg/L Regression equation* RSD, %
Cd 0.9997 0.40–1.50 $A=0.0471C+0.001C$ 1.86
Zn 0.9999 0.50–2.00 $A=0.0367C–0.00049$ 3.06

* A=Absorbance, C=concentration

Table 2: Analytical characteristics of the calibration curves of the analytes.

To verify the accuracy and applicability of proposed CPE procedure, two reference standard materials (lead-zinc sulphide ore OCRO (COD 161-96) and stream sediment SARM 52) were analyzed. The results are given in Table 3. There is no significant difference between the results obtained by the proposed method and the certified results. The relative standard deviation (as a precision) is less than 5%.

Statistical analysis of the results in Table 3 indicate that the preconcentrated samples are not subject to any systematic error i.e., accurate.

Applications

Water analysis: Various water samples from different origins (Tap water, Nile river and sea water samples) were spiked with different amounts of cadmium and zinc and CPE procedures were employed for determination of the recovery of the analytes in these water samples, Table 4. The results of the present CPE method for Cd(II) and Zn(II) are in good comparison with those gained upon using the well-known APDC/MIBK solvent extraction method (SE), Table 5.

Analysis of pharmaceutical samples: Cloud point extraction followed by FAAS determination was applied to determine zinc in some commercial zinc containing pharmaceutical samples. The experimental results agreed well with the given reported values, Table 6.

Application to synthetic mixtures: An aliquot of 10 ml of aqueous sample solution containing different compositions of foreign metal ions (Co(II), Ni(II), Hg(II) and Bi(III), Cr(III) and Pb(II), different amounts of the analytes, 2 × 10⁻⁴ mol L⁻¹ of PPT and 0.05% (v/v) of Triton X-114 at pH 6. The CPE procedures were performed under the previously mentioned optimum conditions and the recovery of the analytes were determined. The results obtained in Table 7 shows high extraction recovery of the analytes from samples containing different synthetic mixtures.

Table 3: Statistical evaluation for analysis of some pharmaceutical vitamin samples after CPE (n=5). Comparison of experimental mean (X̄) with true value (µ) by $t$ test.

| Water sample | Cd(II) added (µg/ml) | Zn(II) added (µg/ml) | Recovery % |
|--------------|----------------------|----------------------|------------|
| Tap water    | 5.00                 | 5.00                 | 98.9       |
| (Our laboratory) | 10.00             | 10.00               | 98.8       |
| Nile water   | 5.00                 | 5.00                 | 99.3       |
| (Mansoura city) | 10.00             | 10.00               | 99.7       |
| Sea water    | 5.00                 | 5.00                 | 98.7       |
| (Ras Elbar city) | 10.00             | 10.00               | 98.6       |

Table 4: Recovery of Cd(II) and Zn(II) in different water samples (n=5).

Table 5: Determination of Cd(II) and Zn(II) in µg/ml in natural water samples by the present cloud point extraction (CPE) and the standard APDC/MIBK solvent extraction methods (SE) (n=6 for both methods).

| Sample                  | Cloud point extraction (PE) | Solvent extraction (SE) |
|-------------------------|-----------------------------|-------------------------|
|                         | Cd(II), µg/l                | Zn(II), µg/l             | RSDs,%       |
| Tap water (Our Lab Mansoura City) | 0.30 185                   | 1.0-5.0                 | 0.32 186.0   | 2.0-3.0 |
| Waste water (Meat Anter, Talkha) | 0.6 80                     | 2.0-4.0                 | 0.62 82.0    | 1.2-4.0 |
| Nile River water (Sherbin City) | 0.13 26                    | 1.5-4.8                 | 0.14 25.5    | 1.5-4.5 |
| Lake water (Manzalah) | 0.15 158                   | 1.0-5.0                 | 0.16 160.0   | 2.0-4.5 |
| Sea water (Ras Elbar city) | 0.12 30.0                  | 2.0-4.0                 | 0.11 31.5    | 1.0-5.0 |

Table 6: Statistical evaluation for analysis of some pharmaceutical vitamin samples after CPE (n=5). Comparison of experimental mean (X̄) with true value (µ) by $t$ test.

| Sample                  | µg/l               | RSD, % |
|-------------------------|--------------------|--------|
| No. 1*                  | 14.95              | 15.0   | 0.05   | 1.5 | 4.2 |
| No. 2*                  | 15.02              | 15.0   | 0.02   | 1.95 | 2.8 |

*Vitmax plus (Gaiox Welco Egypt) capsule; *Totalit (Egypt) tablet. (X̄): experimental value, (µ) true value. For P=0.05 and n=5 (4 degree of freedom)=2.78, from Ref. [29], RSD; %. Relative standard deviation.
Conclusion

In the present study, a simple, low cost and reliable combined cloud point extraction-FAAS methodology using the Triton-X114 nonionic surfactant has been used successfully for the determination of cadmium and zinc in water, vitamin and certified samples. The analytical characteristics of the proposed procedure (LOD, Recovery, % and RSD, %) are obtained. The limit of detection of cadmium and zinc gained is in good comparison or superior to the cloud point procedures reported in literature. Two certified reference materials were used to verify the accuracy of the proposed method.

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