

CHLORMEQUAT SELECTIVE ELECTRODES: CONSTRUCTION, EVALUATION AND APPLICATION AT FIA SYSTEMS

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ABSTRACT

A flow injection analysis (FIA) system having a chlormequat selective electrode is proposed. Several electrodes with poly(vinyl chloride) based membranes were constructed for this purpose. Comparative characterization suggested the use of membrane with chlormequat tetraphenylborate and dibutylphthalate. On a single-line FIA set-up, operating with $1 \times 10^{-2} \text{ mol L}^{-1}$ ionic strength and 6.3 pH, calibration curves presented slopes of $53.6 \pm 0.4 \text{ mV decade}^{-1}$ within 5.0×10^{-6} and $1.0 \times 10^{-3} \text{ mol L}^{-1}$, and squared correlation coefficients > 0.9953 . The detection limit was $2.2 \times 10^{-6} \text{ mol L}^{-1}$ and the repeatability equal to $\pm 0.68 \text{ mV}$ (0.7%). A dual-channel FIA manifold was therefore constructed, enabling automatic attainment of previous ionic strength and pH conditions and thus eliminating sample preparation steps. Slopes of $45.5 \pm 0.2 \text{ mV decade}^{-1}$ along a concentration range of 8.0×10^{-6} to $1.0 \times 10^{-3} \text{ mol L}^{-1}$ with a repeatability $\pm 0.4 \text{ mV}$ (0.69%) were obtained. Analyses of real samples were performed, and recovery gave results ranging from 96.6 to 101.1%.

Keywords: Chlormequat; Growth regulator; Pesticides; Potentiometry; Ion-selective electrode; FIA

INTRODUCTION

Plant growth retardants are applied in agronomic and horticultural crops to reduce unwanted longitudinal shoot growth without lowering plant productivity. Most growth retardants act by inhibiting gibberellin biosynthesis. To date, four different types of such inhibitors are known, among which are the onium compounds such as chlormequat chloride (CMQ-Cl), the 2-chlorethyltrimethylammonium [1].

Chlormequat (CMQ) is extensively used in agriculture for its growth regulating properties as it also presents the ability to control ozone damage to wheat by protecting it against yield reduction [2]. Overall, the attainment of better crops is connected to the mode and concentration of its application. This feature is directly associated with the amount of active compound among the commercial preparations and for which they must be accurately controlled. Furthermore, this widespread practice may

promote contamination of waters, fruits, cereals and vegetables and thus its quantification becomes fundamental.

With respect to residues in food and water, the quantification of CMQ-Cl is extensively described in the literature and some of the methods have also been applied to the analysis of commercial preparations. Due to the overall complexity of the relevant sample matrices, as well as the low concentrations expected in contaminated samples, almost all methods require a detection system capable of producing both quantitative and qualitative information. For this reason mass spectrometry is nearly the only detection system proposed in the literature, and may be associated with liquid chromatography [3–13], thin layer chromatography [14] and electrophoresis systems [15,16]. Independently of the separation approach, the use of mass spectrometry is connected to high capital investment, which makes it unsuitable for routine applications. Conductimetric detection coupled to ion chromatography [17,18] and indirect UV detection coupled to electrophoresis [15,19] are alternative strategies suggested in the literature. In some cases they may be difficult to implement if the equipment is not available in the analytical laboratory. Chemical transformation of the analyte with sodium phenylsulfide [20] or potassium pentafluorothiophenolate [21] has also been exploited in order to enable a gas chromatographic detection. The latter has been considered not sufficiently specific for CMQ and this may be misleading to an analyst using it or adapting it to other procedures [22].

Other proposals in the literature concern electroanalytical systems. Since CMQ is not an electroactive compound, the glassy carbon electrode of a voltammetric detector has been modified with a cation-exchanger membrane [23]. This approach has advantages concerning the environment but it has not been applied to natural samples. Also a potentiometric-based method has been described in the literature and this has advantages regarding emission of effluents of low toxicity [24]. It is not specific for the analyte. It involves a titration with tetraphenylborate and other compounds within samples may react with this chemical and promote analytical errors.

An alternative to previous methods would be a potentiometric determination by means of a CMQ selective electrode. The required equipment is of low cost and is common in analytical laboratories. The system allows a selective detection of the analyte with emission of low toxicity aqueous effluents. In order to obtain adequate working characteristic for performing direct readings of CMQ in various sample matrices, several selective membranes will be proposed and discussed.

The widespread use of growth regulators suggests the need for a routine control of this compound and especially for industries with a daily production of phytopharmaceuticals. In this sense, a simple automatic system such as the flow injection analysis (FIA) technique will be implemented and selective detectors will be constructed with a tubular configuration which is a suitable shape for this flow method.

EXPERIMENTAL

Apparatus

The potential differences between indicating and reference electrodes were measured by means of a Crison mpH 2002 decimillivoltammeter (± 0.1 mV sensitivity) coupled to a Metrohm E 586 recorder. The reference electrode was an Orion (90-02-00)

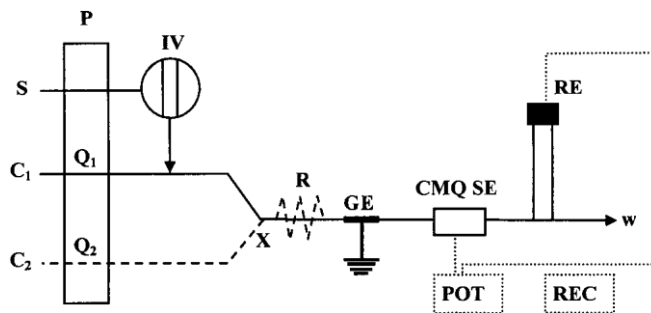


FIGURE 1 FIA set-up for evaluation of working characteristics of the potentiometric detector (channel C_1) and analyzing real samples (channels C_1 and C_2). P: peristaltic pump; S: sample; C_1 : IS or buffer solution, both with $1 \times 10^{-7} \text{ mol L}^{-1}$ of CMQ, or water regarding the analysis of samples; C_2 : buffer solution with $2 \times 10^{-7} \text{ mol L}^{-1}$ of CMQ; IV: injection valve (500 mL); Q_1 $\frac{1}{4}$ Q_2 : 4 mL min^{-1} ; X: confluence point; R: reactor length of 20 cm; GE: ground electrode; CMQ SE: CMQ selective electrode; RE: reference electrode; w: waste; POT: decimillivoltammeter; and REC: recorder.

double junction electrode. The selective electrode, with no internal reference solution presented a tubular configuration and was constructed as described elsewhere [25]. When necessary, pH values were monitored by means of a GAH 110 combined glass electrode connected to a decimillivoltammeter.

The FIA system (Figure 1) comprised a Gilson Minipuls 2 peristaltic pump, fitted with PVC tubing (0.80, 1.60 and/or 2.00 mm i.d.) and a four-way Rheodyne 5020 injection valve holding loops of 100, 200, 500 or 1000 mL. All components were gathered by PTFE tubing (Omnifit, Teflon, 0.8 mm i.d.), Gilson end-fittings and connectors. The support devices for tubular and reference electrodes, as well as the confluence point accessory were constructed in Perspex[®] as reported elsewhere [26].

Reagents and Solutions

All chemicals were of analytical grade and deionised water (conductivity $< 0.1 \text{ mS cm}^{-1}$) was employed. Several reagents were used throughout, namely CMQ-Cl (Fluka), barium chloride (Merck), *o*-phosphoric acid (85%, Merck), and calcium hydroxide (96%, Riedel-deHaeⁿ). For the selective membrane preparation, tetraphenylborate (TPB) sodium (Aldrich), bis(2-ethylhexyl)sebacate (bEHS, Fluka), *o*-nitrophenyl octyl ether (*o*NFOE, Fluka), dibutylphthalate (DBP), poly(vinyl chloride) of high mol-

ecular weight (PVC, Fluka) and tetrahydrofuran (THF, Riedel-deHaeⁿ) were used. Evaluation of the effects of both pH and interfering ionic species at the potentiometric response required sodium hydroxide (Merck), hydrochloric acid (37%, Merck), sodium chloride (Merck), potassium chloride (Merck), magnesium chloride (Riedel-deHaeⁿ), calcium chloride (Merck), ammonium chloride (Merck), hydroxylammonium chloride (Riedel-deHaeⁿ) and diquat (Riedel-deHaeⁿ).

Concentrated standard stock solutions were prepared by rigorous weighing of the corresponding solids and subsequent dilution with ionic strength (IS) adjuster, buffer solution or water. Less concentrated solutions were obtained by careful dilution of the stock solution with the appropriate diluent.

The IS adjuster was a $3.3 \times 10^{-3} \text{ mol L}^{-1}$ BaCl_2 solution, with an IS equal to $1 \times 10^{-2} \text{ mol L}^{-1}$. The simultaneous pH and ionic strength adjustment was obtained with a $\text{H}_3\text{PO}_4/\text{Ca}(\text{OH})_2$ buffer solution having a pH of 6.3 and an IS of $1 \times 10^{-2} \text{ mol L}^{-1}$.

This buffer was prepared by mixing 125.0 mL of a 1×10^{-2} mol L⁻¹ H₃PO₄ solution with 75.0 mL of a 1×10^{-2} mol L⁻¹ Ca(OH)₂ solution, subsequently diluting to a final volume of 500.0 mL. In the double channel set-up (channel C₃ in Figure 1), this buffer solution was prepared with double concentration values but maintaining the same pH. Apart from other chemicals, the carrier solution (either in channel C₁ or C₂ in Figure 1) included an amount of CMQ-Cl equal to 1×10^{-7} or 2×10^{-7} mol L⁻¹, respectively. This small amount of analyte was included to improve baseline stabilization and preserve the selective membrane thus avoiding a continuous leaching of sensor

towards waste. Standard solutions were prepared in carrier solutions, having no CMQ-Cl.

For the study of the effect of pH upon the potentiometric response, a solution of 5.0×10^{-4} mol L⁻¹ CMQ, prepared in IS adjuster, was selected. The interference of other chemicals was studied for 5.0×10^{-5} , 1.0×10^{-4} and 5.0×10^{-4} mol L⁻¹ concentrations of CMQ-Cl and sodium chloride, potassium chloride, magnesium chloride, calcium chloride, ammonium chloride, hydroxylammonium chloride or diquat. All these solutions were made in 3.3×10^{-3} mol L⁻¹ of BaCl₂.

Construction of CMQ Selective Electrodes

The ionic sensor (CMQ-TPB) was prepared by precipitation, mixing 25.0 mL of a 1.0×10^{-2} mol L⁻¹ CMQ-Cl solution and 50.0 mL of a 1.0×10^{-2} mol L⁻¹ sodium TPB solution. The resulting solution was allowed to stand for 24 h, protected from light and in an ice-bath. The precipitate was removed from the solution by filtration. After thorough washing with water, the filtered precipitate was kept in a dark flask inside a dessicator to prevent changes caused by light and humidity.

Sensor solutions were obtained by dissolving about 0.0200 g of the previous ionic sensor in 1.0000 g of oNFOE, BEHS or DBP. The corresponding membrane solutions were prepared by mixing 0.20 mL of the previous solution with 0.09 g of PVC, formerly dissolved in about 2 mL of THF. The overall composition of the previous membranes, designated types A, B and C, respectively, as indicated in Table I.

The selective electrodes were prepared by applying membrane solutions to a tubular conductive support of graphite and epoxy resin [25]. The membrane was allowed to dry for 24 h and then put into 1×10^{-3} mol L⁻¹ CMQ-Cl solution prepared in water. The electrodes were stored in these conditions when not in use.

Procedures

The general working characteristics of the potentiometric detectors were assessed in a single-channel FIA manifold (channel C₁ in Figure 1) using as carrier IS adjuster or

TABLE I Overall membrane composition (% w/w) of the CMQ selective electrodes

Components	A	B	C
CMQ-TPB	1.3	1.3	1.3
oNFOE	67.5	–	–
BEHS	–	66.6	–
DBP	–	–	67.2
PVC	31.2	32.1	31.5

buffer solution to which had been added a small amount of CMQ-Cl. The injection volume was 500 mL and the flow rate 4 mL min^{-1} (Q_1). The results reported are mean values of at least four calibrations. Repeatability was represented by the standard deviation of at least 15 injections of a standard CMQ solution and the response time was estimated in terms of sampling rate.

The influence of pH on the potentiometric response was evaluated for a $5.0 \times 10^{-4} \text{ mol L}^{-1}$ CMQ-Cl solution at different pH values. A small adaptation of the single-channel manifold consisting of a closed loop circuit with continuous circulation of 200 mL of the previous standard solution was required. The pH was altered by addition of small amounts of a saturated NaOH solution or concentrated HCl, and was monitored by an additional potentiometric cell consisting of a combined glass electrode connected to a second decimillivoltammeter. The solutions selected for changing the pH values were highly concentrated in order to prevent significant alterations of the concentration of CMQ in solution. Additionally, and with the exception of Na^{P} , they also avoided the introduction of other chemical species into the system (other than those already present: $\text{Ba}^{2\text{P}}$, Cl^- and CMQ). NaOH was selected instead of

$\text{Ba}(\text{OH})_2$ because of the low solubility of the latter. For each pH measurement an amount of 500 mL was inserted into a 4 mL min^{-1} carrier stream of IS adjuster. Ranges of pH defined as satisfactory allowed a $\pm 5 \text{ mV}$ variation in the analytical signal.

The selectivity was evaluated by the separated solutions method [27], and performed by separately injecting 500 mL of a CMQ-Cl solution and other possible interfering compound solutions into a single-channel FIA manifold with a carrier stream of

$3.3 \times 10^{-3} \text{ mol L}^{-1} \text{ BaCl}_2$, flowing through the detector at 4 mL min^{-1} .

Prior to the analysis of natural samples, total flow-rate and injection volume were optimized in a multivariate way by recording the operating characteristics of the detector for a 1×10^{-6} to $1 \times 10^{-3} \text{ mol L}^{-1}$ concentration range, and under pH and IS adjustment conditions. For this, a double-channel FIA manifold was set-up with 100, 200, 500 or 1000 mL injection volumes, and 3, 4, 5, 6, 7 or 8 mL min^{-1} overall flow-rates. For the analyses of samples, the potentiometric response was recorded in the double-channel FIA manifold, operating simultaneously with channels C_1 and C_2 (Figure 1). Water and buffer solution with a small amount of CMQ-Cl respectively were selected as carriers. This last solution had IS and CMQ concentration values double to those of the single-channel carrier as it would be diluted by half at the confluence point (X, Figure 1).

Sample Preparation

The only phytopharmaceutical preparation of CMQ, commercialized in Portugal as Cycocel[®] was labeled as containing 400 g L^{-1} of CMQ, expressed in the hydrochloride form. To perform recovery trials, and considering the extremely high concentration of CMQ in the sample, a stock solution of sample was first prepared by measuring an exact amount of 39.5 mL and diluting it with water to a final volume of 100.00 mL. A 3000 mL volume of this solution was diluted with water to a final volume of

50.00 mL; the expected concentration was $7.7 \times 10^{-5} \text{ mol L}^{-1}$ CMQ. Recovery trials were then performed with a similarly prepared solution of 3000 mL of a $1.00 \times 10^{-2} \text{ mol L}^{-1}$ CMQ standard solution before completing the final volume up

to 50.00 mL; expected concentration was $7.0 \times 10^{-4} \text{ mol L}^{-1}$ CMQ.

Since it was not possible to find contaminated waters in Portuguese territory, a river water sample was collected and doped with the Cycocel® leading to an expected final

contamination 160 mg L⁻¹. This was considered as the *stock* sample solution. Further dilutions for the potentiometric analysis and recovery trials were performed similarly to those previously mentioned with respect to the phytopharmaceutical preparation.

Results regarding analysis of samples correspond to mean values of eight determinations.

RESULTS AND DISCUSSION

To effect the construction of CMQ selective electrodes with proper working characteristics, three polymeric membranes were prepared, presenting basically the same overall composition but with different plasticizing solvents (*o*NFOE, *b*EHS and *DBP*). Apart from different chemical structures they presented very different polarity characteristics, with dielectric constants equal to 23.6, 6.4, and 4.6 respectively [28]. This property was very important for the ion exchange process within the selective membranes.

The behavior of each CMQ selective electrode was assessed so that the best detector could be selected for the analysis of natural samples.

Working Characteristics of CMQ Selective Electrodes

The working characteristics of the several CMQ selective electrodes, established in a single-channel FIA manifold (Figure 1), were evaluated for both IS and simultaneous pH and IS adjustment conditions. The latter required a pH of 6.3, in agreement with the following pH study. A 500 mL injection volume and a 4 mL min⁻¹ flow-rate were selected with the purpose of establishing experimental conditions close to those in the stationary state. The three types of electrodes presented a very low limit for the linear range, equal to 5 × 10⁻⁶ mol L⁻¹, a similar and low response time, and a quite different repeatability, varying within 0.8 and 1.7 % (Table II).

Regarding the IS adjustment trials, results indicated a slightly higher sensitivity for type C detectors (Table II), suggesting that a low polarity solvent could improve the behavior of the selective electrodes. However, the variation within these results could not probably be attributed only to the physical properties of the solvent, especially considering the small difference between the dielectric constants of the *DBP* and *bEHS*. Supporting this possibility, *Amstrong et al.* reported a decrease of the *o*NFOE dielectric constant to ±12 when incorporated in 33% of *PVC* [29]. This

TABLE II Working characteristics of CMQ selective electrodes with the single-channel FIA manifold

Working characteristics	IS (1 × 10 ⁻² mol L ⁻¹)		pH (6.3) and IS (1 × 10 ⁻² mol L ⁻¹) A			
	B	C	A	B	C	
LLLR ^(a) (mol L ⁻¹)	5.0 × 10 ⁻⁶	5.0 × 10 ⁻⁶	5.0 × 10 ⁻⁶	5.0 × 10 ⁻⁶	5.0 × 10 ⁻⁶	
DL ^(b) (mol L ⁻¹)	2.5 × 10 ⁻⁶	3.9 × 10 ⁻⁶	2.9 × 10 ⁻⁶	2.8 × 10 ⁻⁶	2.2 × 10 ⁻⁶	
Slope (mV decade ⁻¹)	46.9 ± 1.1	42.5 ± 0.3	49.7 ± 0.6	39.5 ± 1.0	51.1 ± 2.6	
Repeatability ^(c) (±mV)	±0.59 (0.9%)	±0.51 (0.8%)	±1.11 (1.2%)	±1.02 (1.7%)	±1.06 (1.4%)	±0.68 (0.7%)
R squared	> 0.9995	> 0.9936	> 0.9989	> 0.9906	> 0.9909	> 0.9953
Sampling rate (h ⁻¹)	80	84	75	85	80	72

pH working range ^(d)	2.3–9.8	4–11	2.3–11.8	-	-	-
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^(a)Lower limit of linear range; ^(b)detection limit; ^(c)at $1.0 \times 10^{-4} \text{ mol L}^{-1}$; ^(d) at $5.0 \times 10^{-4} \text{ mol L}^{-1}$.

phenomenon was explained as the restriction of the free rotation of the mediator sol-vent molecules while in a polymeric matrix.

Studies with solutions of adjusted pH and IS confirmed the better operating characteristics of type C detector (Table II), mainly because of an increase of 8% in sensitivity of the potentiometric response when compared with previous results. Under these conditions, type B working characteristics also improved, particularly in the slope increase of 20%. While slopes of type B and C electrodes improved, those of type A indicated a decrease of 16% in sensitivity. In fact these results demonstrate that low polarity solvents would be more appropriate for preparing CMQ selective electrodes although the chemical structure of DBP must have influenced the potentiometric response leading to the best electrodes.

Operational pH ranges were unusually wide for the three types of electrodes (Table II). Only at very low pH values did the analytical signals show a tendency to a small increase, which would be linked to H^p interference at the potentiometric response. Increasing pH to a value of 10 gave rise to constant potentials at all electrodes, which may be linked to the extent of ionization of CMQ remaining constant even at very high pHs. As the pH increased after 10–12 units, the potentials showed a slight tendency to decrease.

Interference of several ionic species on CMQ determinations was evaluated in terms of potentiometric selectivity coefficients. The three membranes showed a similar behavior (Table III), with a relative order of interference of $Mg^{2p} < Ca^{2p} :: diquat \ll Na^p :: K^p \ll NH_4^p, HONH_3^p$. Basically, interference of all cations concerned was considered negligible. Very low coefficients were recorded for Mg^{2p} and Ca^{2p} , both doubly charged ions. Considering type B and C detectors, potassium showed a slightly stronger effect than sodium, most probably due to its ionic size, closer to that of an ammonium ion. Both ammonium and hydroxylammonium showed the strongest interfering effect, a consequence of the similarity of the ionized group to the CMQ structure. Injection of pure solutions of ammonium ions always gave rise to analytical signals much lower than those produced by the main ion. Because it is included in the same pesticide group as CMQ, diquat interference was evaluated and found to be negligible.

In the single-channel FIA manifold, the preparation of any natural samples would require the use of buffer solution instead of water. If this requirement were only connected to a previous dilution, it would be practically indifferent using either of the two as solvent. However, in a routine application, other samples may be the object of other

TABLE III Potentiometric selectivity coefficients of CMQ selective electrodes

<i>Interference</i> ^(a)	<i>IS</i> (1×10^{-2} mol L ⁻¹)		
	A	B	C
Potassium	-2.350±0.256	-1.877±0.008	-1.699±0.105
Sodium	-2.329±0.143	-2.116±0.012	-2.170±0.113
Magnesium	-4.409±0.160	-4.134±0.027	-4.329±0.248
Calcium	-4.282±0.097	-3.947±0.145	-4.137±0.115
Ammonium	-1.604±0.101	-1.907±0.037	-1.995±0.259
Hydroxylammonium	-0.758±0.256	-1.583±0.024	-2.118±0.126
Diquat	-3.177±0.080	-3.936±0.008	-3.182±0.186

^(a)at $5.0 \times 10^{-4} \text{ mol L}^{-1}$.

treatments that could promote an undesired high consumption of reagents, leading to resources dissipation as well as to higher amount of effluents. For this reason a double-channel manifold that enabled automatic adjustment of required conditions inside the manifold, thus permitting sample preparation with only water, was established.

Optimization of a Double-channel FIA Manifold

Prior to analysis of natural samples, suitable experimental conditions should be selected. Thus, the effect of parameters such as flow-rate and injection volume was object of study. After the confluence point of the two channels (X, Figure 1), a reactor was included in order to guarantee adequate mixing of the solutions. A length of 20 cm was found appropriate for this purpose and kept constant during the several trials.

For each injection volume selected, the increase in flow-rate led to both a slope increase and a lower limit of linear range decrease. These variations were not very perceptible when the flow-rate was changed from 7 to 8 mL min⁻¹. The latter was however selected, since it promoted an increase in sampling rate that ranged from 5% (in case of 100 and 200 mL injection volumes) to 24% (in case of 500 mL loop) therefore reduced the estimated time for analysis.

Though peak heights increased as injection volume was augmented, the recorded sensitivity did not change significantly. The highest slope values were attained with the injection of 500 and 1000 mL sample volumes. Regarding sampling-rate, the former volume enabled the analysis of a larger number of samples per time unit, which decreased to about 42% when 1000 mL was used.

The overall optimization process indicated that satisfactory operating characteristics would be obtained by the injection of 500 mL into a stream crossing the detector at 8 mL min⁻¹. Because the sample plug was here divided by two, the operating characteristics of the type C detectors altered and again characterized giving the mean values indicated in Table IV. Under these conditions, the proposed method has a detection limit of 0.8 mg mL⁻¹ and is able to quantify at least 1.3 mg mL⁻¹. When compared with similar conditions at the single-channel manifold, the slope decreased in about 15% but the sampling rate increased to more than double about 108%.

Analysis of Samples

Before analysis, the potentiometric system was calibrated under the optimum conditions. After dilution with water the mean potentiometric results for the technical concentrate (Cycocel®) were near the labeled amount, at about -4.8% (Table V). They

TABLE IV Working characteristics of CMQ selective electrodes with the double-channel FIA manifold

<i>Working characteristics</i>	<i>pH (6.3) and IS (1 × 10⁻² mol L⁻¹)</i> C
LLLR ^(a) (mol L ⁻¹)	8.0 × 10 ⁻⁶
DL ^(b) (mol L ⁻¹)	5.2 × 10 ⁻⁶
Slope (mV decade ⁻¹)	45.5 ± 0.2
Repeatability ^(c) (± mV)	±0.41 (0.69%)

R squared > 0.9912 Sampling rate
(/h) 150

^(a)Lower limit of linear range; ^(b)detection limit; ^(c)at $1.0 \times 10^{-4} \text{ mol L}^{-1}$.

also presented low standard deviations with a relative value of 2.2%, indicating a good precision. A typical diagram of this analysis is depicted in Figure 2.

River water was collected and doped not with pure standard but with Cycocel®. The estimated amount, expressed in the hydrochloride form, was 160 mg L^{-1} of CMQ. Mean potentiometric results were near this value ($\pm 5.1\%$), and the precision of the method was confirmed by a low standard deviation (Table V).

Because it was not possible to perform the reference procedure, based on ion-chromatography and conductimetric detection [30], the accuracy of the currently proposed method was estimated only by recovery experiments (Figure 2). Recovery trials presented mean values of 97 and 101% with low standard deviations (Table V), demonstrating the accuracy of the proposed method and again confirming good precision.

TABLE V Determination of CMQ in Cycocel® and contaminated river water by the FIA proposed method, the corresponding recovery trials, and the relative error (RE)

Samples	Nominal value (CMQ-Cl)	FIA		RE ^(a)
		CMQ-Cl	Recovery	
Cycocel®	400.0 g L^{-1}	380.1 ± 8.3 (2.2%) g L^{-1}	$101.1 \pm 2.5 \%$	-4.8%
River water	160.0 mg L^{-1}	168.1 ± 7.4 (4.4%) mg L^{-1}	$97.0 \pm 0.5 \%$	$\pm 5.1\%$

^(a)estimated in terms of nominal value.

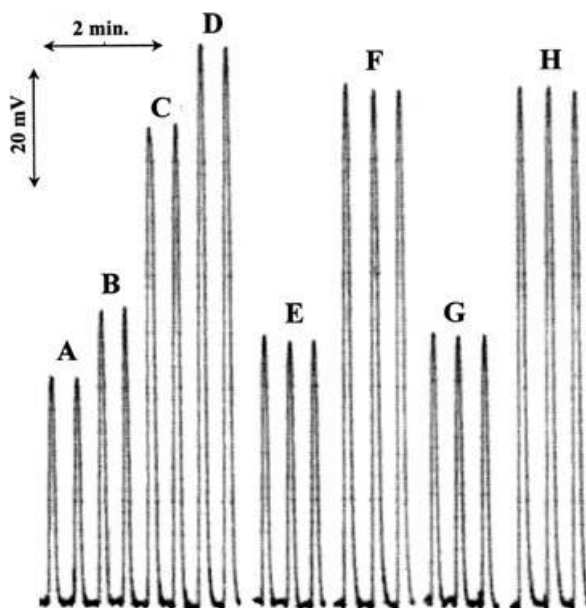


FIGURE 2 Diagram for CMQ standard solutions (A–D), diluted Cycocel® (E,G), and the corresponding recovery trials (F,H). A: $5.0 \times 10^{-5} \text{ mol L}^{-1}$; B: $1.0 \times 10^{-4} \text{ mol L}^{-1}$; C: $5.0 \times 10^{-4} \text{ mol L}^{-1}$; D: $1 \times 10^{-3} \text{ mol L}^{-1}$; E, G: estimated concentration of $7.7 \times 10^{-5} \text{ mol L}^{-1}$; F, H: estimated concentration of $7.0 \times 10^{-4} \text{ mol L}^{-1}$.

CONCLUSIONS

The type C membrane presents a good alternative to other methods previously described in the literature, by CMQ determination in phytopharmaceuticals using potentiometric detection. The proposed system is simple, accurate and inexpensive especially regarding reagent consumption and equipment involved. It is therefore suitable for routine procedures. The sample is prepared by making simple dilution with water, therefore contributing to the preservation of both resources and quality of environment. In terms of inorganic compounds, barium and phosphate generally comprise the majority of the effluents produced at a pH of 6.3. For this reason, an adjustment of pH before discard is not required and the intensity of further pollution from phosphate depends on the receptor; phosphate is of interest to agriculture. The barium may be isolated from solution by a simple precipitation reaction with an appropriate amount of sulphate. Taking into account the need for calibration and assuming 2 readings for each of four standard solutions, the FIA analysis of two samples would produce about 40 mL of effluents. The intrinsic toxicity of the CMQ is low, it is rapidly degraded in soil by microbiological activity and has no influence on soil microflora or fauna [31].

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