Rapid method for determination of glyphosate in groundwater using high performance liquid chromatography and solid-phase extraction after derivatization

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ABSTRACT

The intensive use of pesticides in agriculture has prompted researchers to develop new methods for identifying these pollutants in water. This study sought to validate a high performance liquid chromatography (HPLC) method to determine the concentration of the pesticide glyphosate in groundwater samples by using solid-phase extraction (SPE) filters after derivatization with chloroformate 9-fluorenylmethoxycarbonyl (FMOC-Cl). For the HPLC method, we evaluated the following main validation parameters: linearity, specificity, precision, accuracy, robustness, and limits of detection and quantification. After validation of the method, we determined the concentration of glyphosate in samples from thirteen deep, tubular wells distributed in urban and rural areas in Chapecó, SC, Brazil. The solvent used in the extraction of excess FMOC-Cl was dichloromethane and subsequently filtration was performed on C18 SPE, and injected into the chromatograph column in amino polymer with fluorescence detection. The analytical curve made in ultrapure water was linear, with a correlation coefficient of 0.99. The limits of detection and quantification were 0.24 and 0.07 µg L⁻¹, respectively. Recovery tests in natural waters ranged from 90.37 to 101.70%. Glyphosate was detected in 5 of the thirteen wells evaluated. The highest concentration of glyphosate (6.80 µg L⁻¹) was detected in a countryside well, near the municipal water supply. Despite the low levels of glyphosate detected in our study, any amount present in groundwater samples is worrisome, as these molecules have low ground mobility.

Keywords: agriculture, chromatographic method, pesticides.
Método rápido para determinação de glifosato em água subterrânea usando cromatografia líquida de alta eficiência e extração em fase sólida após derivatização

RESUMO

O uso intensivo de agrotóxicos na agricultura alerta os pesquisadores a desenvolverem novos métodos de identificação destes poluentes em amostras de água. Este estudo teve como objetivo validar um método por cromatografia líquida de alta eficiência (CLAE) para determinar o agrotóxico glifosato em amostras de águas subterrâneas utilizando filtros de extração em fase sólida (SPE) após derivatização com cloroformato de 9-fluorenilmetoxicarbonil (FMOC-Cl). Para o método em CLAE, foram avaliados os principais parâmetros de validação: linearidade, especificidade, precisão, exatidão, robustez, limites de detecção e quantificação. Após validação do método foi determinada a concentração de glifosato em amostras de 13 poços tubulares profundos distribuídos na área urbana e rural no município de Chapecó, SC, Brasil. O solvente utilizado na extração do FMOC-Cl em excesso foi o diclorometano, posteriormente foi realizada filtragem em SPE C18 e injeção no cromatógrafo em coluna polimérica amino com detecção por fluorescência. A curva analítica realizada em água ultrapura apresentou linearidade com coeficiente de correlação de 0,99. Os limites de quantificação e detecção foram de 0,24 e 0,07 µg L⁻¹, respectivamente. Os testes de recuperação em águas naturais variaram de 90,37 a 101,70 %. Dos 13 poços avaliados o glifosato foi detectado em cinco, sendo que a maior concentração de glifosato encontrada foi de 6,80 µg L⁻¹ em um poço localizado na zona rural, próximo à nascente de abastecimento do município. Apesar dos baixos níveis de glifosato detectados em nossa pesquisa, é preocupante o aparecimento destas moléculas em amostras de águas subterrâneas considerando a baixa mobilidade desta molécula no solo.

Palavras-chave: agricultura, método cromatográfico, pesticidas.

1. INTRODUCTION

In 2012, herbicides were the second largest group of pesticides sold globally (Sindiveg, 2012). Among them, the primary and most popularly used is glyphosate [N-(phosphonomethyl) glycine], C₃H₈NO₅P. This powerful agrochemical is used to kill weeds that harm agricultural production, and its deleterious effects on the environment are evident in both aquatic and terrestrial ecosystems, as well as in humans (Amarante Jr. et al., 2002; Pinheiro et al., 2011).

Glyphosate is relatively small in size and has three polar functional groups: carboxyl, amino and phosphonate. The agrochemical has a long half-life, ranging from 49 to 70 days in water, which makes it persistent in soil. High solubility in water (12,000 mg L⁻¹) assists in the transport of glyphosate from terrestrial to aquatic environments. These molecules can be transported to both surface and groundwater, although groundwater samples have not been extensively investigated by the scientific community (Sanchís et al., 2011).

Glyphosate has low acute toxicity to animals because its biochemical mode of action negatively affects shikimic acid, which is found in plants but not in animals (Mirande et al., 2010).

According to data from SINDAG (National Union of Agricultural Products Industry), Brazil leads the world in herbicide and insecticide usage, and the western region of Santa Catarina State has extensive agricultural production in areas that permeate urban centers; this, along with the fact that margin strips of streams and springs are often not respected, causes these chemical compounds to spread easily to aquatic environments. Glyphosate also has
relatively low cost and high efficiency, which results in wide agricultural application. The possibility of resulting environmental contamination must be studied carefully, as there is evidence of harmful environmental effects after prolonged use (Souza et al., 2006).

There are various techniques for determination of glyphosate in environmental samples; however, its high solubility in water, insolvency in organic solvents and low volatility present analytical challenges (Hsu and Whang, 2009). Glyphosate also has high polarity and lacks a chromophore. Detection methods thus require adaptations utilizing derivatization reactions or alteration of some physical property that can be correlated to the amount of glyphosate in the sample (See et al., 2010).

Among the methods used by various authors, the most commonly used derivatization method involves use of FMOC-Cl (9-fluorenylmethyl chloroformate). High fluorescence derivatives can be formed by pre- or post-column derivatization with FMOC-Cl or mercaptoethanol ortho-phthalaldehyde (OPA), respectively (Nedelkoska and Low, 2004). FMOC-Cl can be used for derivatization for both primary and secondary amino groups, and unlike the post-column derivatization, there are no restrictions for mobile phase composition, and the derivation reaction can be better controlled outside of the column (Ghanem et al., 2007).

The Environmental Protection Agency (EPA) recommends technical HPLC with fluorescence detection for analysis of glyphosate, after undergoing column derivatization (USEPA, 2013). However, the pre-column derivatization technique based on 9-fluorenylmethyl chloroformate is currently being employed due to its speed and high sensitivity (Sanchís et al., 2011). To derive allies with this reagent, it is common to use solid phase extraction (SPE) cartridges. Chromatography instruments are used for retention of impurities, filtration processes and analysis. C-18 cartridges are used for extraction of hydrophobic or polar organic analytes in aqueous matrices.

The objectives of this study were (i) to validate the analytical method for rapid determination of glyphosate concentration in groundwater samples, (ii) to optimize the conditions for derivatization of glyphosate, applying SPE to increase the sensitivity of the method and (iii) to test the applicability of the proposed method by evaluating the presence of glyphosate in groundwater samples in Chapecó, SC, Brazil.

2. MATERIALS AND METHODS

2.1. Reagents

Glyphosate (purity 99.5%) was obtained from Chem Service (USA); HPLC grade acetonitrile was obtained from JT Baker (Mexico); and 9-fluorenylmethyl chloroformate (FMOC-Cl), potassium phosphate monobasic, potassium hydroxide (KOH), sodium tetraborate decahydrate and dichloromethane (Chromasolv Plus, 99.9% purity HPLC grade) were purchased from Sigma-Aldrich.

Borate buffer, pH 9 (0.125 M), was prepared with de-ionized water and pH was adjusted with KOH (6 M). The FMOC-Cl (0.005 M) solution was prepared by dissolving the reagent in acetonitrile. A standard glyphosate solution (1000 mg L\(^{-1}\)) was prepared, which was then used to prepare another standard solution of 1 mg L\(^{-1}\). Dilutions were made to cover the range of 5 to 25 µg L\(^{-1}\), all prepared with de-ionized water in polypropylene bottles to prevent any loss of glyphosate on glass surfaces. In this study, we used detection in µg L\(^{-1}\). Water was de-ionized using the Milli-Q system (Millipore, USA). The mobile phase consisted of acetonitrile and phosphate buffer at pH 10, (60:40, v/v). pH was adjusted with 6 M KOH. Figure 1 illustrates the process from sample preparation to liquid chromatography measurement.
2.2. Equipment

The following equipment was used:

- HPLC ProStar 210/215 SDM 270VA. ProStar 410 Autosampler 200VA. ProStar 360 Fluorescence detector 200 VA.
- Analytical columns: NH2, 100A, 5μm, 4.6x2 of 50 mm Kromasil (Switzerland).
- Ultrasound Sonicator Ultra Cleaner 1400 of Unique (Brazil).
- C-18 SPE cartridge and Strata-55um, 70A from Phenomenex (Brazil).

2.3. Validation Method

We used the ANVISA (Brazil) guide to validate analytical and bioanalytical methods. This guide classifies our tests as Class 1. The following tests were performed: specificity, linearity, range, accuracy, repeatability, accuracy, robustness, limit of quantification and detection limit.

The determination of the calibration curve for glyphosate was performed using a polymeric amino column, with the mobile phase consisting of acetonitrile and phosphate buffer (60:40 v/v). The flow of the mobile phase was 0.7 mL min⁻¹, with isocratic method at room temperature. Fluorescence detection was set at a wavelength of 266 nm extinction and 315 nm emission, with a run time of 10 minutes.

The samples were prepared with de-ionized water, with different glyphosate aliquots (5 to 25 µg L⁻¹) to obtain the required concentrations. The total volume per concentration was brought up to 5 mL, after which 0.5 mL of borate buffer was added to maintain alkaline pH.

Derivatization occurred with the addition of 2 mL of FMOC-Cl (0.005 M). To avoid FMOC-Cl precipitation, a volume of water was added prior to reagent addition. The reaction was maintained at room temperature for the following reaction times: 30, 45 and 60 minutes. The excess FMOC-Cl was removed by extraction with 3 mL of dichloromethane, and the end of the supernatant was filtered using a SPE cartridge prior to injection into the HPLC column, as described in Section 2.5.

Specificity/selectivity: they were observed with the standard chromatogram and a sample analysis without glyphosate (blank). To secure and to avoid interference from other compounds in the sample, tests were conducted by passing samples of only water, buffer, FMOC-Cl and the mobile phase through the column, thus ensuring that the interfering peaks cannot be glyphosate.

Precision: it was evaluated through repeatability. It was determined by preparation, under the same conditions, of 9 analytical solutions of the sample at working concentration. Relative standard deviation (RSD) was considered acceptable, according to resolution 899/2003 ANVISA (Brasil, 2003).

Accuracy: it was assessed based on the degree of recovery, expressed as percentage of glyphosate solutions fortified with known amounts of glyphosate in triplicate, determined by the ratio between the experimental and theoretical average concentrations using Equation 1:
\[
\text{Accuracy} = \frac{\text{Experimental average concentration}}{\text{Theoretical average concentration}} \times 100
\]  

Robustness: Tests for robustness were performed with slight variations in the analytical parameters, including the variation in mobile phase, mobile phase composition, temperature, and flow rate.

2.4. Groundwater analyses

After validation of analytical methods, samples were taken from deep wells within the boundaries of Chapecó, Brazil. Thirteen wells were chosen, and some points are located in a rural area with strong agricultural influence. The coordinates described herein are depicted in the UTM system WGS 84 22J area (Figure 2).

2.5. SPE cartridge extraction

The cartridge sorbent material is first activated by passing an appropriate solvent through to condition the solid surface (in this case water). The dichloromethane extraction supernatant is subsequently passed through the cartridge. At this stage, the unreacted FMOC-Cl from the dichloromethane extraction is retained on the stationary phase, allowing passage of the more polar derivatized glyphosate (Figure 3). After this step, washing tests with different volumes

Figure 2. Geographical locations of water samples.
of water (1, 2 and 3 mL) were performed to determine the optimal volume. The optimal volume found was 1mL of water to washing cartridges.

![Figure 3. SPE cartridge extraction overview.](image)

3. RESULTS AND DISCUSSION

3.1. Glyphosate chromatograms

Figure 4 shows the chromatograms of the analyses initially with ultrapure water, known as white, and without (a) and with (b) the use of the cartridge. Samples fortified at a concentration of 5 µg L⁻¹ presented a chromatogram peak time of 6.5 minutes. In (c) the glyphosate peak becomes partially covered by FMOC-Cl reagent around 3 minutes. In (d) the chromatogram is clean and well-defined due to the use of SPE cartridges.

![Figure 4. Chromatograms of real and white analyses with and without SPE cartridge use. (a) Ultrapure water (white) without use of the cartridge, (b) ultrapure water (white) with use of the cartridge, (c) glyphosate (5 µg L⁻¹) partially covered by FMOC-Cl and (d) glyphosate (5 µg L⁻¹) with use of SPE cartridges.](image)
3.2. Specificity and selectivity

The method used to specificity and selectivity of the analysis was specific and selective for glyphosate through comparative analysis of chromatogram with the standard and with a sample analysis without glyphosate (blank); we did not observe any interference in the analyses.

3.3. Linearity of the calibration curve

Linearity analysis of our five different concentrations of analytes resulted in a correlation coefficient (R) of greater than 0.999 for all sample concentrations. ANVISA (Brasil, 2003) recommends a correlation coefficient of 0.99. This coefficient demonstrates that the detector response was linear for the range of concentrations used (Figure 5).

![Analytical curve for glyphosate.](image)

**Figure 5.** Analytical curve for glyphosate, the error bars indicate ±1 standard deviation.

3.4. Precision

To test repeatability, three replicates were performed each for low, medium and high concentrations (5 µg L⁻¹, 15 µg L⁻¹ and 25 µg L⁻¹) (Table 1).

<table>
<thead>
<tr>
<th>Concentration (µg L⁻¹)</th>
<th>Retrieved Concentration (µg L⁻¹)</th>
<th>Average</th>
<th>Standard Deviation</th>
<th>RSD (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>4.9803</td>
<td>4.7788</td>
<td>0.3713</td>
<td>7.7</td>
</tr>
<tr>
<td></td>
<td>5.0059</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>4.3503</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>14.4021</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>15</td>
<td>13.1521</td>
<td>14.1865</td>
<td>0.9453</td>
<td>6.6</td>
</tr>
<tr>
<td></td>
<td>15.0054</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>24.9523</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>25</td>
<td>25.8875</td>
<td>25.4325</td>
<td>0.4681</td>
<td>1.8</td>
</tr>
<tr>
<td></td>
<td>25.4577</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 1. Data obtained from the precision assay.
Considering data obtained from the nine analytical solutions prepared, the method demonstrated repeatability with a relative standard deviation (RSD%) of up to 7.7% among samples prepared on the same day under the same conditions, within the maximum permitted by applicable law (Brasil, 2003).

### 3.5. Limits of detection and quantification

Limits of Detection (LD) and Limits of Quantification (LQ) were estimated using the standard deviation (SD) of the intercept and the value of the slope (SC) of the calibration curve according to Equations 2 and 3:

\[
LD = \frac{SD \times 3.3}{SC}
\]  
(2)

\[
LQ = \frac{SD \times 10}{SC}
\]  
(3)

The limits of detection and quantification obtained in this study are shown in Table 2.

<table>
<thead>
<tr>
<th>Limit of Detection (µg L(^{-1}))</th>
<th>Limit of Quantification (µg L(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water Sample</td>
<td>0.07</td>
</tr>
<tr>
<td></td>
<td>0.24</td>
</tr>
</tbody>
</table>

Sanchís et al. (2011) identified in their studies limits of detection and quantification of up to 3.2 and 9.6 ng L\(^{-1}\), respectively, for glyphosate samples using SPE cartridge extraction methods, but with linearity (R) less than 0.98. Considering the maximum allowed under current Brazilian environmental legislation (500 µg L\(^{-1}\)), these values were considered satisfactory.

### 3.6. Accuracy

The results for accuracy at concentrations of 5, 15 and 25 µg L\(^{-1}\) showed a mean recovery percentage of between 90.37% and 101.79% (Table 3).

<table>
<thead>
<tr>
<th>Glyphosate Recovery in Water Samples.</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 µg L(^{-1})</td>
</tr>
<tr>
<td>15 µg L(^{-1})</td>
</tr>
<tr>
<td>25 µg L(^{-1})</td>
</tr>
<tr>
<td>Water sample</td>
</tr>
<tr>
<td>95.53%</td>
</tr>
<tr>
<td>90.37%</td>
</tr>
<tr>
<td>101.79%</td>
</tr>
</tbody>
</table>

Recovery was within the acceptable limits of 80-120% (Brasil, 2003), indicating that the method used is appropriate for the analysis of glyphosate in water.

### 3.7. Robustness

The same sample with slight variations in the analytical parameters (i.e., mobile phase pH, mobile phase composition, temperature, and flow rate) revealed significant variations in the robustness test results. The optimum pH was 10 and the optimal mobile phase composition ratio was 60:40 acetonitrile:phosphate buffer, as described by Nedelkoska and Low (2004).

### 3.8. Derivatization

Figure 6 depicts the derivatization reaction of glyphosate in pH 9 reagent with FMOC-Cl. This reaction is an aminolysis reaction, in which the acyl chloride (FMOC-Cl) reacts with the secondary amine (glyphosate), producing the corresponding amide (FMOC-glyphosate). The alkaline sequestrant acts as proton-maintaining balanced reaction to the right, through the use of the co-product HCl. In the case of the reaction being performed in acidic or neutral pH,
released HCl consumes the rest of the amine group of glyphosate by protonation, causing it to lose group nucleophilicity and decreasing the reaction yield.

![Figure 6. Reaction between FMOC-Cl and glyphosate.](image)

Test trials with different glyphosate derivatization times with FMOC-Cl were performed. Trials were performed with 30, 45 and 60 minutes of derivatization in a concentration of 10 µg L\(^{-1}\). Increased reaction time resulted in an increase in peak area; however, this increase was not significant after 45 minutes (Table 4).

**Table 4.** Results of peak area in relation to the reaction time for derivatization.

<table>
<thead>
<tr>
<th>Time (min(^{-1}))</th>
<th>Peak area</th>
</tr>
</thead>
<tbody>
<tr>
<td>30</td>
<td>27.67 ± 1.78(^{a})</td>
</tr>
<tr>
<td>45</td>
<td>94.37 ± 2.91(^{b})</td>
</tr>
<tr>
<td>60</td>
<td>91.03 ± 3.11(^{b})</td>
</tr>
</tbody>
</table>

Values followed by the same letter in columns do not differ by Tukey test at 95% confidence level.

**3.9. Determination of glyphosate in water samples**

For the determination of glyphosate in groundwater samples, the same procedures from the analysis method validation guide were used. Glyphosate was detected in five of the thirteen samples tested in low concentrations (Table 5).

**Table 5.** Glyphosate concentration in groundwater samples.

<table>
<thead>
<tr>
<th>Location</th>
<th>Glyphosate (µg L(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>Point 1</td>
<td>0.2459</td>
</tr>
<tr>
<td>Point 2</td>
<td>0.4475</td>
</tr>
<tr>
<td>Point 3</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 4</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 5</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 6</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 7</td>
<td>0.4475</td>
</tr>
<tr>
<td>Point 8</td>
<td>6.7972</td>
</tr>
<tr>
<td>Point 9</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 10</td>
<td>1.5158</td>
</tr>
<tr>
<td>Point 11</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 12</td>
<td>0.0000</td>
</tr>
<tr>
<td>Point 13</td>
<td>0.0000</td>
</tr>
</tbody>
</table>
According to chromatogram analyses, the highest concentrations of glyphosate were found in sample Sections 8 and 10, both of which are located in agricultural areas. Section 8 is located near the source of the São José watershed, which supplies water to the population of Chapecó.

Current Brazilian legislation (Brasil, 2011) establishes a maximum allowable value (MAV) for glyphosate of 500 mg L\(^{-1}\) in post-treatment samples. According to Item 6 of the same ordinance, detection of pesticides in samples above the MAV should be analyzed in conjunction with the historical control of water quality.

CONAMA Resolution No. 357 established on March 17, 2005 (Brazilian law) provides for the classification of water bodies and creation of environmental guidelines for its framework, and establishes the conditions and standards for effluent discharge. Rivers are in Classes 1 and 3, wherein acceptable levels of glyphosate are 65 mg L\(^{-1}\) and 280 mg L\(^{-1}\) respectively. Our values for groundwater samples are thus well below the limits established by law.

The methods presented in this study have a few advantages over traditional methods. First, very low levels of glyphosate can be detected in water (i.e., sensitivity is higher). The cost is also lower, because the reagents are inexpensive and easily prepared. Efficiency is higher and time expenditure is reduced, because the sample is prepared for reading in a shorter amount of time. This facilitates the analysis of greater numbers of samples compared to other methods.

Despite the low levels of glyphosate detected, detection of any amount of this substance in groundwater samples is worrisome. This is because glyphosate is not easily leached and has rapid adsorption to soil, and is therefore unlikely to contaminate groundwater (Amarante Jr. et al., 2002). The pesticide has rarely been detected in water samples; but in general, this is due to the difficulty of separating the compound, as well as a paucity of studies due to lack of concern (i.e., because the substance is not considered a serious water contaminant).

Glyphosate is a highly polar compound, requiring extraction from aqueous samples with metallic cations and inorganic anions, which can be difficult. If chromophore groups are contained, glyphosate does not absorb visible electromagnetic radiation, and thus can only be detected by fluorescence or colorimetric methods indirectly, or by derivatization (Amarante Jr. et al., 2002).

These results reinforce the importance of more stringent legislation regarding permissible levels of pesticides in water. Current legislation also does not provide guidance for monitoring of synergistic outcomes of agrochemicals, which we think should be considered. This is important because dozens of compounds can be used in farming, and combinations of compounds may form byproducts or produce synergistic effects that are harmful to both humans and ecosystem health.

4. CONCLUSION

The use of pesticides in agricultural practices, particularly glyphosate, raises concerns about surface and groundwater quality. Newer methods with greater accuracy and sensitivity should be developed in order to quantify contaminants at low concentrations. Liquid chromatography using solid phase extraction after derivatization with FMOC-Cl was effective for the detection of glyphosate in water, and allows greater sensitivity for quantification of glyphosate in parts per billion (ppb). Our trials measuring glyphosate levels in groundwater in urban and rural areas of Chapecó found concentrations within the limits permitted by law; but their presence in rural areas confirms widespread use of this compound, which may affect the water quality in the municipality. Our study suggests that further research should be...
conducted to analyze groundwater glyphosate in this region in order to aid in the protection of the regional ecosystem and preservation of water supply quality.

5. ACKNOWLEDGEMENTS

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6. REFERENCES


