**Vortex-assisted ionic liquid-based dispersive liquid-liquid micro-extraction and spectrofluorometric determination of Rhodamine B in anti-freeze, lipstick, liquid soap, matches and red pencil core**

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**ABSTRACT:** A simple, centrifugeless vortex-assisted ionic liquid-based dispersive liquid-liquid micro-extraction method was developed for spectrofluorometric determination of Rhodamine B dye. Analytical extraction parameters including pH, type and amount of ionic liquid, salt concentration, vortex rate and time were examined and optimized. Matrix effects of potentially interfering ions and dyes were investigated and their tolerable concentrations were determined by applying optimum conditions. Detection limit and preconcentration factor were determined 0.01 µg L⁻¹ and 30, respectively. Dynamic range of the method was linear at RhB concentration range between 0.025 and 1000 µg L⁻¹. Relative standard deviations were 3.9% for the ten replicates measurements of 0.125 µg L⁻¹ Rhodamine B. The method was successfully applied to different samples including anti-freezes, liquid soap, matches, red pencil core and lipsticks. RhB concentrations of liquid and solid samples were determined between 2.05 µg mL⁻¹ and 10.05 µg mL⁻¹ and 307 µg g⁻¹ and 2.75 mg g⁻¹, respectively.

**Keywords:** Dispersive liquid-liquid micro-extraction, ionic liquid, rhodamine B, spectrofluorometry

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Bu çalışmanın bir kısmı 1-4 Mayıs 2019 tarihlerinde Didim/AYDIN’daki düzenlenen ‘4th International Conference on New Trends in Chemometrics and Applications’ isimli konferansta sözlü sunum olarak sunulmuştur.
INTRODUCTION

Rhodamine B (RhB) is highly water soluble xanthene organic dye containing aromatic benzene rings. Although it has been banned to use in food and different products, it has been still used illegally in cosmetic, food, textile and industrial samples due to its low cost, stability and high effectiveness in dying process (Chen et al., 2014). RhB is harmful if swallowed and inhaled by human and animals. It damages to human skin and eyes if in direct contact (Ranjbari and Hadjmohammadi, 2015). Its toxicity, carcinogenicity, neurotoxicity and chronic toxicity towards human and animals have been proved (Alesso et al., 2012; Qi et al., 2014). For these reasons, detection and determination of RhB in different available products are important and essential.

Therefore, scientists in the world, especially analytical chemists have studied to determine RhB contents of commercially available products by using instrumental analytical techniques and devices including liquid chromatography-mass spectrometry (LC-MS) (Li et al., 2013; Cheng and Tsai, 2016), ultra violet-visible spectrophotometry (Unsal et al., 2014a; Unsal et al., 2014b), spectrofluorometry (Fu et al., 2015; Roostaie et al., 2017), electrochemical methods (Sun and Yang, 2017) and high performance liquid chromatography (HPLC) (Tatebe et al., 2014; Xu et al., 2016). Although chromatographic methods supply sensitive quantitative determination, it could not be available in many laboratories because the instrument price is considerably high and it has needed highly skilled and experienced operators (Bişgin, 2018; Li et al., 2020). Due to their simplicity and lower cost than other instruments, UV-visible spectrophotometry and spectrofluorometry are the attractive techniques (Bişgin, 2019a). However, these techniques have not ensured satisfactory results due to necessity for pretreatments of complicated sample matrix and the insufficient sensitivity (Liang et al., 2015). Therefore, it is very important to develop a sensitive, simple, cost-effective and rapid method for the determination of Rhodamine B (Liu et al., 2013).

Particularly, spectrofluorometry is low-cost, simple, rapid and highly sensitive. It has been widely used for direct determination of fluorescence dye like RhB. Unfortunately, complex matrix interference and low concentration of RhB limit the direct determination of RhB contents in sample. Therefore, it has great importance to apply at least one or more pretreatment process prior to the determination of trace target analyte RhB in complicated samples (Bakheet and Zhu, 2017).

In order to isolate the RhB from complex matrix sample, different separation, enrichment and purification methods including solid-phase extraction (SPE) (Su et al., 2015; Yan et al., 2017), cloud point extraction (CPE) (Ghasemi and Kaykhaii, 2016; Bişgin et al., 2018), deep eutectic solvent extraction (DES) (Wang et al., 2017; Yilmaz and Soylak, 2018) and dispersive liquid-liquid micro-extraction (DLLME) (Unsal et al., 2014a; Bağda and Tuzen, 2017) have been used and reported. DLLME method is interesting because of its simplicity, considerably high preconcentration factor, rapid and effective phase separation. In addition, this method uses only very little amounts of toxic organic solvents (Bişgin, 2019b). Therefore, chemists have developed new methods with using surfactants, ultrasonic bath, vortex and ionic liquids based on DLLME method. These are surfactant assisted dispersive liquid-liquid micro extraction (SA-DLLME) (Sadeghi et al., 2018), ultrasound assisted dispersive liquid-liquid micro extraction (UA-DLLME) (Altunay et al., 2019), vortex assisted dispersive liquid-liquid micro extraction (VA-DLLME) (Elik et al., 2017) for extraction of trace substances.

Aim of this study was to develop a VA-IL-DLLME method for extraction and spectrofluorometric determination of RhB. For this purpose, ionic liquid 1-butyl-3-methylimidazolium hexafluorophosphate
[C₄MIM][PF₆] and vortex were used. Several analytical parameters, affecting recovery results, were investigated and optimized. The optimized method was applied to anti-freezes, lipsticks, liquid soap, matches and red pencil core samples to determine their RhB contents. Finally, analytical characteristics and performance of the method were evaluated and compared with reported studies.

MATERIALS AND METHOD

Chemicals
Analytical grade chemicals were used in the experiments and purchased from Merck (Merck, Darmstadt, Germany) and Sigma (Sigma–Aldrich, St. Louis, MO, USA). In the preparation of buffer solutions H₃PO₄-NaH₂PO₄ for pH 2 and pH 3, HAc-NaAc for pH 4 and pH 5 and NaH₂PO₄-Na₂HPO₄ for pH 6, pH 7 and pH 8 were used.

100 µg mL⁻¹ of RhB stock dye solution was prepared and stored in refrigerator at 4 °C. More diluted dye solutions were prepared daily from the RhB stock dye solution. The chemical structure of RhB is given in Figure 1a.

NaCl, KCl and Na₂SO₄ were used as model electrolytes in the extraction experiments. Extensively used dyes and common metal ions (Table 1) were used to prepare model matrix solutions.

1-butyl-3-methylimidazolium hexafluorophosphate [C₄MIM][PF₆] as an ionic liquid, was used to extract RhB dye molecules from aqueous phase to organic phase. The chemical structure of [C₄MIM][PF₆] is given in Figure 1b. Pure ethanol was used for solving and diluting the bulk ionic liquid rich phase.

![Chemical structures of RhB and [C₄MIM][PF₆]](image)

Figure 1. The chemical structures of RhB (a) and [C₄MIM][PF₆] (b)

Instruments
RhB emission measurements were performed by using Hitachi F-7100 fluorescence spectrophotometer (Hitachi, Tokyo, Japan). pH adjustments of aqueous and buffer solutions were performed with using a digital pH-meter (Hanna Instruments, USA). A Velp ZX3 model advanced vortex mixer (Velp Scientifica, Italy) was used for vortexing aqueous mixture. A double-beam Shimadzu UV-160 A (Shimadzu Corporation, Kyoto, Japan) UV-Vis Spectrophotometer was used to determine UV-Vis spectra of the real samples.

VA-IL-DLLME Procedure
Fifteen milliliters of sample solution containing 0.250 µg of RhB and 0.3 mol L⁻¹ of KCl was buffered to pH 2.5. The solution was decanted to centrifuge tube. 240 µL of [C₄MIM][PF₆] was added to solution.
The mixture containing two clear phases was vortexed for 60 seconds at 2400 rpm. Organic fine droplets containing RhB molecules were collected bottom of the centrifuge tube after vortex process. Aqueous upper phase was removed by a syringe. Analyte rich phase was diluted to 500 µL of final volume with ethanol. RhB dye was excited at 520 nm of excitation wavelength and measured at 576 nm of emission wavelength.

**Pretreatment of real samples**

**Anti-freeze samples:** 1.0 mL of each anti-freeze samples were directly subjected to optimized method after pH adjustments.

**Lipstick Samples:** RhB amounts of each of 25 mg lipstick samples were dissolved with CCl₄. RhB molecules in the organic phase were extracted to water phase by 25 mL of 0.1 mol L⁻¹ NaOH with using a separatory funnel. 5 mL of extract was applied to method after pH adjustments.

**Liquid soap:** An amount of 10.0 g (8 mL) liquid soap sample was dried in an oven at 50 °C. Dried sample was dissolved in water and diluted to 2.0 mL. The solution was analyzed with applying the developed method after filtering through blue band filter paper and adjusting pH.

**Red pencil core:** An amount of 12.5 mg red pencil core was accurately weighed. Then it was dissolved in pure water and subsequently filtered through blue band filter paper. The filtrate was diluted to 25 mL in a volumetric flask. Then 1.0 mL of the solution was applied to the developed method after pH adjustments.

**Matches:** Matches samples were weighed (10 mg) and dissolved in pure water. Then, samples were filtered through blue band filter paper and solution was diluted to 25 mL with distilled water. 1.0 mL of solution was subjected to the developed method after pH adjustments.

**RESULTS AND DISCUSSION**

UV-Vis, fluorescence excitation and fluorescence emission data for the determination of RhB dye in anti-freeze, red pencil core, matches, pure aqueous solution, lipstick and liquid soap samples are given in Fig. 2, Fig. 3 and Fig. 4, respectively.
Influence of pH

Aqueous media pH is important for pH sensitive and dependent dye such as RhB. The influences of pH on the extractions of RhB molecules into ionic liquid organic phase were investigated between pH 2 and pH 8 by using 15 mL of model solutions containing 0.250 µg RhB. The results are presented in Fig. 5 with standard deviations. Quantitative extractions of RhB were observed in the pH range of 2-3. In acidic media, RhB molecules were surrounded by ionic liquid molecules. Thus, RhB dye was quantitatively (>95%) extracted from large aqueous phase into small ionic liquid phase. At higher pH values, the cationic form of RhB molecules turned into to the neutral one and its recovery value decreased. Similar results were observed in literature value (Soylak et al., 2011). Therefore, all subsequent studies were performed at pH 2.5 by using H₃PO₄//NaH₂PO₄ buffer.
**Effect of ionic liquid amount**

Ionic liquid amount is important parameter for complete extraction of RhB molecules from aqueous phase to IL rich phase. In order to extract RhB dye, [C4MIM][PF6] and [C4MIM][BF4] ionic liquids were used. In presence of [C4MIM][BF4] extraction was not performed because blurred solution formed and phase separation was not obtained. Therefore, [C4MIM][PF6] was selected and then used in all further extraction studies. IL volume was examined between 60-270 µL. Results are depicted in Fig. 6 with standard deviations. Recovery values increased with increasing IL amount up to 210 µL and then remained constant. Quantitative extraction was obtained between 210 and 270 µL. Hence, 240 µL of [C4MIM][PF6] was chosen optimum amount for further studies.

**Effect of salt concentration**

Extraction efficiency usually increases with increasing ionic strength of aqueous solution because of salting out effect (Chen and Zhu, 2016). The effect of ionic strength was evaluated by using NaCl, KCl and Na2SO4 as model electrolytes on the extraction efficiency. In presence of KCl, stable, repeatable and the quantitative extraction value was obtained. Therefore, KCl was selected for subsequent experiments. Salt
concentration was examined a concentration range of 0-0.04 mol L\(^{-1}\). results are given in Fig. 7 with standard deviations. Over this concentration interval, the cloudy solution was formed. RhB recoveries increased with increasing KCl concentration up to 0.2 mol L\(^{-1}\) and remain stable above this value. Therefore, 0.3 mol L\(^{-1}\) of KCl concentration was applied in all further experimental studies.

![Figure 7. Effect of KCl concentration, N=4](image)

**Influence of vortex time**

Optimization of vortex time is necessary to achieve equilibrium and entire dispersion of IL in aqueous phase by resulting in complete extraction of RhB molecules from aqueous phase to IL droplets. Optimized enough vortex time is the minimum time to obtain equilibrium between aqueous and IL rich phase by ensuring higher extraction efficiency (Bağda and Tuzen, 2017). Vortex time was investigated at 2400 rpm between 10 and 80 seconds. 2400 rpm was selected an optimum because enough agitation was not achieved below than this value. Results are presented in Fig. 8 with standard deviations. Recoveries increased with increasing vortex time. Quantitative extraction and sufficient formation of ionic droplets were obtained only after 50 seconds. Therefore, 60 seconds vortexing at 2400 rpm was chosen optimum for further experimental studies.

![Figure 8. Influence of vortex time, N=4](image)
Effect of matrix

It is necessary that effect of matrix components on the trace determination of analyte should be examined in detection and determination process. Interference effects of matrix components including widely used dyes, potentially available anions and cations were also examined. Results are given in Table 1 with standard deviations. Dyes and ions at concentrations levels, which are given in Table 1, did not interfere the analysis at the optimized conditions of the method. At the optimum conditions of the developed method, ionic liquid \([\text{C}_4\text{MIM}]\text{[PF}_6]\) extracted only RhB molecules as quantitative from aqueous sample solution in presence of other matrix dyes given in Table 1. This circumstance has provided the selectivity to the proposed method. In addition, fluorescence determination provided selectivity to the developed method because RhB has the fluorescence characteristic among other matrix dyes, which are given in Table 1. The used dyes in matrix study have not fluorescence characteristics therefore, any emission wavelengths, which could be interfered the fluorescence determination, were not observed by fluorescence excitation with using fluorescence spectrometer. Thus, only RhB molecules were determined applying both excitation and emission wavelength. In addition, differences between excitation and emission wavelength of RhB increased the selectivity. Moreover, excitation wavelengths and emission wavelengths are specific for each dye that has fluorescence characteristic.

Table 1. Matrix effect of the potentially interfering components, N=4

| Dyes or Ions  | Added species      | Concentration, µg mL\(^{-1}\) | Recovery, % |
|--------------|--------------------|-------------------------------|-------------|
| Tartrazine   | -                  | 4                             | \(^{99}\pm2\) |
| Sunset yellow| -                  | 3                             | 98\pm3      |
| Brilliant blue| -                 | 4                             | 97\pm3      |
| Ponceau 4R   | -                  | 1                             | 97\pm4      |
| Allura red   | -                  | 1                             | 100\pm1     |
| Co\(^{2+}\)  | Co(NO\(_3\))\(_2\).6H\(_2\)O | 100                           | 99\pm2      |
| Ni\(^{2+}\)  | Ni(NO\(_3\))\(_2\).6H\(_2\)O | 100                           | 98\pm3      |
| Na\(^{+}\)   | NaNO\(_3\)         | 1000                          | 97\pm4      |
| Mg\(^{2+}\)  | Mg(NO\(_3\))\(_2\).6H\(_2\)O | 1000                          | 96\pm2      |
| Cu\(^{2+}\)  | Cu(NO\(_3\))\(_2\).5H\(_2\)O | 100                           | 98\pm1      |
| Cr\(^{3+}\)  | Cr(NO\(_3\))\(_3\).3H\(_2\)O | 25                            | 99\pm4      |
| Cd\(^{2+}\)  | Cd(NO\(_3\))\(_2\).6H\(_2\)O | 100                           | 99\pm2      |
| K\(^{+}\)    | KNO\(_3\)          | 1000                          | 98\pm3      |
| Mn\(^{2+}\)  | Mn(NO\(_3\))\(_2\).4H\(_2\)O | 100                           | 97\pm3      |
| Ca\(^{2+}\)  | CaCl\(_2\)         | 1000                          | 96\pm4      |
| Pb\(^{2+}\)  | Pb(NO\(_3\))\(_2\) | 100                           | 96\pm1      |
| Al\(^{3+}\)  | Al(NO\(_3\))\(_3\).9H\(_2\)O | 50                            | 95\pm2      |
| Cl\(^{-}\)   | NaCl               | 1000                          | 98\pm3      |
| PO\(_4^{3-}\) | Na\(_3\)PO\(_4\)  | 1000                          | 100\pm2     |
| SO\(_4^{2-}\) | Na\(_2\)SO\(_4\)  | 100                           | 99\pm4      |
| NO\(_3^{-}\) | NaNO\(_3\)        | 1000                          | 97\pm3      |

\(^{a}\)Mean\pmstandard deviation

Real sample applications

This VA-IL-DLLME method was applied to determine RhB contents of different samples including anti-freezes, liquid soap, lipsticks, matches and red pencil core. The standard addition technique was also used for the determination of real samples by adding 0.125 µg and 0.250 µg of RhB. First, RhB amount of each aqueous solution belonging the real samples including anti-freezes, liquid soap, lipsticks, matches and
red pencil core were determined. Then, 0.125 µg and 0.250 µg of RhB dyes were separately added to the each of same solutions of the real samples. Finally, the solutions were subjected to developed method. Recovery results and RhB contents of the analyzed samples are given in Table 2 with standard deviations. Calculated recovery results with using standard addition technique were between % 96 and % 101. RhB concentrations of anti-freeze, liquid soap, lipsticks, matches and red pencil were separately determined and calculated for each samples according to applied pretreatment procedures and necessary dilutions which were given in “Pretreatment of real samples” part. RhB contents of liquid and solid samples were determined between 2.05 µg mL⁻¹ and 10.05 µg mL⁻¹ and 307 µg g⁻¹ and 2.75 mg g⁻¹, respectively. The accuracy of the method with satisfactory quantitative results indicated the applicability of the method for determination of RhB in different matrix samples.

Table 2. Recoveries and RhB contents of real samples with standard addition technique, N=4

| Sample          | RhB added (µg) | RhB found (µg) | % Recovery | RhB content |
|-----------------|---------------|---------------|------------|-------------|
| Anti-freeze 1   | -             | 0.112±0.005   | 98±4       | 6.52±0.05 (µg mL⁻¹) |
|                 | 0.125         | 0.233±0.011   | 99±3       |             |
|                 | 0.250         | 0.359±0.018   | 99±3       |             |
| Anti-freeze 2   | -             | 0.165±0.007   | 96±4       | 10.05±0.12 (µg mL⁻¹) |
|                 | 0.125         | 0.279±0.014   | 99±3       |             |
|                 | 0.250         | 0.411±0.019   | 99±3       |             |
| Liquid soap     | -             | 0.202±0.008   | 97±3       | 2.05±0.04 (µg mL⁻¹) |
|                 | 0.125         | 0.318±0.012   | 98±4       |             |
|                 | 0.250         | 0.443±0.017   | 98±4       |             |
| Lipstick 1      | -             | 0.245±0.009   | 99±2       | 465±15 (µg g⁻¹) |
|                 | 0.125         | 0.368±0.013   | 98±4       |             |
|                 | 0.250         | 0.485±0.023   | 98±4       |             |
| Lipstick 2      | -             | 0.230±0.010   | 97±3       | 307±11 (µg g⁻¹) |
|                 | 0.125         | 0.345±0.016   | 97±3       |             |
|                 | 0.250         | 0.461±0.021   | 97±3       |             |
| Matches         | -             | 0.175±0.007   | 96±4       | 1.07±0.06 (mg g⁻¹) |
|                 | 0.125         | 0.289±0.011   | 99±3       |             |
|                 | 0.250         | 0.421±0.014   | 99±3       |             |
| Red pencil core | -             | 0.150±0.006   | 98±2       | 2.75±0.03 (mg g⁻¹) |
|                 | 0.125         | 0.269±0.010   | 101±3      |             |
|                 | 0.250         | 0.404±0.014   | 101±3      |             |

aMean±standard deviation

Analytical characteristics of the method

Analytic specifications of the proposed method were investigated. Limit of detection (LOD) of the method was determined as 0.01 µg L⁻¹ for RhB dye. LOD was calculated according to (BT+3SD/m)/PF where BT is average of 21 blank solutions, SD is standard deviation of 21 blank solutions, m is slope of the calibration curve and PF is preconcentration factor. Linear dynamic range (LDR) was obtained at the concentration range between 0.025 µg L⁻¹ and 1000 µg L⁻¹ with A=631.06 C – 0.0083 equation and R²=0.9998 correlation coefficient. In the equation, A and C are absorbance unit and RhB concentration, respectively. Preconcentration factor was obtained as 30. Relative standard deviations (%RSD) was obtained as %3.9 for the ten replicates measurements of 0.125 µg L⁻¹ Rhodamine B. The method analytical characteristics were comparatively given in Table 3.
Table 3. Comparison of the method analytical features with recent reported studies

| Method     | Instrument            | Sample                                      | LOD (µg L⁻¹) | LDR (µg L⁻¹) | RSD (%) | PF | Reference                           |
|------------|-----------------------|---------------------------------------------|--------------|--------------|---------|----|-------------------------------------|
| SPE        | UV-Vis                | Matches tips, water, dish washing liquid    | 3.94         | 20-300       | 1.16    | 26 | Nekoeinia et al., 2016             |
| SPE        | Fluorescence          | Matches tips, shampoo, eye shadow, pencil lead | 0.1          | 0.35-5.0     | 8.2     | 33 | Bagheri et al., 2013              |
| CPE        | UV-Vis                | Lipstick, water, anti-freeze, waste-water   | 0.70         | 50-5000      | 5.0     | 40 | Bişgin et al., 2018               |
| DLLME      | UV-Vis                | Black tea, red wine, chili powder           | 1.48         | 5-450        | 4.7     | 20 | Xiao et al., 2014                 |
| CPE        | UV-Vis                | Tap water, waste-water                      | 0.04         | 30-300       | 6.75    | 28 | Ghasemi and Kaykhaii, 2016        |
| DES-LPME   | UV-Vis                | Lipstick, cologne, river water, nail polish cleaner | 2.2          | -            | 2.3     | 25 | Yilmaz and Soylak, 2018           |
| CPE        | UV-Vis                | Soap, soft pastel, matches tips             | 1.3          | 5-550        | 2.40    | 8.5 | Pourreza et al., 2008             |
| VA-IL-DLLME| Fluorescence          | Lipstick, soap, anti-freeze, red pencil core, matches | 0.01         | 0.025-1000   | 3.9     | 30 | Present work                      |

SPE: Solid-phase extraction
CPE: Cloud point extraction
DLLME: Dispersive liquid-liquid micro-extraction
DES-LPME: Deep eutectic solvent liquid phase micro-extraction
VA-IL-DLLME: Vortex assisted ionic liquid-based dispersive liquid-liquid micro-extraction

CONCLUSIONS

In the present proposed study, VA-IL-DLLME method was developed for extraction and spectrofluorometric determination of RhB in real samples. VA-IL-DLLME offers high extraction efficiency within a short time. The developed method is fast, simple inexpensive and uses very little amounts of toxic organic solvents when compared with other preconcentration methods such as SPE. Obtained quantitative satisfactory recovery results with analyte addition technique proved the applicability of the method to real liquid and solid samples including lipsticks, matches, anti-freezes, red pencil core and liquid soap. Moreover, present method determines the RhB concentrations of samples without applying centrifuge. At the optimum conditions of the developed method, ionic liquid [C₄MIM][PF₆] extracted only RhB molecules as quantitative from aqueous sample solution in presence of other matrix dyes given in Table 1. This circumstance has provided the selectivity to the proposed method. Because of fluorescence feature of RhB and usage of spectrofluorometer in determination step, LOD value of the method was determined as too low because fluorescence detection is very sensitive and RhB has considerably high molar absorptivity coefficient. Moreover, excitation wavelengths and emission wavelengths are specific for each dye that has fluorescence characteristic. In addition, fluorescence determination provided selectivity to the developed method because RhB has the fluorescence characteristic among other matrix dyes that are given in Table 1. The used dyes in matrix study have not fluorescence characteristics therefore, any emission wavelengths, which could be interfered the fluorescence determination, were not observed by fluorescence excitation with using fluorescence spectrometer. Thus, only RhB molecules were determined applying both excitation and emission wavelength. In addition, differences between excitation and emission wavelength of RhB increased the selectivity.
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