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Hexafluorophosphate salts with tropine-type cations in the extraction of alkaloids with the same nucleus from radix physochlainae†

Bing Dong, Jie Tang, Alula Yonannes and Shun Yao *

Ionic liquids (ILs) have been widely used in the field of extraction of natural bioactive compounds because of their advantages compared to traditional organic solvents. In this study, the new 'like dissolves like' mode was designed and seven types of tropine-based ionic liquids were used to extract tropane alkaloids from radix physochlainae, and then the relationship between their performance and structures together with the effects of main extraction conditions were explored. It was found that 0.05 mol L⁻¹ [C₃tr][PF₆] aqueous solution had the ideal selectivity and high extraction efficiency of 95.1% at 75 °C when the extraction time was 55 min and the solid–liquid ratio was 1 : 35, which was superior to that of 85% ethanol–water and 0.1% hydrochloric acid–water. There was no decomposition and racemization of products occurring in the mixture solution when above extraction solvent was applied. In addition, the extraction behavior and mechanism using an ionic liquid aqueous solution was tentatively studied through thermodynamics experiments, near-infrared/infrared spectroscopy (NIR/IR), scanning electron microscopy (SEM), and thermogravimetric analysis (TG), and subsequent back-extraction could be efficiently used to further separate alkaloids and ILs. In the developed 'like dissolves like' mode, the extraction process of target alkaloids was found to be endothermic and spontaneous through the specific interaction between them and the solvent molecules with the same nucleus.

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1. Introduction

Radix physochlainae is the dried root of *Physochlainae infundibularis* Kuang (Solanaceae), which has been used for a very long time in China and has been officially listed in the Chinese Pharmacopoeia.¹ It has the unique curative effects of relieving coughs and asthma as a traditional herbal medicine. The tropane series of alkaloids in radix physochlainae mainly include five kinds of alkaloids (anisodamine, scopolamine, atropine, scopoline and aposcopolamine), which are widely used for the treatment of expectorant, antiasthmatic, antitussive and diseases of the central nervous system.² In a previous study, Peng and coworkers³ used 75% ethanol to extract total tropane alkaloids from flos daturae with final yield of 0.32% through a twice soaking extraction (75 h every time) and a yield of 0.38% through twice backflow extraction (2 h + 1.5 h). In another study, a microwave-assisted method was employed and 87% tropane alkaloids was extracted under the conditions if 1 : 10 solid–liquid ratio (w/v), 25 min, 730 W microwave power and 65 °C.⁴ Ultrasonic wave (40 Hz) assisted extraction was also adopted by researchers⁵ and the yield of 0.44% was finally

obtained when the extraction was carried out with 85% ethanol on 50 °C for 75 min. However, there are some problems including long extraction time, low efficiency and high consumption in the traditional reflux or immersing extraction. On the other hand, new technologies such as ultrasonic extraction and microwave extraction cannot be easily applied to large-scale production in industry because of the limitations of devices and high cost. More importantly, no matter what kind of these mentioned methods is used, traditional alcohol–water/acid–water solvent system will extract most of constituents in herbs without high selectivity, and the great amount of coexisting impurities can result in the difficulty of obtaining target compounds. Therefore, it is necessary to develop more new extraction methods that can promote extraction selectivity, efficiency and practicability meanwhile overcome the problems resulting from the use of traditional solvents.

'Like dissolves like' is a very classical expression frequently used by chemists to understand how some solvents work in the dissolution of solutes. It is generally recognized that this view is simplistic meanwhile it is a useful rule of thumb. The overall solvation capacity of a solvent depends primarily on its polarity. In the field of liquid-phase extraction and liquid–liquid partition, the rule of 'like dissolves like' generally refers to that target compounds tend to dissolve in the solvents with similar polarity. Guided by this rule, researchers can select a series of potential solvents for specific objects on the basis of

School of Chemical Engineering, Sichuan University, Chengdu, 610065, China. E-mail: cusack@scu.edu.cn; Fax: +86-028-85405221; Tel: +86-028-85405221

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accumulated experience and then compare their performance through further experiments. More researches are expected for their valuable contribution to develop new theories and explore deep mechanism, especially for the results when the target compounds meet the extractants with similar structures. Ionic liquids (ILs) is such a kind of designable solvents with variable structures, which have a lot of unique properties (*e.g.* high solubility, vapor pressure, and good stability, *etc.*) different from those of volatile organic solvents (VOS).⁶ They are attracting more and more attention from researchers in the fields of organic synthesis, catalysis and separation science.^{7–9} In recent years, ionic liquids have been widely used in the field of extraction and separation of natural products; they can competitively generate hydrogen bonds with cellulose for the maximum damage to the H-bonding network of cellulose microfibrils, which makes them more easily enter into the plant cell and improve the extraction efficiency. In addition, the solubility of ILs and their specific interactions with different objects can be regulated by adjusting the combination of anions and cations. Finally, the ionic liquid also can be applied in ultrasonic and microwave-assisted modes to shorten the extraction time and promote the extraction efficiency, which is very meaningful in the protection of natural products from destruction under severe conditions. For instance, Cao and his workers¹⁰ used a series of 1-alkyl-3-methylimidazolium ionic liquids to extract piperine from white pepper powders with ultrasonic wave, and they found that the increase of ILs hydrophilicity was beneficial for the improvement of extraction yield. Therefore the yield of 35.7 mg g⁻¹ was obtained by 2 mol L⁻¹ [BMIM][BF₄] after 30 min in 500 W ultrasonic wave, meanwhile only 19.60 mg g⁻¹ yield could be achieved by 75% methanol. Ma and his workers¹¹ developed microwave-assisted extraction to obtain three alkaloids from lotus leaves. The extraction efficiency of [BMIM][Br] was highest compared with other six kinds of ILs because its acidity was more conducive to generate salts with alkaloids, and the yield of three alkaloids by using [BMIM][Br] were all higher than that of 80% ethanol. The research group of Zu found that the extraction efficiency of ILs for three alkaloids from *Catharanthus roseus* increased when the alkyl chain length of the cation was changed from C₂ to C₈,¹² which was due to the hydrophobic effect between ionic liquids and alkaloids and it became the main driving force in the extraction process. In summary, most of these studies focused on the application of ionic liquids in existing extraction modes. Their structural designability is always underutilized and related advantages cannot be fully exploited. Moreover, the inner relationship between their unique physicochemical properties and extraction performance needs further exploration.

According to above research status, the rule of 'like dissolves like' can be simply interpreted as that the hydrophilic cations or anions are needed when the extracted alkaloids are low-polar. At a deeper level, since the formation of hydrogen bonds, dipole-dipole interaction, van der Waals' force and Coulomb force between the ionic liquid and alkaloids can strengthen the selective recognition ability of IL and promote their binding; those ILs with the same nucleus with alkaloids should be more

suitable and efficient in the extraction of the latter. This hypothesis is supposed to be the generalized 'like dissolves like' rule, which will be further evaluated and validated in the following study. Here radix physochlainae was selected as typical herbal object containing tropane alkaloids, and seven kinds of tropane-types ionic liquids were used to extract target alkaloids from it in this new mode. Accordingly, the major research objectives are focused on as followings: (1) what will happen when alkaloids meet the ILs with similar structure? (2) Will the "like dissolve like" rule work on the basis of structural similarity? (3) Can high efficiency and selectivity be obtained when the ILs interact with their analogues in such a complex composition from herbal sample? Related main properties of ionic liquids were first explored, and then the extraction conditions were investigated and the extraction mechanism using ionic liquid aqueous solution was studied. Obviously, this reinterpreted rule can maximize the ILs advantages of designability and selectivity, while it cannot be made full use by those common undesignable solvents.

2. Experiment

2.1 Materials

Herbal raw materials of radix physochlainae originating in Shaanxi province were obtained from local pharmacy and identified as the dried roots of the Solanaceae plant by Prof. Fangyan Li in our department. They were milled and dried and then the sample powders passed through a stainless steel sieve with the particle size of 60 mesh. All chemicals involved in this study were at least of analytical grade and purchased from Kelong Chemical reagents factory (Chengdu, China). Chromatographic methanol used for high performance liquid chromatography (HPLC) was purchased from Chengdu Chemical reagents factory and filtered with microporous membrane (0.45 μm) before use. Standard compounds of anisodamine, *D/L*-atropine, scopolamine, aposcopolamine and scopoline (see Fig. 1) used for HPLC were purchased from National Food and Drug Inspection Institute (Beijing, China) or Research Chemicals Inc. (Toronto, Canada) and all of their purities ≥98.0%. Experimental deionized water was obtained from ultra-pure water purification system (0.4 mm filter) manufactured by Millipore Co., Ltd. (Bedford, USA). Ionic liquids in Fig. 1 were synthesized and purified according to the previous method,¹³ and the purity (all above 97.9%) was analyzed by HPLC method with evaporative light-scattering detector. All samples were stored in closed desiccators until use.

2.2 Apparatus

Nicolet 6700 Fourier transform infrared spectrometer (Abbr. FT-IR, Thermo Scientific, Madison, USA), JSM-7001F scanning electron microscopy (Abbr. SEM, JEOL, Tokyo, Japan), STA 449 F3 simultaneous thermal analyzer (NETZSCH, Selb, Germany) were used to observe morphology of herbal powders and residue after extraction. NIR spectrometer (NIRQUEST512), deuterium light source (DH-2000-BAL) and fiber optic probes (T300-UV-VIS) were provided by Ocean Optics (Dunedin, USA), which



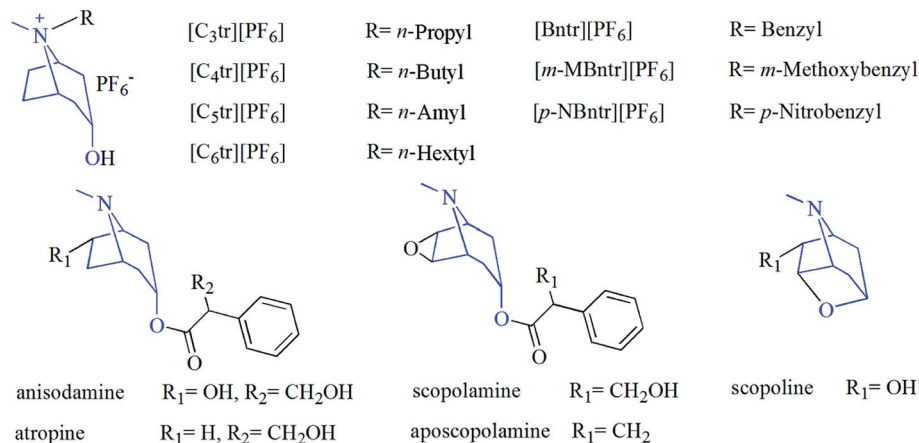


Fig. 1 Structures of tropane alkaloids and hexafluorophosphate salts with same nucleus.

was used to detect the extraction of alkaloids from radix physochlainae. Extraction experiments were carried out using HH-S constant temperature water bath with an uncertainty of ± 0.05 K (Wanke Instrumental Co. Ltd., Jiangsu, China) and KQ-2200DA ultrasonic extractor (Shumei Co. Ltd., Jiangsu, China). DDS-12A conductivity meter (Kangyi Scientific instrument Co. Ltd., Shanghai, China) and 2800 UV-Vis spectrometer (Hengping instrument Co. Ltd., Shanghai, China) were used to measure conductivity and polarity of ionic liquids. HPLC analysis was performed with EC2006 HPLC (Elite, Dalian, China) consisting of UV1201 UV-Vis detector and ELSD 2000 evaporative light-scattering detector (Alltech, Kentucky, USA). Refractive index of alkaloids extraction solution was determined by WAY-2S Abbe refractometer (Shengguang instrument Co., Ltd., Shanghai, China) to study thermodynamics and dynamics of alkaloids extracted from radix physochlainae.

2.3 Determination of ionic liquid polarity and conductivity

The polarity of ionic liquids was measured by Reichardt's dye method¹⁴ as following: 2.5 g ionic liquids were accurately weighed and dissolved in 100 mL of methanol, then 0.0375 g 2,6-diphenyl-4-(2,4,6-triphenyl-*N*-pyridino)phenolate was added into above IL methanol solution. When the dye molecule was completely dissolved and mixed with the ionic liquids, methanol in the mixture was completely removed under vacuum for 48 h, and then the dried product was analyzed by full wavelength scanning (200–800 nm). Conductivity was measured using DDS-12A digital conductivity meter with automatic balance bridge, and the conductance electrode constant was determined as 1.0474 cm^{-1} by standard aqueous solution potassium chloride (0.01 mol L^{-1}). Seven kinds of ionic liquids were measured from 5°C to 85°C with interval of 20°C under stirring to reach complete equilibrium. The electrode and conductivity cell with self-made double jacketed glass device were washed with distilled water, redistilled water and acetone respectively, and then dried by cold air before the measurement.

2.4 Extraction procedure

2.4.1 Extraction by ionic liquids aqueous solution. 1.00 g powders (60 meshes) of radix physochlainae were weighed accurately and placed in 50 mL Erlenmeyer flask, which was mixed with 10 mL 0.05 mol L^{-1} aqueous solution of hexafluorophosphate salt ($\text{pH} = 5.3$). The target constituents were extracted at 70°C in constant temperature water bath for a certain time. At the end of extraction, the solution was filtered and the filtrate was concentrated and its volume was set with methanol of chromatographic grade to 5 mL. Then the sample solution was filtered with a microporous membrane of $0.45 \mu\text{m}$ before HPLC analysis.

2.4.2 Extraction by traditional solvents. Four common solvents were selected to extract the target alkaloids from the same powders (60 meshes) of radix physochlainae with comparison to ionic liquids, which included pure deionized water (1), the system of 0.1% hydrochloric acid and water (2), the mixture of 85% ethanol and water (3), and the system of 85% ethanol and 0.1% hydrochloric acid with water (4). The extraction was carried out with solid-liquid ratio (g mL^{-1}) of 1 : 10 at 70°C for 50 min; after extraction, the solution was filtered and the filtrate was evaporated and then diluted with methanol of chromatographic grade to 10 mL. Various samples were filtered with $0.45 \mu\text{m}$ microporous membrane and the content of total alkaloids was analyzed by HPLC to calculate extraction efficiency (%).

2.5 Analytical method

2.5.1 HPLC conditions and standard curves of five alkaloids. HPLC analysis based on the reported method¹⁵ for five representative alkaloids was carried out on Welch C_{18} chromatographic column ($4.6 \times 250 \text{ mm}$, $5 \mu\text{m}$) at column temperature of 25°C ; the mobile phase was composed of 20 mmol L^{-1} ammonium acetate solution (A) and methanol (B) and the flow rate was 1.0 mL min^{-1} . Detection wavelength was set at 210 nm and injection volume was $5 \mu\text{L}$, and the gradient procedure was operated as: 0–10 min, A : B = 60 : 40 (v/v); 10–30 min, A : B = 30 : 70 (v/v); after 30 min, A : B = 20 : 80 (v/v).



The concentration of five standard alkaloids had good linear relationship with peak area in the investigated range of content, and corresponding standard curves of anisodamine, atropine, scopolamine, aposcopolamine and scopoline were determined as $y_1 = 12\,261.40x_1 - 368.19$ ($R^2 = 0.9998$, linear range: 0.50–3.50 μg , 95% confidence interval of slope and intercept: ± 152.80 & 4.26), $y_2 = 3560.60x_2 - 619.55$ ($R^2 = 0.9997$, linear range: 0.24–1.28 μg , 95% confidence interval of slope and intercept: ± 38.21 & 8.08), $y_3 = 1628.80x_3 - 135.65$ ($R^2 = 0.9999$, linear range: 0.17–1.73 μg , 95% confidence interval of slope and intercept: ± 17.98 & 2.53), $y_4 = 2188.80x_4 - 266.52$ ($R^2 = 0.9997$, linear range: 0.15–0.40 μg , 95% confidence interval of slope and intercept: ± 23.17 & 4.32), $y_5 = 110.59x_5 - 1.84$ ($R^2 = 0.9998$, linear range: 0.20–1.20 μg , 95% confidence interval of slope and intercept: ± 1.98 & 0.09) successively, where x was their amount (μg) and y was the UV absorbance value of related peak area, respectively.

2.5.2 Quantitative analysis of tropane alkaloids in herbal material. 1.00 g powders (60 mesh) of radix physochlainae were weighed accurately and pretreated with ammonia, and then were extracted by chloroform with solid–liquid ratio of 1 : 10 (g mL^{-1}) for 3 times.¹⁶ The extract was concentrated and its volume was set with methanol to 10 mL, and 20 μL sample solution was injected into HPLC after filtration by 0.45 μm microporous membrane. As the result, the chromatogram of total extract from radix physochlainae indicated that five alkaloids could be separated well with other coexisting compounds (see Fig. 2a); their amount could be determined according to the peak area and standard curves introduced in the last section. It showed that anisodamine (0.632 mg g^{-1}) and scopoline (0.509 mg g^{-1}) are the first and second main alkaloids in our sample respectively, meanwhile the content of aposcopolamine was only 0.093 mg g^{-1} (shown in Fig. 2b) and total mass fraction of five alkaloids was 0.173% in herbal material. The amount of target constituents (m_1 , mg) in extraction solution was obtained by the developed HPLC method and used to calculate the extraction efficiency (%), which could be determined through the following equation:

$$\text{Extraction efficiency } (E, \%) = (m_1/m_0) \times 100 \quad (1)$$

where m_0 (mg) is the total alkaloids amount in herbal powders determined as above.

2.6 Mathematical tools

2.6.1 First-order model. A general first-order model has been used to study the kinetics of solid–liquid extraction, which can provide an acceptable representation of solid–liquid extraction as evident.^{17,18} The parameters in the first order kinetic model can be easily elucidated from the extraction kinetics, using the analytical solution obtained by integral analysis. The same model has been extended to study the extraction of alkaloids from radix physochlainae by ionic liquids aqueous solution and other traditional organic solvents. The first order kinetic model can be represented as:

$$\frac{dC_t}{dt} = \kappa(C_s - C_t) \quad (2)$$

where C_t (mg mL^{-1}) is the concentration of alkaloids at any time (t), C_s (mg mL^{-1}) is the saturation concentration of alkaloids and k (min^{-1}) is the first order rate constant. Then eqn (2) is solved with the boundary conditions as $C_t|_{t=0} = 0$ and $C_t|_{t=t} = C_t$ to be:

$$\ln(C_s - C_t) = \kappa t + A \quad (3)$$

The effect of temperature on extraction rate constant provides information on the activation energy that must be overcome for extraction. The extraction rate constant is related to the activation energy (E_a) and temperature (T) as following:

$$\kappa = \kappa_0 \times e^{\frac{-E_a}{RT}} \quad (4)$$

where a plot of $\ln(\kappa)$ vs. $1/T$ yields a straight line with (E_a/R) as the slope, from which activation energy can be determined.

2.6.2 Effective diffusion coefficient of extraction of alkaloids from radix physochlainae. To study the effective diffusion coefficient of alkaloids from radix physochlainae, the diffusion

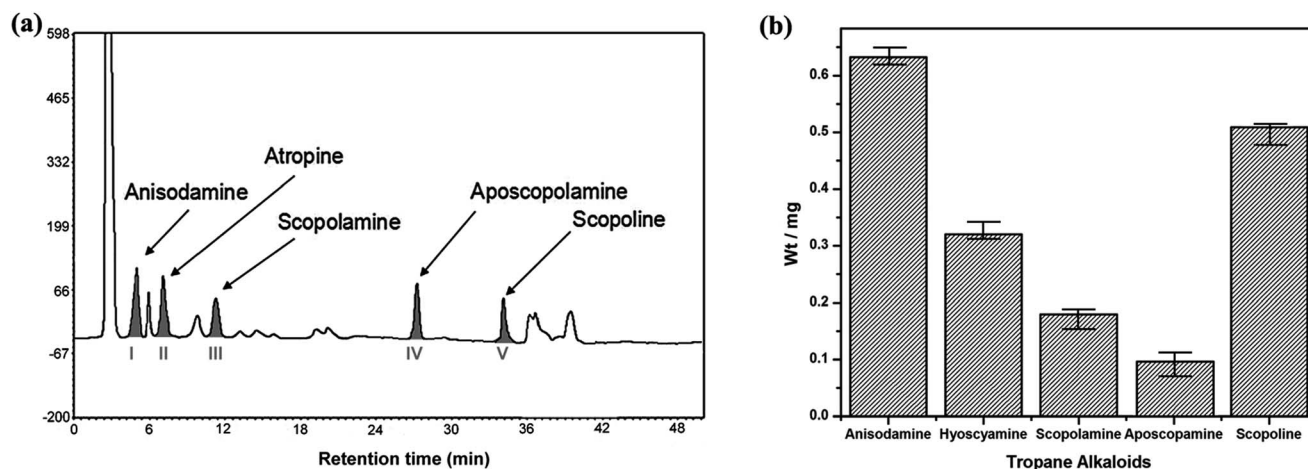


Fig. 2 The chromatogram (a) and composition (b) of five tropane alkaloids in radix physochlainae.



model based on Fick's second law was used in this study. The law for extraction of alkaloids from spherical particles is expressed as following:¹⁹

$$\frac{\partial M}{\partial t} = D_e \left[\frac{1}{r^2} \frac{\partial}{\partial r} \left(r^2 \frac{\partial M}{\partial r} \right) \right] \quad (5)$$

where M (kg m^{-3}) is the alkaloids concentration in radix physochlainae particles, r (m) is the distance from the center of herbal particles, D_e is the effective diffusion coefficient of alkaloids and t (min) is time. According to the ref. 20, the solution of eqn (5) using the initial and boundary conditions can be transformed to:

$$\frac{Y_t}{Y_s} = 1 - \frac{6}{\pi^2} \exp\left(-\frac{D_e \pi^2 t}{R^2}\right) \quad (6)$$

where Y_t (mg g^{-1}) is the yield of total alkaloids extracted from radix physochlainae at time (t), Y_s (mg g^{-1}) is the yield of total alkaloids extracted at saturation. The effective diffusion coefficient can be calculated from time-yield data using eqn (6).

2.6.3 Thermodynamics of extraction of alkaloids from radix physochlainae. Thermodynamic analysis of extraction of solute from solid using extraction has been carried out through determination of ΔG° , ΔH° and ΔS° . Van't Hoff equation²¹ can be used to determine ΔG° and ΔS° as following:

$$\ln K_e = -\frac{\Delta H^\circ}{RT} + \frac{\Delta S^\circ}{R} \quad (7)$$

where K_e is the equilibrium constant for leaching and is defined as the ratio of amount of extracted solute to that of unextracted solute.²¹ In the present case, it is the ratio of the amount of extracted alkaloids to the amount of unextracted alkaloids as given below:

$$K_e = \frac{Y_s}{Y_{\max} - Y_s} \quad (8)$$

where Y_s (mg g^{-1}) is the total alkaloid yield extracted using ionic liquid aqueous solution as the solvent at a temperature T (K) and Y_{\max} is the total alkaloid yield (mg g^{-1}) determined after an exhaustive extraction using chloroform as the solvent. ΔG° is calculated from ΔH° and ΔS° as eqn (9):²¹

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ \quad (9)$$

3. Results and discussion

3.1 Structure, properties and extraction performance of ILs for tropane alkaloids

The physical and chemical properties of ionic liquids can be significantly affected by the combination of cations and anions,²² so their extraction performance is very different for specific compounds. Generally, the structural features of target constituents and the special characteristics of ILs as polar organic salts should be fully considered before the extraction, and then the potential tropane-based ionic liquids are selected according to the generalized principle of 'like dissolves like' to achieve efficient and highly-selective extraction. Meanwhile,

considering the lower polarity of tropane alkaloids, $[\text{PF}_6]^-$ was chosen as the most commonly used hydrophobic anion. As the result, seven ionic liquids were used to extract alkaloids with the same nucleus and the comparison of extraction efficiency was performed under the same extraction conditions of 0.1 mol L^{-1} ionic liquids aqueous solution, 1 : 10 solid-liquid ratio (g mL^{-1}), 50 min and 70°C . As shown in Fig. 3a, it can be found that the order of extraction efficiency is same as that of association constant of the studied ionic liquids (see Table 1), and both of them is $[\text{C}_3\text{tr}][\text{PF}_6] > [\text{C}_4\text{tr}][\text{PF}_6] > [m\text{-MBntr}][\text{PF}_6] > [\text{C}_6\text{tr}][\text{PF}_6] > [\text{C}_5\text{tr}][\text{PF}_6] > [p\text{-NBntr}][\text{PF}_6] > [\text{Bntr}][\text{PF}_6]$. Obviously, one of the key factors for high extraction efficiency is the strong association action resulting from intermolecular interactions between ionic liquids and alkaloids, and there are two competitive effects between ionic liquids-water and ionic liquids-alkaloids in extraction system. If the solvation of water for ionic liquids is weaker, the extraction efficiency will be higher because of the stronger association effect of ionic liquids with alkaloids. Here the association constant provides an ideal prediction index for the extraction performance of various ILs.

At the same time, the polarity of ionic liquids is also one of the important properties that affect the extraction rate of alkaloids. So it was investigated through the method described in Section 2.3 and the data have been reported in our previous study.¹³ As the result, the polarity of four kinds of ionic liquids branched with linear carbon chain were found higher than that with aromatic hydrocarbons except for $[p\text{-NBntr}][\text{PF}_6]$,²³ which was due to that strong electron withdrawing group in $[p\text{-NBntr}][\text{PF}_6]$ could make the charge center of the molecule more unevenly distributed and the molecular polarity of $[p\text{-NBntr}][\text{PF}_6]$ was higher than that of $[\text{C}_n\text{tr}][\text{PF}_6]$ ($n = 5$ or 6). Because the nonpolar alkyl chain is shorter, the polarity of $[\text{C}_3\text{tr}][\text{PF}_6]$ and $[\text{C}_4\text{tr}][\text{PF}_6]$ is higher than that of $[\text{C}_5\text{tr}][\text{PF}_6]$ and $[\text{C}_6\text{tr}][\text{PF}_6]$, and their extraction performance was found higher than the latter two ILs.

In the further investigation for inherent relationship between ILs and individual alkaloids with same nucleus, the extraction rule of four kinds of ionic liquids branched with linear carbon chain ($\text{C}_3 \sim \text{C}_6$) for anisodamine and scopoline was firstly studied (shown in Fig. 3b), which had the first and second highest content among these alkaloids. It was found that extraction rate of anisodamine became higher with the increase of polarity of ionic liquids, while the trend for scopoline was exactly the opposite. From the viewpoint of the 'like dissolves like' extraction for single alkaloid compound, it could be found that they can be extracted more efficiently when their polarity is more similar to the polarity of ILs. Anisodamine has more polar groups in its structure and shorter retention time in reverse-phase system than scopoline, which can indicate that the polarity of anisodamine is higher than that of the latter; so the high extraction efficiency of anisodamine and scopoline can be achieved by $[\text{C}_3\text{tr}][\text{PF}_6]$ as the IL with highest polarity and $[\text{C}_6\text{tr}][\text{PF}_6]$ as the IL with lowest polarity, respectively. Moreover, it should be noticed that $[\text{C}_3\text{tr}][\text{PF}_6]$ had higher efficiency than $[\text{C}_4\text{tr}][\text{PF}_6]$ for the extraction of scopoline, and this result was also attributed to the structure of the IL cation. Alkyl chain with the carbon number <4 is more beneficial for intermolecular



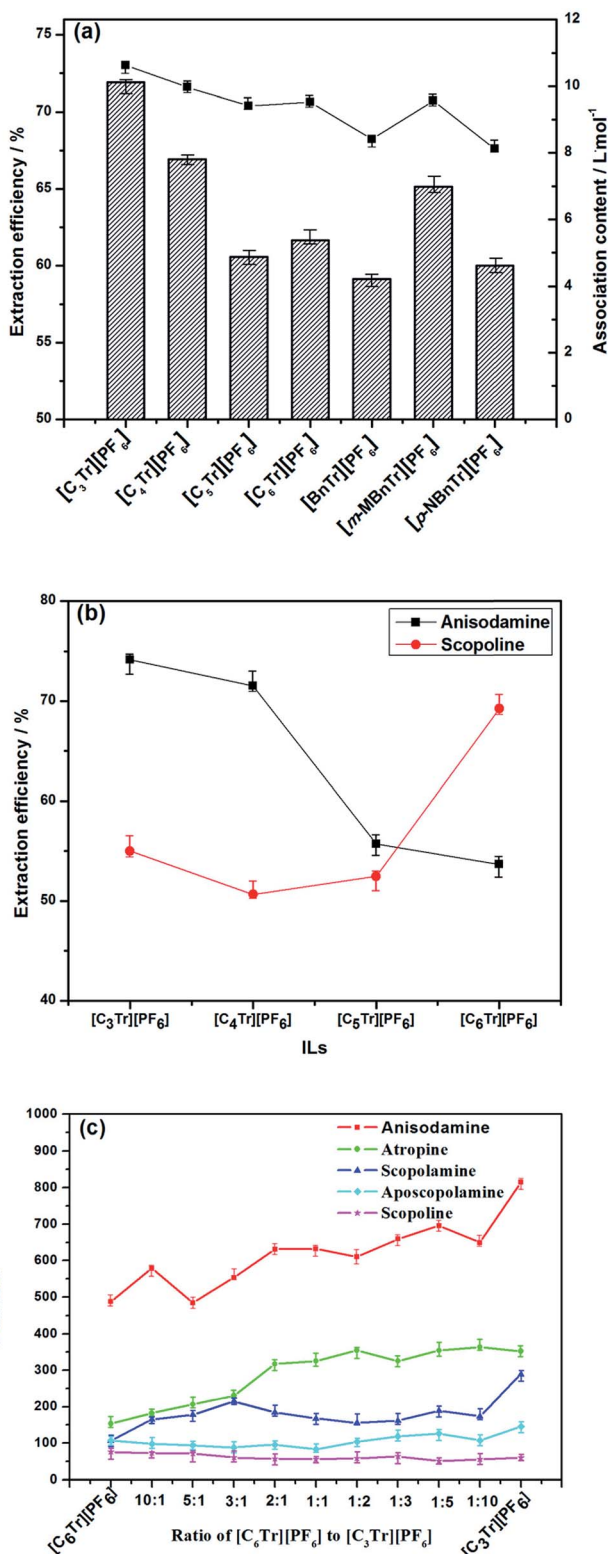


Fig. 3 The extraction efficiency of total alkaloids (a) and two main alkaloids (b) by ionic liquids aqueous solution and five tropane alkaloids using [C₃Tr][PF₆] + [C₆Tr][PF₆] + H₂O ternary system as extraction solvents (c).

interaction because of its small volume and hindrance, and related ILs have higher absolute value of Gibbs free energy of association (ΔG). When the carbon number is beyond 4, the polarity effect will dominate the extraction process. As for the mixture of target alkaloids with different polarity (just like their coexisting status in herbal cells), those with higher polarity can raise the polarity level of total alkaloids, especially when they have superior quantity and content. So [C₃Tr][PF₆] aqueous solution has shown the highest extraction efficiency in seven types of ionic liquids for raw material of radix physochlainae.

Finally, there is little investigation and application of more than one ionic liquid in similar process in previous studies. Since the most ideal IL for the extraction of various individual alkaloids is different, here the possible synergistic effect of two kinds of ILs mixed solution was explored and the combination of [C₃Tr][PF₆]-[C₆Tr][PF₆] was selected which possessed the highest and lowest polarity in the series of [C_nTr][PF₆] respectively. The experimental conditions are as follows: 0.1 mol L⁻¹ ionic liquids mixture solution, 1 : 10 solid-liquid ratio, 50 min on 70 °C for comparing single ionic liquids and IL mixture with the different ratios ([C₆Tr][PF₆] : [C₃Tr][PF₆] = 1 : 0, 10 : 1, 5 : 1, 3 : 1, 2 : 1, 1 : 1, 1 : 2, 1 : 3, 1 : 5, 1 : 10, 0 : 1, w/w). As shown by the results in Fig. 3c, it can be found that the content of anisodamine will become higher with the increase of [C₃Tr][PF₆] in the mixture, meanwhile the content of scopoline will decrease. On the other hand, the extraction performance of the mixture of [C₆Tr][PF₆] and [C₃Tr][PF₆] aqueous solution was always worse than that of [C₃Tr][PF₆] aqueous solution and no synergistic effect was found. This result could be ascribed to the unavoidable association action between [C₆Tr][PF₆] and [C₃Tr][PF₆], which can result in their nonideal mixing and nonlinear variation of physico-chemical properties.²⁴⁻²⁶ It was not conducive to their interaction with alkaloid molecules and improvement of extraction efficiency. Therefore, the following studies were carried out with [C₃Tr][PF₆] as the extraction solvent.

3.2 Effects of various conditions on the extraction of tropane alkaloids

Concentration of ionic liquid solution has great influence on the swelling and destruction of cell wall together with mass transfer process of extraction solution, which can directly determine the extraction efficiency of extractant on the target alkaloids. In the following investigation, 10 mL of 0.05 mol L⁻¹ ~ 0.175 mol L⁻¹ [C₃Tr][PF₆] aqueous solutions were compared in the extraction of 1.0 g herbal powders for 50 min at 70 °C. As shown in Fig. 4a, the extraction efficiency firstly increases and then gradually decreases with the increase of IL concentration. It is due to that with the continuous increase of concentration of ionic liquid, over high viscosity of extractant will hinder radix physochlainae powders to well disperse in the ionic liquid solution for full extraction and the permeation of the latter into plant cells together with mass transfer become more difficult.²⁷

Extraction time is another important factor needed investigation and optimization. After reaching complete extraction, excessive time will be unnecessary and unfavorable due to the instability of ester bond or other bond in the structures of



Table 1 Association constant (K_A), limiting molar conductivity (Λ^∞) and Gibbs free energy of IL association (ΔG)

$T/^\circ\text{C}$	$[\text{C}_3\text{tr}][\text{PF}_6]$			$[\text{C}_4\text{tr}][\text{PF}_6]$			$[\text{C}_5\text{tr}][\text{PF}_6]$		
	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$
5	61.36	5.01	-3212	56.18	4.78	-3618	51.55	4.01	-2864
25	88.03	6.51	-4253	85.96	6.25	-4491	84.24	6.12	-3847
45	129.87	8.15	-5312	125.00	8.13	-5543	121.95	7.55	-5549
65	169.49	10.63	-6645	151.52	9.97	-6302	139.31	9.41	-6465
85	263.16	15.83	-8224	250.01	14.87	-6964	240.9	12.81	-8038

$T/^\circ\text{C}$	$[\text{C}_6\text{tr}][\text{PF}_6]$			$[\text{Bntr}][\text{PF}_6]$			$[m\text{-MBntr}][\text{PF}_6]$		
	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$
5	50.51	3.45	-3727	46.73	3.45	-2864	48.78	4.54	-3499
25	82.64	5.62	-4503	78.13	4.46	-3706	81.97	5.32	-4323
45	118.15	7.45	-5347	118.91	6.08	-4774	128.23	7.92	-5474
65	125.31	9.52	-6335	161.29	8.14	-6127	164.67	9.56	-6627
85	232.22	10.37	-7594	208.33	11.5	-7272	204.08	12.85	-7603

$T/^\circ\text{C}$	$[p\text{-NBntr}][\text{PF}_6]$		
	$\Lambda^\infty/\text{S cm}^2 \text{ mol}^{-1}$	$K_A/\text{L mol}^{-1}$	$\Delta G/\text{J mol}^{-1}$
5	47.40	5.14	-3786
25	80.65	5.44	-4244
45	120.77	6.79	-5067
65	161.29	8.12	-5888
85	208.33	9.11	-6579

tropine alkaloids. In order to avoid possible decomposition and racemization of products in the solution, 1.00 g herbal powders were extracted with 10 mL of 0.05 mol L⁻¹ [C₃tr][PF₆] aqueous solutions from 10 to 80 min. It was found the complete extraction could be achieved within 50 min by heating in water bath (see Fig. 4b). The fast-growing stage of extraction efficiency was the first 20 min; after 50 min, it gradually increased and extraction process approached equilibrium state. In order to further analyze related stages, kinetics research will be performed in Section 3.5.1. Compared to other alkaloids extraction by water,²⁸ it took less time for above ionic liquid aqueous solution to extract alkaloids from natural plants. Structural similarity makes them extract tropane alkaloids more easily from radix physoclainae in 'like dissolves like' mode.

Besides the concentrations of ionic liquids and extraction time, the solid-liquid ratio of the extraction system was also studied. Excessive ionic liquid will result in the increase of solvent consumption, and then increase the burden of IL recovery. Different ratios of liquid to solid (40, 30, 20 and 10 mL g⁻¹) have been investigated and the extraction was carried out with 0.05 mol L⁻¹ [C₃tr][PF₆] aqueous solution for 50 min on 70 °C (see Fig. 4c). The results indicated that the solid to liquid ratio grew fast in the range of 1 : 10–1 : 30; in the range of 1 : 30–1 : 40, extraction efficiency of tropane alkaloids was increased slowly and then unchanged. In the

fast stage of extraction, the significant concentration difference inside and outside the cells can promote mass transfer and diffusion effectively, the equilibrium concentration will become lower and more alkaloids can be extracted when the volume of solvent system is greater; if the excessive IL solution is used, heat will be absorbed by the large amount of solvent and could not be exerted onto the herbal material in the second stage.²⁸ The solid-liquid ratio of 1 : 30 is enough for complete extraction of tropane alkaloids in the radix physoclainae powders.

Finally, extraction temperature should be investigated because the thermal effect is the driving force when the object is completely dispersed into the extractive solution, which also determines the contact area and probabilities between the ionic liquid and the target alkaloids. The extraction efficiency is shown in Fig. 4d through the comparative experiments under different temperature of 40, 50, 60, 70, 80 and 90 °C. It can be found that the increase of temperature is linearly proportional to the extraction amount of tropane alkaloids for incomplete dispersion in solution which can affect mass transfer at lower temperature in the range of 40–70 °C; but in the range of 70–90 °C, a linearly proportional decline appears because of product instability on high temperature.²⁹ In conclusion, the temperature around 70 °C should be more appropriate for the extraction of target compounds according to the data and figure.



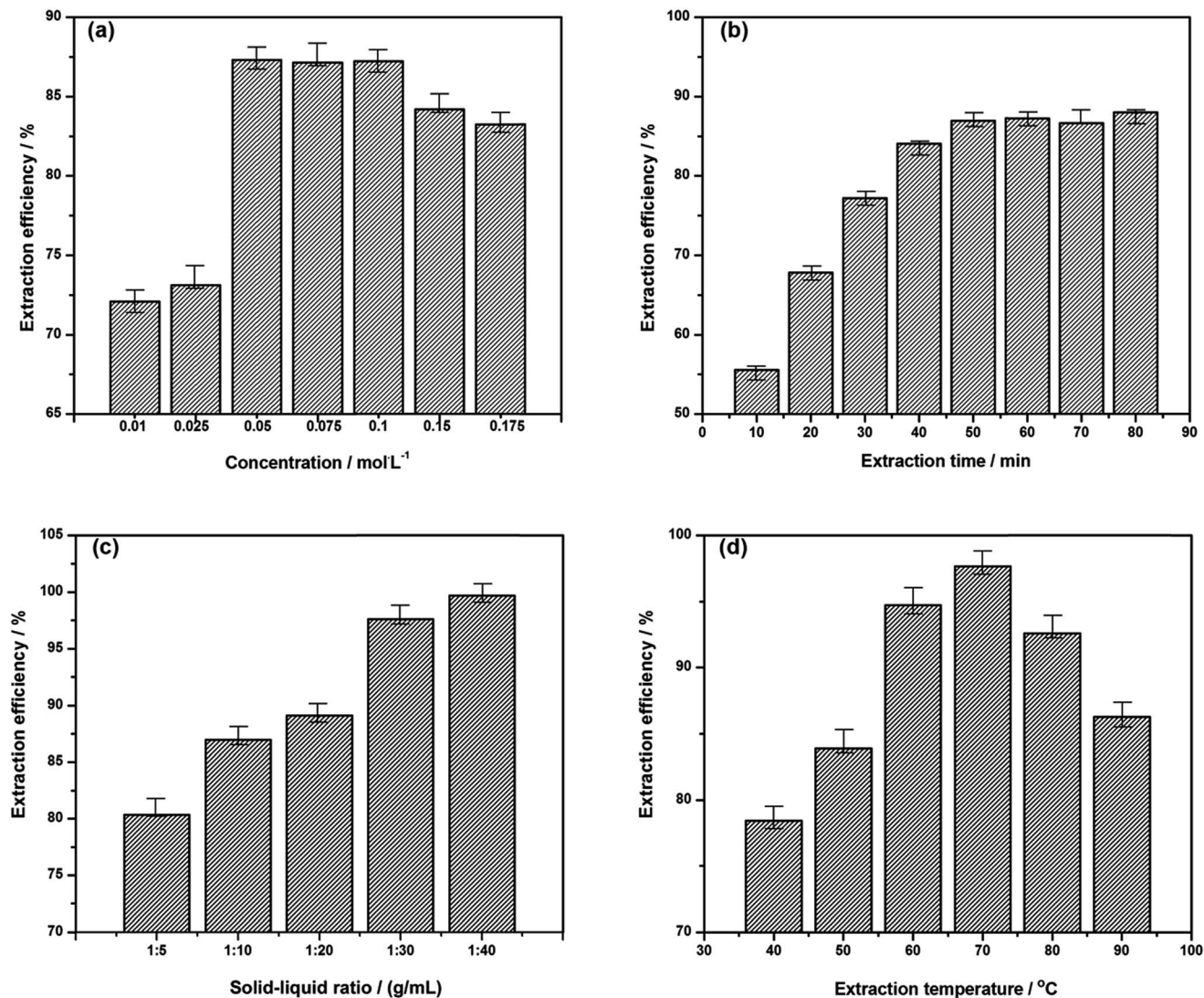


Fig. 4 Effects of various concentrations of ionic liquid (a), time (b), solid-liquid ratio (c) and temperature (d) on the extraction of total alkaloids.

3.3 Central composite design for interaction analysis and optimization

In order to further study the relationship among extraction factors and optimize extraction conditions, central composite design (CCD) was selected as the most commonly used method of response surface analysis.³⁰ The experimental design was completed by Design Expert 7.0 on the basis of optimal extraction conditions, which was also used to analyze the results through response surface. Three related parameters were included for selection as various independent variables (each variable representing three levels). There was no obvious difference in the yield of alkaloids when the concentration of [C₃tr][PF₆] from 0.05 to 0.1 mol L⁻¹, so extraction time, solid-liquid ratio and extraction temperature were selected which had more significant influence on the extraction efficiency. As the result, the regression second-degree complete model for the extraction efficiency (*Z*, %) in terms of coded factors is presented as the following equation:

$$Z = 93.80 + 3.54A + 1.16B + 0.56C + 0.27AB - 0.19AC + 0.14BC - 1.50A^2 - 2.54B^2 - 2.68C^2 \quad (10)$$

where *A*, *B* and *C* represent solid-liquid ratio, extraction temperature and extraction time, respectively. The analysis results of response surface method are shown in ESI 1† (RSM optimization). The significance of each variable to the response value was judged by *F* and *P* values. It can be found that the response surface model is highly significant (*F* = 48.29 and *P* < 0.0001); at the same time, the lack of *F* value of the model is 1.78, which proves that the result is not significant and no missing factor in the regression equation. In addition, *A*, *B*, *C*, *A*², *B*² and *C*² are significant items. The analysis results of response surface method can prove the feasibility of the regression second-degree complete model. Furthermore, the credibility analysis of the regression equations also indicates the model can be used to predict the extraction results effectively. The coefficient of variation (C.V.) of the model is 0.73%, which is in the acceptable range and shows good repeatability.



In addition, the “Adj *R*-Squared” of 0.9638 and R^2 of 0.9841 indicate that the model can explain the change of 96.38% response value, and the fitting of the model can better reflect the actual situation. In summary, the optimal extraction efficiency of 95.1% can be obtained, the optimal extraction temperature was 75 °C, the extraction time was 55 min and the best solid–liquid ratio was 1 : 35; meanwhile the final purity of total alkaloid reached 92.8% under above optimized conditions, which proved the highly selective extraction of IL for ‘like dissolves like’ mechanism. Moreover, the influential order of the three variables is: solid–liquid ratio ($P < 0.0001$) > extraction temperature ($P = 0.0016$) > extraction time ($P = 0.0479$, shown as 3D plots in ESI †), which can prove that the solid–liquid ratio plays a very crucial role in extraction.

3.4 Comparison with traditional methods and recovery of alkaloids

After exploring the optimal conditions, the extraction efficiency of alkaloids resulting from the developed method was compared with corresponding performance of other common solvents from previous research³¹ and Chinese Pharmacopoeia. In the following study, the system of 0.1% hydrochloric acid–water, 85% ethanol–water, and the system of 85% ethanol–0.1% hydrochloric acid–water were compared with 0.05 mol L⁻¹ [C₃Tr][PF₆] aqueous solution. Because imidazolium ionic liquids with the same anion of [PF₆]⁻ are hydrophobic and cannot be dissolved in water well, their performance was not compared with above systems. Instead, 0.05 mol L⁻¹ [Bmim][BF₄] was also selected in the list of solvents as a frequently-used imidazolium ionic liquid in extraction to validate the generalized ‘like dissolves like’ rule here. The traditional extraction was carried out with solid–liquid ratio of 1 : 10 on 70 °C for 50 min. It should be noted that microwave or ultrasonic wave-assisted extraction was not considered in the study because they were found unfavourable for the selective extraction; although these methods can enhance mass transfer and shorten extraction time at some extent, they will lower the purity of products significantly (<82%) according to our experiments. Even if 40 Hz ultrasonic wave was used in the extraction of tropane alkaloids, 75 min was still needed to extract them completely in previous study.⁵ On the basis of the comparison under the same conditions (see Fig. 5a), the extraction efficiency of tropane ionic liquid aqueous solution is highest; water shows the worse extraction performance for the investigated hydrophobic alkaloids, but a great difference can be observed after adding a little [C₃Tr][PF₆] into water. However, when we used [Bmim][BF₄] instead of [C₃Tr][PF₆], it was found the former could extract the alkaloids with higher polarity meanwhile those with lower polarity could not be extracted well, and total extraction efficiency became lower. Moreover, though ethanol also can improve the solubility of related objects in extraction system, its high concentration must be required and the extraction efficiency of [C₃tr][PF₆]-water still cannot be transcended by it. Meanwhile, the purity of total alkaloids extracted by ethanol was only 49.2%, while the selective interaction from tropine-based IL makes the product purity reach 92.8% in new ‘like

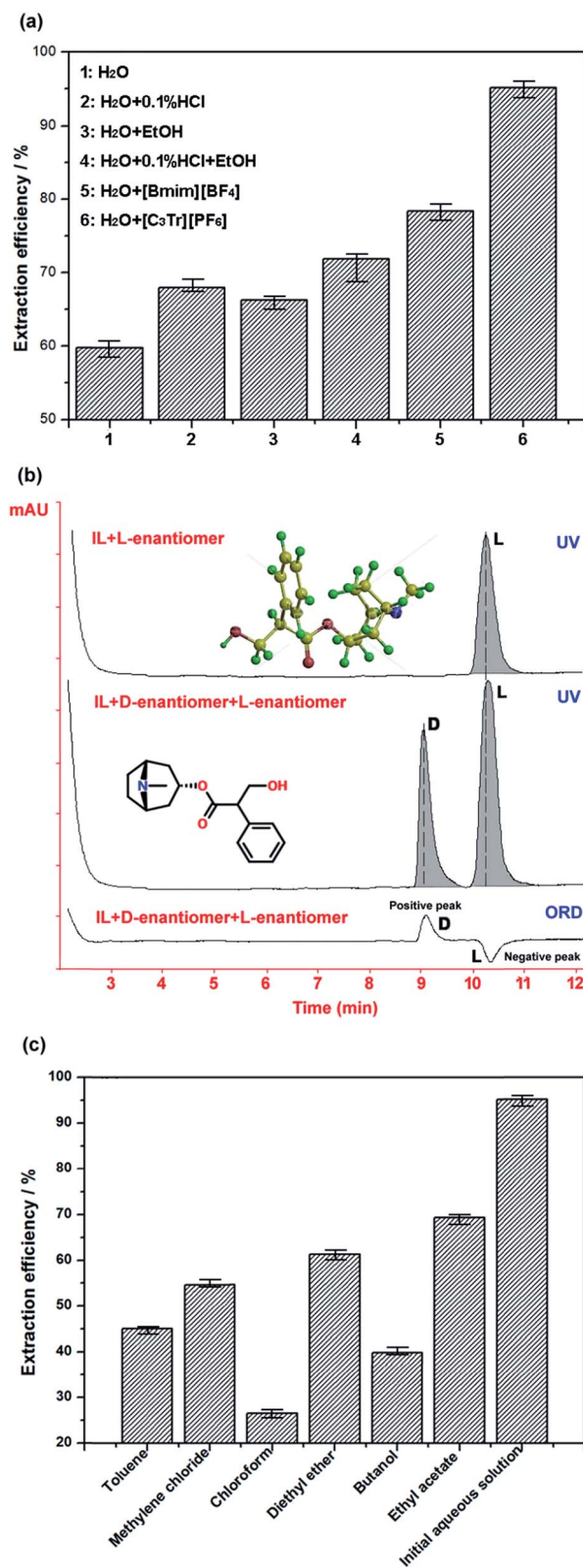


Fig. 5 Performance comparison of different extraction solvents on extraction efficiency (a), the chromatograms in racemization study by using chiral HPLC–UV/ORD (b) and the extraction efficiency of various solvents in recovery of tropane alkaloids (c).



Table 2 Comparison of specific rotation with IL aqueous solution, scopolamine aqueous solution and scopolamine-IL aqueous solution

Temperature/°C	$[\alpha]_{\lambda=598}^t$		
	IL aqueous solution	Scopolamine aqueous solution	Scopolamine-IL aqueous solution
25	0.18	29.00	28.00
35	0.18	22.00	22.00
70	0.06	16.00	13.00

dissolves like' mode. The extraction efficiency of 0.1% hydrochloric acid-water is not very ideal because the basicity of tropane alkaloids is not so strong that they cannot easily become water-soluble salt and fully extracted by this acidic aqueous solution. $[C_3tr][PF_6]$ belongs to near-neutral IL, which will not rely on salifying process to realize extraction. Furthermore, though the unit price of ionic liquids is a little higher, its dosage is very small and can be efficiently reused after efficient recovery through the suitable method.

In addition, because target tropane alkaloids usually contain chiral center in their structures, they are optically active (*e.g.* atropine existing in natural plants with its *L*-configuration). The ester bond in their structures cannot be easily affected under weakly acidic and near-neutral conditions, and it is the most stable in the pH range of 3.5–4. However, in the process of traditional extraction, it is difficult to obtain their original configuration in the raw materials because the alkaline solution or the long extraction time will cause the racemization of target alkaloids.^{32,33} Alkaloids are often found in the form of their salts in herbs for their binding with coexisting acids. In the pre-treatment process in Section 2.5.2, ammonia was frequently used to transform alkaloids from their ionic state to molecular state for ideal extraction result. How to eliminate influence of related conditions on the structure and configuration of target alkaloids was focused on by researchers in the past time. Considering the tropane-based ionic liquids were employed in the new developed extraction mentioned above, the off-line Autopol IV polarimeter (Rudolph Research Analytical, Hackettstown, USA) and on-line LC2000P optical detector (JASCO, Tokyo, Japan) coupled with HPLC were applied to observe the possible racemization phenomenon in the mixture of atropine-0.05 mol L⁻¹ $[C_3tr][PF_6]$ aqueous solution on 70 °C for 50 min. The chromatographic analysis was performed on Chromtech Chiral-AGP column (150 mm × 2 mm I.D., 5 μm, Congleton, UK) at room temperature and mobile phase was composed of 1% acetonitrile-10 mmol aqueous ammonium acetate buffer (pH 5.8); UV detection wavelength was 230 nm and flow rate was controlled at 0.3 mL min⁻¹. Under these conditions, the samples of the mixture of *D,L*-atropine standard compounds and IL (as sample 1) together with the mixture of IL and *L*-atropine prepared above (as sample 2) were analyzed under the UV and ORD detection. As the result, no *D*-enantiomer appeared in the mixture after the *L*-atropine was mixed with $[C_3tr][PF_6]$ under extraction conditions (see Fig. 5b) and related off-line optical data in Table 2 also proved there was no racemization occurred

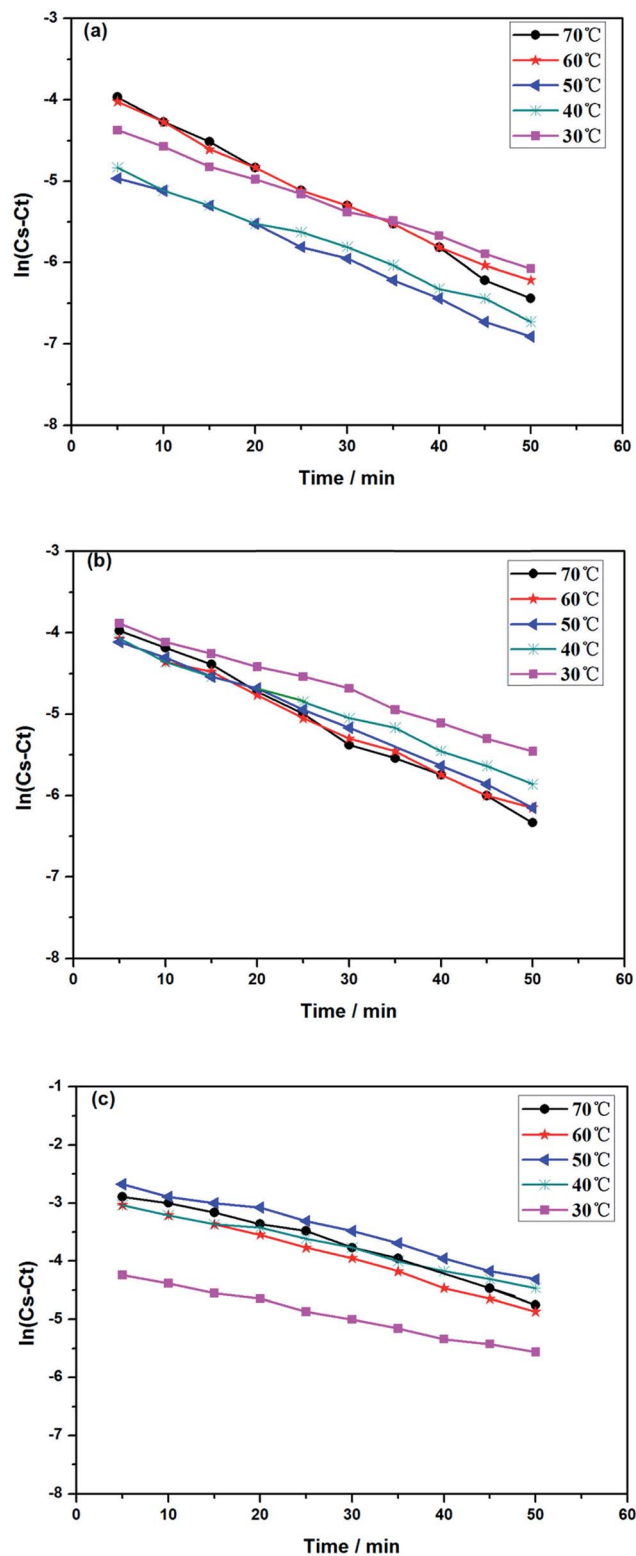


Fig. 6 Relationship between $\ln(C_s - C_t)$ and t through different extraction solvents ((a) 0.05 mol L⁻¹ $[C_3tr][PF_6]$ aqueous solution, (b) 85% ethanol-water, (c) 0.1% hydrochloric acid-water).

in the solution. By comparison, the alkaline condition could result in the decomposition of ester bond and transformation of atropine configuration indeed. In summary, the developed



method can not only realize the selective extraction efficiently, but also maintain the structure and configuration of products through specific intermolecular interactions.

Finally, considering the simplicity and efficiency of the method, the back extraction with common organic solvents (toluene, chloroform, dichloromethane, ethyl ether, *n*-butyl alcohol and ethyl acetate) was used to separate alkaloids from the ionic liquid and realize their recovery after the removal of water. These candidate solvents were mixed with the solid extract with solid-liquid ratio of 1 : 3 (g mL⁻¹) under stirring, respectively; then the organic solvent containing alkaloids was obtained by filtration and then concentrated under vacuum. Finally, the back-extracted products were dissolved in chromatographic methanol and analyzed by HPLC. The quantitative results in Fig. 5c indicate that the order of different solvents for alkaloids extraction efficiency is: chloroform > *n*-butyl alcohol > dichloromethane > toluene > ethyl ether > ethyl acetate. So chloroform was the best selection to recover alkaloids from their mixture with ILs. It should be noted that, although VOS was used in above post-treatment process of extraction, its consumption was much less for the back extraction of solid product than that in back extraction of extracted solution or the direct extraction of herbal materials with chloroform (*e.g.* the authoritative method in China Pharmacopoeia, solid-liquid ratio of radix physochlainae and chloroform is 1 : 20, g mL⁻¹), and here 1 : 3 was the ideal solid-liquid ratio of the solid extract and chloroform.

3.5 Extraction mechanism and process analysis

3.5.1 Kinetics of alkaloids extracted from radix physochlainae. The extraction rate constant (κ) and kinetic order for heating extraction of alkaloids from radix physochlainae by ionic liquid aqueous solution and other organic solvents can be identified through the study of extraction kinetics. In this study, Abbe refractometer was used to measure the dynamic concentration of alkaloids in radix physochlainae according to the linear relationship between the refractive index and the solid concentration in the solution within the test range.³⁴ Target

alkaloids in 1.00 g of radix physochlainae powders with 60 mesh was extracted with 35 mL 0.05 mol L⁻¹ [C₃tr][PF₆] aqueous solution, 85% ethanol-water and 0.1% hydrochloric acid-water, and the refractive index of samples was determined every five minutes within 1 h. It is found that $\ln(C_s - C_t)$ is linear with time (t , min) from Fig. 6 and Table 3. The results can indicate that the extraction kinetic equation derived from eqn (2) is in good agreement with the experimental results, and the dissolution process of alkaloids from radix physochlainae accords with the first-order kinetic model. The first-order kinetic model is obtained based on Fick's second law for linear diffusion when the $C_{t=0}$ of alkaloids is 0, which is suitable for component extraction without decomposition and volatilization and the herbal particles with regular spheres. Zhao and his coworkers³⁵ studied the extraction process of ester-series alkaloids from aconiti kusnezoffii Reichb in water bath and found that the kinetic equation could be well fit by the first-order kinetic model. In the study, extraction rate constant increased with the increase of temperature, and the results accorded with Fick's second law which showed that higher temperature resulted in the stronger penetration ability of solvents together with diffusion ability of those involved alkaloids. Similarly, the extraction rate constant (κ) can also become greater with the increase of temperature when using [C₃tr][PF₆] aqueous solution, 85% ethanol-water and 0.1% hydrochloric acid-water as extraction solvents, which can prove that high temperature is conducive to extract alkaloids from radix physochlainae. The order of extraction rate constant (κ) with three kinds of solvents is: [C₃tr][PF₆] aqueous solution > 85% ethanol-water > 0.1% hydrochloric acid-water, because [C₃tr][PF₆] can enter cells and interact with alkaloids with the same nucleus from radix physochlainae more easily than the others.

Furthermore, the fitting equation of $\ln \kappa$ and $1/T$ had good linear relationship in the extraction of alkaloids, and the average activation energy (E_a) was calculated according to the linear equation in different solvents. The results were summarized in Table 3, the average activation energy (E_a) of [C₃tr][PF₆] aqueous solution was least which could prove that it was more easy to extract alkaloids from radix physochlainae, and the

Table 3 Regression equation of $\ln(C_s - C_t)$ and t under different extraction solvents

Extraction solvents	Temperature/K	Fitting equation	R^2	k/min^{-1}	$E_a/\text{kJ mol}^{-1}$
[C ₃ tr][PF ₆] aqueous solution	303.15	$\ln(C_s - C_t) = -0.0371t - 4.2214$	0.9958	0.0371	8.2178
	313.15	$\ln(C_s - C_t) = -0.0402t - 4.6686$	0.9938	0.0402	
	323.15	$\ln(C_s - C_t) = -0.0445t - 4.6732$	0.9964	0.0445	
	333.15	$\ln(C_s - C_t) = -0.0490t - 3.8264$	0.9967	0.0490	
	343.15	$\ln(C_s - C_t) = -0.0541t - 3.7104$	0.9963	0.0541	
85% ethanol-water	303.15	$\ln(C_s - C_t) = -0.0346t - 3.7200$	0.9956	0.0346	9.1792
	313.15	$\ln(C_s - C_t) = -0.0380t - 3.9244$	0.9943	0.0380	
	323.15	$\ln(C_s - C_t) = -0.0455t - 3.8419$	0.9974	0.0455	
	333.15	$\ln(C_s - C_t) = -0.0471t - 3.8425$	0.9965	0.0471	
	343.15	$\ln(C_s - C_t) = -0.0528t - 3.6729$	0.9952	0.0528	
0.1% hydrochloric acid-water	303.15	$\ln(C_s - C_t) = -0.0300t - 4.0918$	0.9963	0.0300	8.9923
	313.15	$\ln(C_s - C_t) = -0.0321t - 2.8549$	0.9934	0.0321	
	323.15	$\ln(C_s - C_t) = -0.0370t - 2.4380$	0.9879	0.0370	
	333.15	$\ln(C_s - C_t) = -0.0413t - 2.7675$	0.9954	0.0413	
	343.15	$\ln(C_s - C_t) = -0.0445t - 2.5018$	0.992	0.0445	



average activation energy (E_a) was also lower than those reported values (e.g. 22.33 kJ mol⁻¹ in previous study³⁵). Meanwhile extraction rate constant of alkaloids extraction from radix physochlainae did not significantly change with temperature, so the temperature could not have important influence on the extraction efficiency of alkaloids according to the influential order of three variables (solid-liquid ratio, extraction temperature and time). The average activation energy (E_a) values of alkaloids extraction from radix physochlainae were below 20 kJ mol⁻¹, indicating that this extraction process was controlled by diffusion. According to the previous study for the fitted E_a values of the solid-liquid extraction of bioactive compounds from natural products, when the activation energy value lay in the range between 20 and 40 kJ mol⁻¹, indicating that the extraction process from herbal materials was governed by a mixed reaction and diffusion regime; and if the value was higher than 40 kJ mol⁻¹, the process was controlled by the solubilization reaction.³⁶

3.5.2 Effective diffusion coefficient. Effective diffusion coefficient (D_e) can reflect the diffusion ability of substances and depends mainly on the type of diffusion material and medium together with temperature and pressure. According to above study, the alkaloids concentration changed with time and corresponding molecules diffused in an unstable state; accordingly; the diffusion behavior of related alkaloids accorded with Fick's second law. Furthermore, the diffusion coefficient (D_e) between liquid and solid phases was calculated according to eqn (5) by experimental study, and it can be observed from Table 4 that effective diffusion coefficient will increase with the rising temperature and has good linear relationship with temperature. This trend should be due to the mechanism that the infiltration, diffusion, dissolution of alkaloids molecules could be accelerated when the temperature became higher. On the other hand, extraction rate is proportional to the extraction temperature, the resistance of alkaloids delivering through the solid-liquid interface will be lowered when the solution viscosity reduces with the rise of temperature, which is the key to promote to extract alkaloids from radix physochlainae and improve the mass-transferring rate of

alkaloids. It was found that the diffusion coefficient (D_e) of extraction by ionic liquid was higher than that in traditional solvents at every temperature, which proved that the extraction efficiency of alkaloids by ionic liquid was higher than 85% ethanol-water and 0.1% hydrochloric acid-water. The result was supported by the theory of 'like dissolves like' rule, and the tropane alkaloids are hydrophobic and easily dissolved because of selective interactions provided by the IL with similar structure. Moreover, it was easier to transfer alkaloids by using 85% ethanol-water than 0.1% hydrochloric acid-water for the tropane alkaloids were prone to dissolve in organic solvents. Consequently, the diffusion coefficient (D_e) in 85% ethanol-water was higher than that in 0.1% hydrochloric acid-water.

3.5.3 Thermodynamics of alkaloids extraction. The thermodynamic parameters of alkaloids from radix physochlainae are summarized in Table 5. The positive value of the change in enthalpy (ΔH°) indicates that the extraction of alkaloids from radix physochlainae is endothermic. The positive value of the change in entropy (ΔS°) indicates the increase in disorder of the studied solid-liquid system due to the solute transfer from more ordered solid phase to the less ordered liquid phase.³⁷ At the beginning of extraction, the target molecules were dispersed in plant cells and underwent two-dimensional movement; then they entered into the solvent system and moved in three-dimensional space during continuous extraction. The negative values of change in Gibbs free energy (ΔG°) indicate the extraction process to be spontaneous.³⁸ Due to the subtle change of pressure and volume in the extraction process, the ΔH° value is approximately equal to the component extraction energy (ΔE°), which is absorbed by the system for extraction of target ingredients. Heating extraction of water bath is one way of heat transfer from outside to inside, and the ionic liquid can competitively form hydrogen bonds with hydroxyl groups on cellulose skeleton and affect the structure of plant cells, so the component extraction energy ΔE° of ionic liquid was higher than that of 85% ethanol-water and 0.1% hydrochloric acid-water. At the same time, the entropy (ΔS°) will rise for the increase of randomness of molecules. With the increase of temperature, the Gibbs free energy (ΔG°) becomes slower,

Table 4 Values of D_e with different solvents and the exponential equation between D_e and T

Extraction solvents	Temperature/K	Exponential equation	R^2	$D_e/10^{-12}$ (m ² min ⁻¹)
[C ₃ tr][PF ₆] aqueous solution	303.15	$10^{-12} D_e = 1.1417 e^{0.0096T}$	0.9971	21.1089
	313.15			22.8204
	323.15			25.1024
	333.15			27.955
	343.15			30.8075
85% ethanol-water	303.15	$10^{-12} D_e = 0.8128 e^{0.0106T}$	0.9751	19.9678
	313.15			21.8078
	323.15			25.9582
	333.15			27.3845
	343.15			30.2370
0.1% hydrochloric acid-water	303.15	$10^{-12} D_e = 0.7312 e^{0.0104T}$	0.9877	17.1153
	313.15			18.2563
	323.15			21.1089
	333.15			23.3909
	343.15			25.3877



Table 5 Thermodynamic parameters of alkaloids from radix physochlainae

Extraction solvents	Temperature/K	K_c	$\Delta H^\circ/\text{kJ mol}^{-1}$	$\Delta S^\circ/\text{J (mol K)}^{-1}$	$\Delta G^\circ/\text{kJ mol}^{-1}$
[C ₃ tr][PF ₆] aqueous solution	303.15	2.0461	53.7965	180.6594	-0.9705
	313.15	2.1862			-2.7770
	323.15	4.6619			-4.5836
	333.15	8.2589			-6.3902
	343.15	24.3876			-8.1968
85% ethanol–water	303.15	1.4382	4.5703	17.9579	-0.8736
	313.15	1.4591			-1.0532
	323.15	1.5927			-1.2328
	333.15	1.6650			-1.4123
	343.15	1.7551			-1.5919
0.1% hydrochloric acid–water	303.15	1.4620	10.8986	39.2689	-1.0051
	313.15	1.7852			-1.3985
	323.15	1.8603			-1.7912
	333.15	2.3060			-2.1839
	343.15	2.4123			-2.5766

which indicates that the entropy (ΔS°) increases more significantly than the enthalpy (ΔH°). In conclusion, the extraction of alkaloids was entropy-driven.³⁹

The thermodynamic parameters for extraction of alkaloids from radix physochlainae with traditional methods have not been reported before, here it can be found the equilibrium constant by [C₃tr][PF₆] aqueous solution is higher than that in 85% ethanol–water and 0.1% hydrochloric acid–water. This result indicates that the solute transferring from the solid phase to liquid phase is more favorable by [C₃tr][PF₆] aqueous solution. In addition, in comparison with other two traditional organic solvents, the change in entropy is higher when [C₃tr][PF₆] aqueous solution is used as extractive solvent. Hence tropane-series alkaloids are more feasible to be extracted by [C₃tr][PF₆] aqueous solution than that by the other two conventional solvents on the basis of the change in Gibbs free energy in the extraction process with [C₃tr][PF₆] aqueous solution, 85% ethanol–water and 0.1% hydrochloric acid–water.

3.5.4 Extraction process characterized by near infrared spectroscopy (NIR). Near infrared spectroscopy (NIR) has been widely used in many fields as one of useful analytical tools, especially for quantitative and qualitative analysis of traditional Chinese medicine.^{40,41} It has the advantages of high speed, convenience, accuracy and non-destruction, and online process monitoring can be easily realized with fiber spectrometer. In this section, the extraction process of alkaloids in 75 °C water bath was monitored by NIR, and target system in 0.05 mol L⁻¹ [C₃tr][PF₆] aqueous solution after extraction for 0 min/30 min/55 min were analyzed. The probe with 2 cm optical length was selected and two optical fibers were used to transmit NIR radiation to a NIRQUEST512 spectrometer; the NIR spectra of extracts were collected when the integration time was 280 ms and the reference value of water solvent was stored as the background before the detection. Consequently, the NIR spectra are shown in ESI 2† and it can be found that the absorbance band of 1400–1500 nm increases gradually with the development of extraction, which belongs to the fragment of benzene ring in the structure of tropane alkaloids and indicates they are continuously extracted from radix physochlainae

during this period. At the same time, the absorbance of 1500–1600 nm also increases, which belongs to the characteristic area of hydroxyl groups.⁴² The increase of –OH concentration also can prove the gradual extraction of tropane alkaloids (containing more hydroxyls than [C₃tr][PF₆]) from radix physochlainae. Above two absorbance bands are the main characteristic regions for the process analysis of the studied extraction, which are ascribed to the major difference of related substituent groups in the structures of tropine-based IL and tropane-series alkaloids. They can be regarded as key observation points for on-line monitoring of this extraction process.

3.5.5 Analysis of herbal powders before and after extraction. In order to comprehensively investigate the extraction behavior in the studied process, FI-IR, SEM and TG were successively used to characterize the samples of radix physochlainae and its residue after extraction by ionic liquids. It can be found from the infrared spectrum (see Fig. 7a) that the wavenumber of main characteristic peaks remains unchanged basically and no new absorbance is generated, which indicates the extraction conditions are mild and tropane ILs cannot deconstruct or strongly combine with the cellulose fiber skeleton of herbal materials. So the coexisting impurities will become much less and ILs can be recovered easily. Meanwhile, compared with the sample before extraction, the absorption intensity of characteristic peak at 1400 cm⁻¹ of C–N bond becomes weak significantly. It can prove that most of tropane alkaloids in radix physochlainae have been selectively extracted by ionic liquids so that the absorbance of their characteristic functional groups disappears. On the other hand, it is obviously shown that the surface structure of radix physochlainae have changed after extraction through the SEM observation (see Fig. 7b). The surface of raw materials is relatively smooth, but the surface of residues after extraction becomes rough and porous. This cellular morphology resulting from IL is very beneficial for permeation and diffusion, and the improved mass transfer is closely related with high extraction efficiency. It is realized through the mechanism that the ionic liquid can competitively form hydrogen bonds with hydroxyl groups on cellulose and affect network of cellulose microfibril.⁴³ According



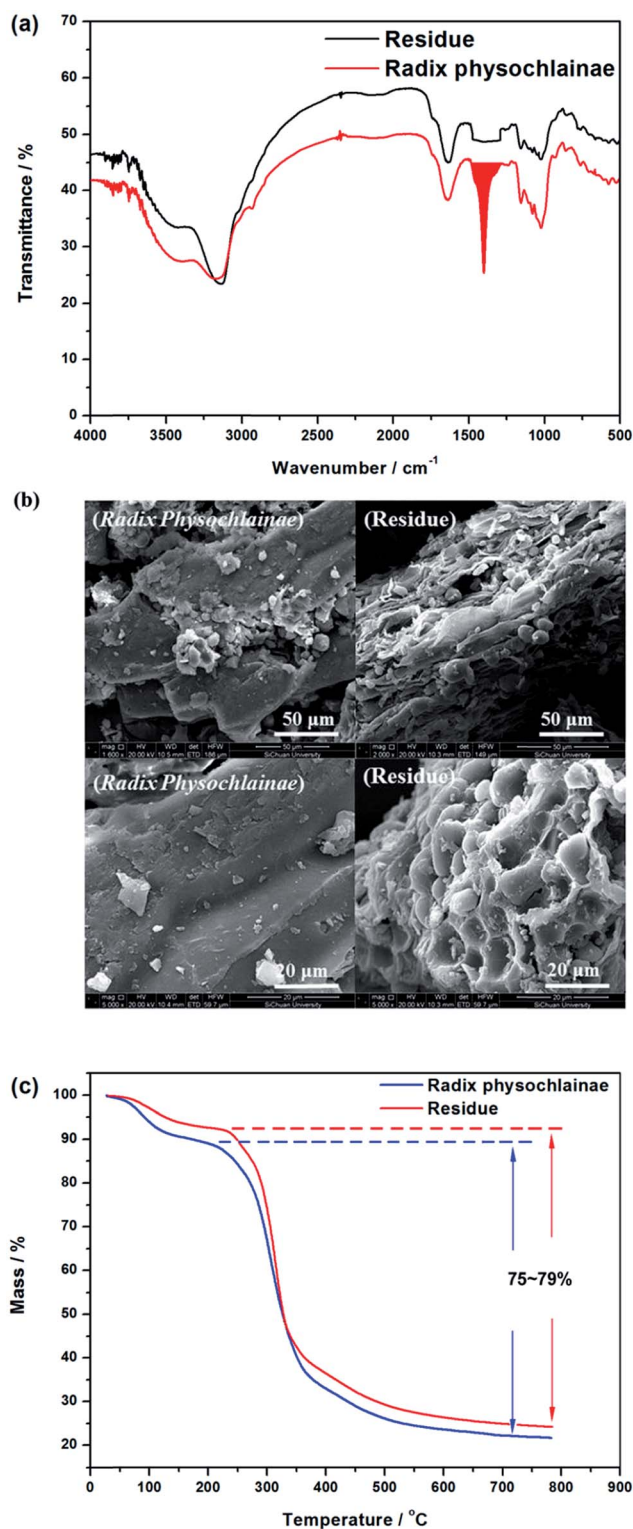


Fig. 7 FT-IR spectra (a), SEM images (b) and TG (c) of radix physochlainae and the extraction residue.

to the TG curves in Fig. 7c, it is found that glucose and fructose⁴⁴ will decompose under 150 °C and the quality of glucose and fructose in herbal material is higher than that in its residue, so the result can be validated that bioactive compounds have been

extracted from radix physochlainae powders. The thermal decomposition of the residue and radix physochlainae starts at 220 °C, and there is a rapid devolatilization process from 300 °C to 320 °C. The weight loss decreases gradually above 500 °C, and the maximum weight loss is between 75% and 79% at the temperature of 700 °C, which results from the decomposition of ionic liquids, cellulose and other coexisting constituents. The temperature of raw material and residue is around 300 °C and 320 °C on maximum mass loss-rate, respectively; it can be ascribed to that the pyrolysis temperature would shift to high temperature through extraction on high temperature.⁴⁵

4. Conclusions

In this paper, the principle of 'like dissolves like' has been practiced through structural similarity between extractant and objects, and seven tropine-based ionic liquids were employed and compared with traditional solvents to extract tropane alkaloids from radix physochlainae, and the cations of ILs had significant influence on their prosperities together with extraction efficiency of alkaloids. [C₃tr][PF₆] aqueous solution was found as the best extractant to achieve high extraction efficiency and purity simultaneously and there was no racemization of the alkaloids occurred in ionic liquid aqueous solution. So this kind of IL can not only realize the selective extraction efficiently, but also maintain the structure and configuration of products through specific intermolecular interactions as extraction solvent. At the same time, the extraction mechanism was tentatively studied through thermodynamic and kinetic analysis together with characterization of radix physochlainae and its residue with FI-IR, SEM and TG technology. It was found that the extraction of alkaloids from radix physochlainae was endothermic, the extraction process was spontaneous and alkaloids are more feasible to be extracted by [C₃tr][PF₆] aqueous solution than 85% ethanol-water and 0.1% hydrochloric acid-water. In addition, a very small volume of chloroform was used to achieve the separation of involved compounds extracted in IL aqueous solution and tropane alkaloids could be easily extracted but ionic liquid did not. As the result, the hypothesis has been successfully proved in this study that the extractant with the same nucleus with objects should be more suitable and efficient in the extraction of the latter, which can be deemed as the generalized 'like dissolves like' theory.

Conflicts of interest

The authors declare no conflict of interest.

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