Chemical Diversity Investigation of Hepatotoxic Pyrrolizidine Alkaloids in Qianliguang (Senecio scandens) and Related Species by UHPLC-QTOF-MS

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ABSTRACT

Objective: Qianliguang (Senecio scandens) is a common Chinese medicinal herb. Qianliguang-containing herbal proprietary products are registered as over-the-counter remedies in China and exported to Western countries. The presence of hepatotoxic pyrrolizidine alkaloids (PAs) has raised concerns about the safety of using Qianliguang and its products. The present study aims at investigation of different types of PAs present in Qianliguang collected from representative locations in China.

Methods: In this study, a simple but specific UHPLC-QTOF-MS method for the determination of toxic PAs was developed, based on the characteristic fragment ions specific to different types of PAs. It was successfully applied for the identification and distinguishing of PAs present in Qianliguang and related Senecio species growing in different locations of China.

Results: Significant diversity of the PA types and quantities were revealed among the samples tested. The estimated total amounts of toxic PAs in three of the samples exceed the toxic limits of PA intake restricted by WHO, demonstrating the timely and highly demand for regulating both types and quantities of PAs present in Qianliguang.

Conclusions: This study provides the methodology for simultaneous identification and quantification of PAs present in herbs without requiring corresponding standards, which could be further used for more systematic investigations of the PA distribution in Qianliguang and other PA-containing herbs.

Key Words: Qianliguang (Senecio scandens), Pyrrolizidine alkaloids, UHPLC-QTOF-MS, Qualification, Quantification

INTRODUCTION

The popular and persistent misconception that nature is always ‘good, clean and healthy’ denies that there are harmful naturally-occurring chemicals, and consequently some toxins-containing medicinal herbs, foods, and dietary supplements are wrongfully accepted as safe. Recent years have witnessed increasing concern about the safety of herbal supplements and botanical products. Among the natural toxins, pyrrolizidine alkaloids (PAs) are probably one of the most significant groups, because PAs are widely distributed across the plant kingdom in up to 13 distantly related angiosperm families covering about 3% of flowering plants worldwide, and about half of them are hepatotoxic and/or tumorigenic to humans[1–3]. PAs are esters of three types of necines: retronecine (including its 7-enantiomer), otonecine, and platynecine[4, 5]. The former two types containing an unsaturated necine bases and are hepatotoxic and can cause hepatic veno-occlusive disease and may ultimately lead to liver cancer in humans, while the platynecine-type PAs with a saturated necine base are generally non-toxic[6]. To date, about 660 PAs have been identified from more than 6000 plant species, and mainly from three families: Asteraceae (Compositae), Boraginaceae and Fabaceae (Liguminosae)[7]. A large number of these plant species have been traditionally used as herbal remedies worldwide.

Qianliguang, derived from the aerial parts of Senecio scandens Buch.-Ham. (Asteraceae family) (Figure 1)[8], is a common Chinese medicinal (CM) herb documented in Pharmacopoeia of P.R. China. This medicinal herb is commonly used to treat bacterial dysentery, enteritis, conjunctivitis, and respiratory tract infections[9–12]. It has also been used either solely or in compound formulae with other herbs in 12 proprietary CM formulations authorized by the State Food and Drug Administration of China as over-the-counter (OTC) remedies, such as Qian Bai Biyan Pian

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1 Hepatotoxic Pyrrolizidine Alkaloids in Qianliguang (Senecio scandens)
Our group has previously established the characteristic mass fragments for identifying and distinguishing different types of PAs, such as ions at m/z 120 and 138 for retronecine-type PAs (RET-PAs); m/z 150 and 168 for otonecine-type PAs (OTO-PAs); and m/z 122 and 140 for platynecine-type PAs (PLA-PAs)[26]. We have also demonstrated that these characteristic mass fragmentation ions can be utilized to determine the presence of PAs in herbal plants and herb-derived products, and to specify the types of PAs[27–32]. In the present study, this method was further applied to the systematic investigation of different types of PAs present in Qianliguang samples collected from representative locations in China, and two additional plant samples, which are different species in Senecio genus and also used as medicinal herbs.

MATERIAL AND METHODS

1. Chemicals

Senecionine and seneciphylline were purchased from Extrasynthese (Genay, France). Senkirkine was from Chromadex (California, USA). Retrorsine was obtained from Sigma (Milwaukee, USA). The mixture of platyphylline and neo-platyphylline (w/w, 3:1, purity > 98%) was a gift from Prof. Hai-Shen Chen from the Department of Phytochemistry, The Secondary Military Medical University, China. Clivorine and integerrimine were isolated in our laboratory by the previously reported method[33, 34]. Riddelliine was a gift from Dr. Po-Chuen Chen, the U.S. National Toxicology Program (NTP).

2. Plant materials

Five samples of Qianliguang (Senecio scandens) were collected from: (i) Conghua, Guangdong Province in September 2007 (SS1); (ii) Nanning, Guangxi Province in July 2005 (SS2); (iii) Damao Mountain, Hong Kong in 2006 (SS3); (iv) Dapu, Hong Kong in 2006 (SS4); and (v) Bozhou, Anhui Province in July 2005 (SS5). Another two samples of Senecio species, Senecio cannabifolius var integri folius Kitam. (SC6) and Senecio nemorensis L. (SN7) were collected from Changbai Mountain, Jilin Province in August 2005 and August 2008, respectively. All the plants used for this study were authenticated by Prof. Zhong-Zhen Zhao, a pharmacognosy professor, School of Chinese Medicine, Hong Kong Baptist University. All the voucher specimens were deposited at School of Chinese Medicine, Hong Kong Baptist University.

3. Sample preparation

The water extract of each herbal sample was prepared by boiling the sample powder (1000 g) with 6 L distilled water under reflux for 1 h. The same extraction procedure was repeated two more times. The aqueous extracts were combined and centrifuged at 3000 g for 10 min. The supernatants were concentrated and then lyophilized to afford a powdered water extract (~26% yield). 1.5 g of the water extract was then dissolved in 50 ml of 0.01% aqueous sulfuric acid solution and centrifuged at 3000 g for 10 min.
An aliquot (25 mL) of the supernatant was adjusted to pH 2–3 with 0.1% aqueous sulfuric acid solution, then extracted with 25 mL of hexane for three times to remove the non-alkaloid components. The resultant aqueous layer was adjusted to pH 9–10 with ammonium hydroxide, and then extracted with 25 mL of dichloromethane for three times. The dichloromethane extracts were combined and dried under reduced pressure by a rotary evaporator. The residue was dissolved in 1 mL of methanol and filtered with a 0.45 μm PTFE membrane filter prior to UHPLC-MS analysis. Reference standards of seneciphylline and senkirkine were dissolved individually in methanol to obtain the corresponding stock solution with a concentration of 500 μM, and diluted to appropriate concentrations for the construction of the calibration curve prior to the UHPLC-MS analysis.

4. UHPLC-MS analysis

UHPLC was performed with a Waters UHPLC system which was equipped with a binary solvent delivery system and coupled to a Bruker MicroOTOF-Q Mass Spectrometer with an ESI source. The chromatography was performed on a Waters Acquity UHPLC BEH shield RP18 column (2.1 × 100 mm, 1.7 μm). A gradient mobile phase system containing (A) 0.03% aqueous diethylamine and (B) acetonitrile was used as follows: 0–1 min, 95%-90% of A; 1–16 min, 90%-65% of A; 16–18 min, 65%-35% of A; 18–18.1 min, 35%-0% of A, and then kept for 2 min. The flow rate was 0.35 mL/min and the injection volume was 2 μL. The mass spectrometer was operated in positive ion mode under full-scan conditions. The operating parameters were set as follows: spray voltage 4500 V, nebulizer 1.5 bar, dry heater 190 °C, dry gas 8.0 L/min, collision energy 25.0 ev, scan range m/z 50-1000.

RESULTS AND DISCUSSION

1. UHPLC-MS analysis of reference PAs

In the present study, RET-PAs (riddelliine, retrorsine, seneciphylline, integerrime and seneconine), OTO-PAs (senkirkine and clivorone) and PLA-PAs (platyphylline and neoplatyphylline) were analyzed to optimize the UHPLC-MS conditions and also obtain the retention times (tR) and MS characteristics for the identification of corresponding PAs in the samples. Under the stated conditions, each PA exhibited the expected protonated molecular ion \([M+H]^+\) along with two specific fragmentation ions: for RET-PAs at m/z 120.0813 and 138.0919, for OTO-PAs at m/z 150.0919 and 168.1025, and for PLA-PAs at m/z 122.0970 and 140.1075, respectively (Figure 2). These characteristic fragmentation ions are in good agreement with our previously established MS patterns for identifying and distinguishing the types of PAs of interest[26]. To be simplified, the MS data of these characteristic fragmentation ions shown below are described as the nominal mass m/z 120 (for m/z 120.0813), m/z 138 (for m/z 138.0919), m/z 150 (for m/z 150.0919), m/z 168 (for m/z 168.1025), m/z 122 (for m/z 122.0970) and m/z 140 (for m/z 140.1075). The structures of the fragment ions specific to the three types of PAs are shown in Figure 3.

2. Qualitative analysis of Qianliguang and related Senecio species

The qualitative identifications of PAs in the extracts of Qianliguang and related Senecio species were obtained from the extracted ion chromatograms (EICs), based on the characteristic fragment ions specific to three different types of PAs. In the EICs of Qianliguang (SS1) collected from Guangdong Province (Figure 4A), two peaks, eluted at 4.2 min and 5.2 min, had the characteristic fragmentation ions of OTO-PAs at both m/z 150 and 168, suggesting that two OTO-PAs were present in this herbal sample. The PA eluted at 5.2 min had the protonated molecular ion at m/z 366.1917 and was unequivocally identified as senkirkine by the comparison of retention time, MS characteristics and molecular weight (MW) to those of the reference standard (Table 1). The MW of the PA eluted at 4.2 min was 2 units less than senkirkine, suggesting the loss of two protons, thus it was tentatively identified as dehydrosenkirkine. In addition, there are other two PAs, eluted at 7.4 min and 12.2 min, possessing the characteristic ions of PLA-PAs at m/z 122 and 140. The peak at 12.2 min had the protonated molecular ion at m/z 340.2124, which was 2 units more than that of platyphylline (or neoplatyphylline). Thus it was tentatively identified as dihydroplatyphylline (or dihydronapotyphylline). The other PLA-PA, eluted at 7.4 min, had the protonated molecular ion at m/z 240.1600, which matched that of six PLA-PAs previously isolated from Senecio species, namely 7 (or 9)-angeloylplatynecine, 7 (or 9)-senecioylplatynecine, and 7 (or 9)-tigloylplatynecine (Supplemental Figure 1)[35]. Therefore, this PLA-PA could be one of these six PAs.

For the other four Qianliguang samples (SS2–SS5) collected from different locations in China, OTO-PAs senkirkine and dehydrosenkirkine were also detected in SS2–4 but not in SS5 (Table 1). No other types of PAs were detected in SS2 (collected from Guanxi Province). In the two Qianliguang samples collected from Hong Kong (SS3 and SS4), a RET-PA eluted at 6.5 min exhibited two characteristic ions at m/z 120 and 138, and was detected in both of them. It had the same MW (349 Da) as riddelliine and other six RET-PAs previously identified in various Senecio species (structures shown in Supplemental Figure 2)[24, 36–40]. Due to the different retention time of riddelliine (tR 6.9 min), this RET-PA could be one of the other six PAs. Additionally, in sample SS3, seneciphylline eluted at 9.2 min was unequivocally identified by comparison with the reference standard. Two other RET-PAs eluted at 8.2 min and 8.8 min were found in SS3. They had a same MW of 385 Da, which is 2 units more than that of jaconine[36], and thus were tentatively identified as dihydrojaconine and its isomer.

In sample SS4, there were two additional PLA-PAs, eluted at 7.4 min and 11.7 min, with the same MW of 239 Da. As described for sample SS1, they might be 7 (or 9)-angelyloylplatynecine, 7 (or 9)-senecioylplatynecine, or 7 (or 9)-tigloylplatynecine[37]. In the Qianliguang collected from Anhui (SS5), seneciphylline, neoplatyphylline and seneconine were eluted at 9.1 min, 10.7 min and 11.0 min, respectively, and were unequivocally identified by comparison with the reference standards. Another PA, eluted at 3.4 min, had a MW
Figure 2. Mass spectra of (A) Riddelliine, (B) Retrorsine, (C) Seneciphylline, (D) Integerrimine, (E) Senecionine, (F) Senkirkine, (G) Clivorine, (H) Platyphylline, (I) Neoplatyphylline
of 365 Da and the two characteristic ions of RET-PAs. Its MW was identical to that of adonifoline\textsuperscript{[24]}, which makes it a candidate for this PA.

For both of two herbs of Senecio species (SC6 and SN7), a peak was eluted at 6.9 min with the two characteristic ions of RET-PAs at \(m/z\) 120 and 138 (Figure 4B and 4C). It had the same MW, 351 Da, as three RET-PAs (jacobine, retrorsine and usaramine) reported from Senecio species\textsuperscript{[35]}. Due to the different chromatographic property of retrorsine (\(t_R\) 8.2 min), usaramine and jacobine were more likely candidates for this PA. Another RET-PA present in both samples was eluted at 13.4 min, with a MW of 335 Da, corresponding to that of integerrimine (\(t_R\) 10.8 min) and senecionine (\(t_R\) 11.0 min). Therefore it was tentatively identified as an isomer of integerrimine or senecionine. In addition, another isomer of integerrimine or senecionine with the same MW of 335 Da but eluted at 13.8 min, was detected in SN7 (Table 1). Also in SN7, a PLA-PA was eluted at 14.1 min with the protonated molecular ion at \(m/z\) 338.1967 corresponding to that of platyphylline (or neoplatyphylline) but having a different retention time. Therefore it was tentatively identified as an isomer of platyphylline or neoplatyphylline. In SC6, there was another PLA-PA eluted at 11.7 min with a MW of 239 Da, which was the same as that detected in sample SS4.

As described above, the diverse chemical profiles of the PAs present in Qianliguang and related Senecio species, which were collected from different locations and harvested at differential times of the year, were observed. It is important to note that among the unequivocally identified PAs, senecionine (RET-PA), seneciphylline (RET-PA), and senkirkine (OTO-PA) are known hepatotoxins and tumorigens, and have been reported to induce liver tumors in experimental animals\textsuperscript{[41]}. On the other hand, adonifoline was only found in one sample (SS5), while additional 15 PAs were identified in different samples. Notably, some of toxic PAs had significantly higher quantities than adonifoline in the samples tested, indicating that the regulation of only adonifoline in Qianliguang by Pharmacopoeia of P.R. China (2010 edition) is obviously not adequate.
3. Quantification of toxic PAs in Qianliguang and related Senecio species

Because the hepatotoxicity of PA-containing herbs occurs only with the unsaturated PAs\(^3\), i.e. RET-PAs and OTO-PAs, we therefore further quantified these two toxic types of PAs in all samples tested. Due to the unavailability of most of the reference standards, RET-PAs and OTO-PAs were semi-quantified using seneciphylline and senkirkine as the representative standard for each type, respectively. The peaks corresponding to the protonated molecular ion of seneciphylline at \(m/z\) 334.1649, senkirkine at \(m/z\) 366.1911, and internal standard (clivorine) at \(m/z\) 406.1860 were extracted. The calibration curves were constructed using the peak area ratio (analyte/internal standard) over the concentration ranges of 0.03–10 µM for seneciphylline and 0.03–30 µM for senkirkine. The regression equations were \(y = 1.322 x + 0.0708\), \(R^2 = 0.9997\) for seneciphylline, and \(y = 0.7233 x + 0.4126\), \(R^2 = 0.9952\) for senkirkine, with good linearity at

![EICs of Qianliguang samples](image)

Figure 4. EICs of Qianliguang samples: (A) Qianliguang (SS1) collected from Guangdong Province; (B) Senecio cannabifolius var integrifolius Kitam. (SC6) collected from Jilin Province; (C) Senecio nemorensis L. (SN7) collected from Jilin Province.
the concentration ranges tested. The limit of quantification (LOQ) was 0.03 µM for both PAs. The protonated molecular ions of RET-PAs and OTO-PAs identified in qualitative analysis were extracted and then quantified. The results are presented in Table 2 and Figure 5. Two OTO-PAs, namely senkirkine and dehydrosenkirkine, were found to be the major toxic PAs present in Qianliguang from Guangdong (SS1), Guangxi (SS2), Damao Mountain, Hong Kong (SS3). By contrast, Qianliguang from Anhui (SS5), and two herbs of Senecio species (Senecio cannabifolius var integrifolius Kitam. (SC6) and Senecio nemorensis L. (SN7)) contained only RET-PAs. More importantly, quantity of toxic PAs in these samples varied significantly. Three samples, including SS1, SC6 and SN7 contained high quantities (100–300 µg/g) of toxic PAs, while Qianliguang from Dapu, Hong Kong (SS4) had a lowest amount (about 0.2 µg/g) of toxic PAs. It is also worthy to note that the two samples of different Senecio species (SC6 and SN7) contained significantly high amounts of toxic RET-PAs.

S. cannabifolius var integrifolius Kitam. (SC6) also called ‘Daye Fanhuncao’ is widely used in China as medicinal herbs for the treatment of maladies such as phlegm and asthma. S. nemorensis L. (SN7) called ‘Linyin Qianliguang’ is used for the treatment of carbuncles and furuncles. Therefore, such herbal plants in Senecio genus have also posed potential risk for the PA-intoxication and require related regulations.

Based on the estimated quantity of toxic PAs in the tested herbs and the dosage (0.25–0.50 g/kg/day) of Qianliguang recommended by Pharmacopoeia of P. R. China[9], amount of total toxic PAs by the intake of Qianliguang and Senecio species investigated in this study was calculated and shown in Table 2. Among them, the estimated dosages of SS1, SC6 and SN7 exceeded the toxic dose of PA intake in humans (15 µg/kg, b.w./day based on the use of comfrey, a PA-containing herb) specified by the World Health Organization (WHO)[42]. Furthermore, if calculated by the daily consumption of QBBP recommended by Pharmacopoeia of P.R. China (i.e. 9–12 tablets of QBBP, equivalent to 21.8–29.1 g of Qianliguang)[8], the daily intake of SS1, SC6 and SN7 exceeded the short term limit of PAs in QBBP restricted by the United Kingdom Medicines and Healthcare Product Regulatory Agency (1000 µg/day for less than 2 weeks)[14].
Calculated intake

| Amounts | SS1 | SS2 | SS3 | SS4 | SS5 | SC6 | SN7 |
|---------|-----|-----|-----|-----|-----|-----|-----|
| RET-PAs (µg/g) | 105.9 | 0.57 | 3.47 | 0.03 | Nil | Nil | Nil |
| OTO-PAs (µg/g) | 105.9 | 0.57 | 3.95 | 0.19 | 6.18 | 303.4 | 101.3 |
| Total toxic PAs (µg/g) | 26.48 ~ 52.95 | 0.14 ~ 0.28 | 0.98 ~ 1.97 | 0.05 ~ 1.54 ~ 3.09 | 75.85 ~ 151.70 | 25.32 ~ 50.65 |
| Calculated dosage (µg/kg. b.w./day)** | 14.58 ~ 26.48 | 0.28 | 61.48 | 5.28 | 303.4 | 101.3 |
| Calculated intake (µg/day)*** | 2580.62 | 12.42 | 86.11 | 4.14 | 134.72 | 6614.12 | 2208.34 |

* b.w.: body weight; ** Calculated by the dosage (0.25–0.50 g/kg/day) of Qianliguang recommended by the Pharmacopoeia of P. R. China; *** Calculated by the daily consumption of QBPP recommended by the Pharmacopoeia of P. R. China (i.e. 9–12 tablets of QBPP, equivalent to 21.8–29.1 g of Qianliguang).

CONFLICTS OF INTEREST

The authors have declared no conflict of interest.

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Supplemental Figure 1. Six PLA-PAs with MW of 239 from Senecio species.
Supplemental Figure 2. Six RET-PAs with MW of 349 from Senecio species.