Marker compounds contents of *Salvia miltiorrhiza* Radix depending on the cultivation regions

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Received: 25 February 2019 / Accepted: 19 March 2019 / Published Online: 30 June 2019
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Abstract *Salvia miltiorrhiza* Radix is cultivated in Korea and China and is traditionally used to treat cardiovascular diseases. In this study, we developed and validated a quantitative analysis method for *S. miltiorrhiza* Radix using high-performance liquid chromatography (HPLC). Identification was performed using ultra performance liquid chromatography-tandem mass spectrometry. For quantitative analysis, we used seven marker compounds. Separation conditions for HPLC were optimized using an ODS column with gradient conditions of 1% formic acid in distilled water and 1% formic acid in acetonitrile, with a flow rate of 0.8 mL/min and a detection wavelength of 280 nm. This method showed good linearity ($R^2=0.9998$), precision (relative standard deviation $\leq 3.3\%$), accuracy (recovery of 94.16–102.89%), limit of detection (7.53 $\mu\text{g/mL}$), and limit of quantification (23.71 $\mu\text{g/mL}$). This approach successfully quantified marker compounds in *S. miltiorrhiza* Radix. The individual marker compounds were identified by comparing the molecular masses and retention times with does standard compounds. Marker compound contents of *S. miltiorrhiza* Radix were investigated with different cultivation regions. Seven marker compounds were detected and quantified in all samples. Among them, salvianolic acid B showed the highest contents and it ranged from 4.13 to 7.15%. The salvianolic acid B content (7.15%) of marker compound was the highest in Bonghwa, and the tanshinone IIA content (1.90%) was the highest in Pohang. The results of marker compounds and developed

method were intended to provide a favorable reference for the study of *S. miltiorrhiza* Radix from different regions of Korea.

Keywords Cultivation region · High-performance liquid chromatography · Liquid chromatography-mass spectrometry · *Salvia miltiorrhiza* Radix · Salvianolic acid B · Tanshinone IIA

Introduction

Salvia miltiorrhiza Radix is cultivated in China and Korea [1] and is used as a traditional medicine to treat coronary, cerebrovascular diseases, hepatitis, cardiovascular diseases, and chronic renal failure [2,3]. Its various biological effects, including inhibition of lipid peroxidation, and anticancer, antioxidant, antimicrobial, and immune activities, have also been studied and reported [4-7]. These biological effects have been found to be mainly caused by marker compounds including hydrophilic phenolic acids and lipophilic diterpenes [8-10]. For quality control in its medicinal products, salvianolic acid B, tanshinone IIA, cryptotanshinone, and tanshinone I have been selected as chemical markers by the Chinese Pharmacopoeia. Salvianolic acid B has also been selected as a chemical marker by the Korean Pharmacopoeia. Therefore, since rapid and repeatable quantification of salvianolic acid B and tanshinone IIA contents are essential for evaluating quality, a suitable analysis method is required. Several analytical methods have been used to analyze *S. miltiorrhiza* Radix and its products [11-13]. Most plants contain different marker compounds whose activities depend on the variety, plant part, growth time, harvest time, and production region [14-16]. In this study, an HPLC approach was developed and validated for quantifying marker compounds, which were then identified using ultra performance liquid chromatography-tandem mass spectrometry (UPLC-MS). We also investigated geographical differences in these marker compounds occurring in *S. miltiorrhiza* sourced from different cultivation regions.

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Materials and Methods

Chemicals and materials

S. miltiorrhizae Radix were collected from different regions in Korea well-known for their cultivation of medicinal plants: Bonghwa (K-1), Gochang (K-2), Jangheung (K-3), Pohang (K-4), and Yeongyang (K-5). A Chinese-originating sample of *S. miltiorrhizae* Radix was purchased from a traditional medicine local market in Daegu, Korea. Standard compounds were purchased from Sigma-Aldrich (St. Louis, MO, USA). Water was triply distilled. HPLC grade acetonitrile was obtained from J.T. Baker (Phillipsburg, NJ, USA). All other reagents and solvents were of analytical grade.

Sample preparation and HPLC analytical conditions

S. miltiorrhizae Radix was dried at room temperature in the shade, then pulverized and sieved to the powder using >60 μm mesh. To prepare samples for analysis, 0.3 g of powder was sonicated with 50 mL of 75 % ethanol for 30 min and filtered through a 0.45 μm filter. We used an HPLC system (Waters Corp., Milford, MA, USA) with an ODS H80 analytical column (4.6 $\times$ 250 mm ID, 4 μm , YMC Co., Kyoto, Japan). The mobile phases consisted of distilled water (solvent A) and acetonitrile (solvent B), both containing 1% formic acid (v/v). Gradient elution for HPLC analysis was conducted for chromatographic separation, using gradient programs of 0.00–10.00 min, 25–25% B; 10.00–20.00 min, 25–60% B; 20.00–25.00 min, 60–85% B; and 25.00–40.00 min, 85–85% B. Flow rate was 0.8 mL/min, with detection wavelength at 280 nm, and injection volume of 10 μL . Marker compounds of the sample were identified by comparing their retention times with those of standards. Compounds were tentatively quantified on the basis of dry weight using the linear regression equation obtained from the standard calibration curves.

HPLC analytical method validations of marker compounds

Validation was carried out according to internationally accepted criteria [17]. The parameters evaluated were system suitability, selectivity, linearity, precision and accuracy, limits of detection and quantification, and system suitability [18]. The linearity of the marker compounds was determined using external standards. Reference solutions of salvianolic acid B at concentrations of 25–400 $\mu\text{g/mL}$, salvianolic acid A at concentrations of 50–200 $\mu\text{g/mL}$, and the other five compounds at concentrations of 25–200 $\mu\text{g/mL}$ were analyzed under optimized chromatographic conditions. For each sample analyte, LOD and LOQ were defined as the lowest concentration that could be determined with a signal to noise ratio (S/N) of three. The accuracy of the proposed method was indicated by the percent recovery of the four different concentrations of marker compounds extracted from the prepared sample solution using a stock solution. The percent relative standard deviation was calculated as a measure of method

precision. Accuracy and precision were determined by multiple analysis of samples prepared at different concentrations on a single day and different days ($n=3$).

Identification of marker compounds using UPLC-MS

Marker compounds were identified using UPLC-MS-MS, consisting of a Waters UPLC system with a Waters Xevo TQ-S micro MS with electrospray ionization source. Chromatography was performed on an Acquity UPLC BEH ODS column (2.1 $\times$ 100 mm ID, 1.7 μm). The mobile phase consisted of 0.1% formic acid in distilled water (solvent A) and acetonitrile (solvent B), with a flow rate of 0.392 mL/min and an injection volume of 5 μL . The gradient elution for UPLC analysis was conducted for chromatographic separation, with gradient programs of 0.00–1.70 min, 25–25% B; 1.70–3.40 min, 25–60% B; 3.40–4.25 min, 60–85% B; and 4.25–6.80 min, 85–85% B. The electrospray ionization interface was conducted both positive and negative modes.

Statistical analysis

Data are expressed as mean $\pm$ standard deviation ($n=3$). Duncan's multiple range tests with significance judged at $p < 0.05$ were performed to determine significant differences between the means using SAS software (SAS, Inc., Cary, NC, USA).

Results and Discussion

Validations of marker compounds in *S. miltiorrhiza* Radix using HPLC

Retention times of marker compounds were 3.8, 10.6, 13.7, 18.9, 30.5, 30.8, and 33.9 min for salvianic acid A, rosmarinic acid, salvianolic acid B, salvianolic acid A, cryptotanshinone, tanshinone I, and tanshinone IIA, respectively (Fig. 1). Salvianic acid A, rosmarinic acid, salvianolic acid B, and salvianolic acid A correspond to hydrophilic phenolic compounds, whereas tanshinone IIA, tanshinone I, and cryptotanshinone represent lipophilic diterpene compounds [19]. Therefore, all compounds in *S. miltiorrhizae* Radix have broad ranges of polarity [20]. Regression equations for linearity, LOD, and LOQ of the seven compounds were calculated based on the mean peak areas versus the standard concentrations (Table 1). Each coefficient of correlation (R^2) was ≥ 0.9998 , as determined by the least square analysis, suggesting good linearity. The LOD and LOQ of standard compounds were 2.56–7.83 and 7.76–23.71 $\mu\text{g/mL}$, respectively. Intra-day samples showed intra-day assay precision below 1.24% and accuracies ranging from 98.73 to 102.17% (Table 2). Inter-day samples showed precision below 3.30% and accuracies ranging from 94.16 to 102.89%. Variations in precision were less than 3.3%, and variations in accuracy ranged from 94.16 to 102.89%. These results indicate that this HPLC method provides good accuracy, consistent with previous reports [12].

Fig. 1 HPLC chromatograms of standard compounds (A) and *S. miltiorrhiza* Radix extract (B)

Table 1 Linearity, LODs, and LOQs of marker compounds

Compounds	Regression equation ^a	Correlation coefficient (R^2)	Linear range ($\mu\text{g/mL}$)	LOD ($\mu\text{g/mL}$)	LOQ ($\mu\text{g/mL}$)
salvianic acid A	$Y=3.9135x+11.417$	0.99995	25-200	2.56	7.76
rosmarinic acid	$Y=8.8977x+13.407$	0.99986	25-200	4.57	13.86
salvianolic acid B	$Y=5.0297x-24.748$	0.99980	25-400	7.83	23.71
salvianolic acid A	$Y=3.2754x-2.85$	0.99987	50-200	5.23	15.85
cryptotanshinone	$Y=13.219x-55.054$	0.99992	25-200	3.42	10.37
tanshinone I	$Y=10.204x+48.804$	0.99981	25-200	5.29	16.02
tanshinone IIA	$Y=7.1288x-26.053$	0.99991	25-200	3.69	11.19

^ay = mean peak area; x = concentration of the standard solution (g/mL)

Identification of marker compounds in *S. miltiorrhiza* Radix using UPLC-MS

Separated marker compounds of *S. miltiorrhiza* Radix were identified by comparison with standards using UPLC retentions time and mass spectra analyses (Table 3). The retention times of marker compounds were 0.83, 1.64, 1.93, 3.89, 5.78, 5.80, and 6.24 min for salvianic acid A, rosmarinic acid, salvianolic acid B, salvianolic acid A, cryptotanshinone, tanshinone I, and tanshinone IIA, respectively, which were the same as those of standards. Chemical structures and MS/MS spectra are shown in Fig. 2. The mass spectrum of marker compounds showed significant ions at m/z values of 197.04, 359.15, 717.25, 493.04, 277.16, 297.17, and

295.23, which matched with the molecular weights of marker compounds, respectively [21].

For phenolic acid compounds, small molecules such as carbon dioxide, carbon monoxide, and water were produced by the monomers because of the presence of carboxyl, carbonyl, or hydroxyl groups [22]. Salvianic acid A showed as deprotonated molecules at m/z 197.04. The fragment ions corresponded to CO and H_2O loss from the parent ion. Rosmarinic acid showed as deprotonated molecules at m/z 359.15. Fragment ions at m/z 161.07 and 179.22 corresponded to salvianic acid A or caffeic acid loss from the parent ion. Salvianolic acid B showed as deprotonated molecules at m/z 717.25. The fragment ions m/z

Table 2 Intra-day and inter-day precision (RSD, %) and accuracy (recovery, %) of marker compounds

compounds	Concentration (g/mL)	Intra-day (n=3)		Inter-day (n=3)	
		precision (RSD, %)	accuracy (recovery, %)	precision (RSD, %)	accuracy (recovery, %)
salvianic acid A	200	1.01	100.29	1.27	99.47
	100	0.68	99.14	1.44	99.87
	50	1.16	99.65	1.91	100.48
	25	1.24	99.58	3.30	97.85
rosmarinic acid	200	0.95	99.52	1.43	100.72
	100	0.63	99.43	2.50	99.69
	50	0.89	100.19	2.19	98.79
	25	0.78	102.17	1.55	101.57
salvianolic acid B	400	0.93	99.78	1.37	101.91
	200	0.92	99.83	1.31	101.87
	100	0.43	100.51	0.86	102.89
	50	0.75	100.72	1.64	102.72
	25	0.62	100.19	1.08	99.87
salvianolic acid A	200	0.92	100.61	1.54	99.88
	150	0.36	101.60	1.26	98.50
	100	0.90	100.57	2.23	102.61
	50	0.60	100.11	1.22	101.19
cryptotanshinone	200	0.41	101.35	1.41	99.53
	100	0.48	100.26	1.49	99.28
	50	0.15	100.34	1.50	95.56
	25	0.19	100.20	0.52	94.16
tanshinone I	200	0.23	99.79	1.67	100.86
	100	0.68	99.47	2.28	101.77
	50	0.64	99.78	2.94	99.52
	25	0.62	102.08	2.89	104.11
tanshinone IIA	200	0.39	99.71	2.39	97.04
	100	0.31	98.73	2.30	101.15
	50	0.95	100.59	1.62	98.67
	25	0.26	99.61	2.64	101.00

321.15 and 519.13 corresponded to one or two ionized molecules of salvianic acid A (m/z 198, 198×2) loss from the parent ion. Salvianolic acid A showed as deprotonated molecules at m/z

493.04. Fragment ions at m/z 295.22 were formed by the loss of salvianic acid A (m/z 198) from the parent ion. Tanshinone I showed as protonated molecules at m/z 277.16. The primary fragment ions at m/z 249.17 corresponded to CO loss from the parent ion. Fragment ions at m/z 221.17 and 234.23 were formed by the losses of CO and CH_3 from the parent ion. Cryptotanshinone showed as protonated molecules at m/z 297.17. The primary fragment ion at m/z 254.30 corresponded to CO and CH_3 loss from the parent ion. Other fragment ions at m/z 251.24, 268.24, and 282.12 were formed by the losses of CO, CH_3 , and H_2O from the parent ion. Tanshinone IIA showed as protonated molecules at m/z 295.23. The primary fragment ion of tanshinone IIA at m/z 206.23 corresponded to H_2O , CO, and CH_3 loss from the parent ion. Secondary fragment ions at m/z 234.11, 249.30, 262.18, and 277.31 were formed by the losses of CO, CH_3 , and H_2O from the parent ion. Based on these fragmental patterns and comparison with reference standards and literature data [23,24], the marker compounds could be identified using this method.

Contents of marker compounds in *S. miltiorrhiza* Radix from different cultivation regions

The marker compound contents of *S. miltiorrhiza* Radix from different cultivation regions were calculated using regression equations (Table 4). Marker compounds were detected and quantified in the samples. The contents of salvianic acid A, rosmarinic acid, salvianolic acid B, salvianolic acid A, cryptotanshinone, tanshinone I, and tanshinone IIA contents were 0.00–0.03, 0.24–0.55, 4.13–7.15, 0.04–0.13, 0.07–0.57, 0.00–0.47, and 0.14–1.90%, respectively. High contents of salvianolic acid B were detected in all samples, with the K-1 sample (7.15%) containing the most. The salvianolic acid B content of *S. miltiorrhiza* Radix from different cultivation regions in Korea (5.53%) was higher than that of the Chinese samples (4.42%). Tanshinone IIA, tanshinone I, and cryptotanshinone contents were also higher in the Korean samples than in the Chinese samples ($p < 0.05$). Salvianolic acid A, which is one of the marker compounds of *S. miltiorrhiza* Radix, was not detected in this study; ordinarily, its content is low, though it can be converted from salvianolic acid B under high temperature and pressure [25–

Table 3 Identification of marker compounds in *S. miltiorrhiza* Radix samples by ultra-performance liquid chromatography-tandem mass spectrometry (UPLC-MS-MS) analysis

Compounds	Retention time (min)	Masses of $[\text{M}-\text{H}]^-$ or $[\text{M}+\text{H}]^+$ (m/z)	Fragmentation of ions (m/z)	
			ESI (–)	ESI (+)
salvianic acid A	0.83	197	179, 135	
rosmarinic acid	1.64	359	197, 179, 161	
salvianolic acid B	1.93	717	717, 519, 321	
salvianolic acid A	3.89	493	493, 295	
tanshinone I	5.78	277		249, 234, 221
cryptotanshinone	5.80	297		282, 268, 254, 251
tanshinone IIA	6.24	295		295, 277, 262, 249, 234, 206

Fig. 2 Chemical structures and MS/MS spectra of marker compounds in *S. miltiorrhiza* Radix

27]. Other studies have also shown that salvianolic acid B shows the highest content compared to other marker compounds [28], accounting for 80-95% of the total contents. Overall, the contents quantified in this study were similar to or higher than those

reported previously [29-31]. The present results enable quality control of traditional medicinal preparations containing *S. miltiorrhiza* and provide a reasonable strategy for developing novel functional food products.

Table 4 Contents (dry weight basis, %) of marker compounds in *S. miltiorrhiza* Radix depending on different cultivation regions

Samples	salvianic acid A	rosmarinic acid	salvianolic acid B	salvianolic acid A	cryptotanshinone	tanshinone I	tanshinone IIA
K-1 ^a	0.00±0.01 ^C	0.55±0.07 ^A	7.15±0.98 ^A	0.13±0.01 ^A	0.13±0.01 ^{BCD}	0.01±0.01 ^C	0.37±0.03 ^{CD}
K-2	0.03±0.03 ^A	0.46±0.09 ^{AB}	5.88±1.11 ^{AB}	0.08±0.02 ^{ABC}	0.18±0.02 ^B	0.09±0.03 ^B	0.68±0.11 ^B
K-3	0.00±0.00 ^C	0.33±0.08 ^{BC}	5.08±0.98 ^{BC}	0.08±0.08 ^{ABC}	0.15±0.02 ^{BC}	0.01±0.01 ^C	0.41±0.07 ^C
K-4	0.00±0.00 ^C	0.24±0.04 ^C	4.34±0.68 ^C	0.04±0.00 ^C	0.57±0.08 ^A	0.47±0.09 ^A	1.90±0.29 ^A
K-5	0.00±0.00 ^C	0.41±0.02 ^{AB}	5.23±0.31 ^{BC}	0.10±0.01 ^{AB}	0.15±0.00 ^{BC}	0.03±0.00 ^C	0.47±0.02 ^C
K	0.01±0.02	0.40±0.12	5.53±1.22	0.09±0.04	0.24±0.17	0.12±0.19	0.77±0.61
C-1	0.00±0.00 ^C	0.40±0.08 ^B	4.14±0.76 ^C	0.06±0.01 ^{BC}	0.07±0.00 ^D	0.00±0.00 ^C	0.19±0.02 ^{DE}
C-2	0.02±0.01 ^{AB}	0.43±0.13 ^{AB}	4.13±0.12 ^C	0.07±0.02 ^{ABC}	0.07±0.00 ^D	0.00±0.00 ^C	0.16±0.03 ^E
C-3	0.01±0.01 ^{BC}	0.41±0.06 ^{AB}	5.00±0.62 ^{BC}	0.07±0.02 ^{ABC}	0.10±0.04 ^{CD}	0.00±0.00 ^C	0.14±0.01 ^E
C	0.01±0.01	0.41±0.08	4.42±0.66	0.07±0.02	0.08±0.02	0.00±0.00	0.17±0.03

Means followed by the same superscript letters (A-E) within the column are not significantly different ($p < 0.05$)

^aK-1, Bonghwa; K-2, Gochang; K-3, Jangheung; K-4, Pohang; K-5, Yeongyang

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