


Cite this: *RSC Adv.*, 2020, 10, 44050

# Optimization of flash extraction, separation of ginsenosides, identification by HPLC-FT-ICR-MS and determination of rare ginsenosides in mountain cultivated ginseng†

Lei Xu,<sup>‡a</sup> Jing Xu,<sup>‡a</sup> Guohui Shi,<sup>a</sup> Shengnan Xiao,<sup>a</sup> Rongke Dai,<sup>a</sup> Shao Wu,<sup>a</sup> Baoshan Sun,<sup>\*a</sup> Xiaoshu Zhang<sup>\*a</sup> and Yuqing Zhao<sup>id</sup> <sup>\*ab</sup>

In this paper, we used the flash extraction method (FEM) to extract ginsenosides from mountain cultivated ginseng (MCG), optimized the FEM process by response surface methodology (RSM), and separated 23 kinds of ginsenosides from MCG, including rare ginsenoside Rg<sub>3</sub>, 20(R/S)-Rg<sub>2</sub>, Rk<sub>3</sub>, 20(S)-Rh<sub>2</sub>, 20(R)-Rh<sub>1</sub>, F<sub>1</sub> and Rg<sub>6</sub>. Among them, notoginsenoside R<sub>1</sub> was isolated from MCG for the first time. Additionally, we established an HPLC-FT-ICR-MS method to accurately identify 20 ginsenosides in MCG, and quantitatively analyzed the differences in the content of rare ginsenosides in MCG and Garden-Cultivated Ginseng (CG) by HPLC-UV. The results showed that the chemical components of MCG and CG were similar, but the ginsenoside content of MCG was double that of CG. Notably, the content of ginsenoside 20 (S)-Rh<sub>2</sub> and 20 (R)-Rh<sub>1</sub> had the largest difference, and the content in MCG was 33 and 24 times higher than that in CG, respectively. Through quantitative analysis, we clarified the reason why the activity of MCG is stronger than that of CG, which provided a theoretical basis for clinical application and further research of MCG.

Received 1st September 2020  
Accepted 21st November 2020

DOI: 10.1039/d0ra07517e

rsc.li/rsc-advances

## 1 Introduction

*Panax ginseng* Meyer is a slow-growing perennial plant belonging to the Araliaceae family and is widely grown in regions of Korea, China, and Asian countries. As a famous tonic in traditional Chinese medicine in the northeast of China, ginseng has been used for a long time in the treatment of splenic asthenia, wheezing, insomnia and spontaneous perspiration. It has been used as a medicinal plant in China for more than 2000 years.<sup>1</sup> Recent reports have shown that ginseng possesses numerous biological activities, including anti-cancer,<sup>2</sup> immunomodulatory effects,<sup>3</sup> anti-oxidant,<sup>4</sup> anti-aging,<sup>5</sup> anti-inflammation,<sup>6</sup> anti-diabetic.<sup>7</sup> Studies have shown that these beneficial effects are attributed to ginsenosides, polysaccharides, and volatile oils. Ginsenosides in particular have been shown to exhibit significant anti-oxidant, anti-cancer, and anti-apoptotic activity.<sup>8,9</sup>

In recent years, due to the increased destruction of forest resources and the predatory excavation of ginseng, the

availability of wild ginseng resources had plummeted. Mountain cultivated ginseng (MCG), which simulates the growth habit and ecological environment of wild ginseng, can be regarded as an imitation of wild ginseng.<sup>10</sup> Many results showed that the MCG had stronger pharmacological activities than Garden-Cultivated Ginseng (CG).<sup>11</sup> MCG has been collected in the Chinese pharmacopoeia since 2005,<sup>12</sup> considered as the sources of ginseng, using the same standard with CG. But at present there are no clear methods for the identification of two kinds of ginseng, which can easily lead to ginseng market chaos. In the Chinese pharmacopoeia and many existing ginseng standards, the extraction of ginseng requires a long time of heating with some ginsenosides may be transformed.<sup>13</sup> Therefore, there is an urgent need for a room temperature and fast extraction method in order to better extract and distinguish MCG and CG.

The flash extraction method (FEM) is a convenient and efficient extraction process that can effectively shorten room temperature extractions.<sup>14</sup> FEM performs high-speed stirring, strong vibration, and negative pressure percolation to achieve extractions, which are conducive to the extraction of heat-sensitive components.<sup>15</sup> Song<sup>16</sup> used FEM to rapidly isolate 27 compounds in *Nitraria sibirica* fruit, and reported the benefits of FEM over conventional extraction processes. Kan<sup>17</sup> used FEM to extract the seed oil of *Elaeagnus mollis*, which compared to the Soxhlet extraction method, improved oxidative stability in

<sup>a</sup>School of Functional Food and Wine, Shenyang Pharmaceutical University, Shenyang 110016, China. E-mail: zyq4885@126.com; xiaoshu2397@163.com; sunbaoshan@syphu.edu.cn; Fax: +86 24 43520300; Tel: +86 24 43520309; +86 24 43520301

<sup>b</sup>Key Laboratory of Structure-based Drug Design & Discovery, Ministry of Education, Shenyang Pharmaceutical University, Shenyang 110016, China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ra07517e

‡ Two authors contributed equally to this work.



volatile oil. Zhang<sup>18</sup> showed that using FEM, the structure of soybean flour could be protected from temperature induced damage, improving the analysis of its structural, physical and chemical properties. However, FEM has not been used for the rapid extraction and separation of ginseng.

High performance liquid chromatography (HPLC) and Fourier transform ion cyclotron resonance mass spectrometry (FT-ICR-MS) can provide accurate molecular mass and structural information due to their high sensitivity, resolution, and rapid scanning speed.<sup>19</sup> In this study, the efficient extraction of ginsenoside at room temperature was reported for the first time. The extraction conditions were optimized using response surface methodology (RSM). Further study, the semi preparative HPLC, MCI gels, and other chromatographic methods were used to isolate the components of ginsenosides in MCG. Meanwhile, more rigorous identification methods for ginseng (HPLC-FT-ICR-MS) were proposed and the ginsenosides in MCG and CG were qualitatively detected. We finally clarified the superior performance of MCG over CG, which provided a theoretical basis for the application of MCG in future studies.

## 2 Materials and methods

### 2.1 Reagents and materials

Semipreparative high performance liquid chromatography was carried out on an Agela Innoval ODS-2 column (5  $\mu$ m, 10  $\times$  250 nm, China) with an LC3000 instrument (Beijing Tong Heng Innovation Technology Co., Ltd. China). NMR spectra: Bruker ARX-600 spectrometer in pyridine-*d*<sub>5</sub> with trimethylchlorosilane as internal standard.

The flash extractor was obtained by All Herbal Sciencetech, LLC (JHBE-20A, Beijing, China). HPLC-FT-ICR-MS was performed with auto-sampler, thermostatic column chamber, online degasser, quadruple pump on Agilent 1260 system and diode-array detector (Agilent Corporation, Santa Clara, CA, USA) with ESI interface and a 7.0 tesla magnet. HPLC analyses were performed on CXTH LC-3000 (Beijing, China). Octadecyl silica chromatographic column [Agilent 5 HC-C<sub>18</sub> column (250  $\times$  4.6 mm)]. Chromatographic grade methanol, acetonitrile and purified water were purchased from Fisher Scientific International Inc. and Wahaha (Hangzhou, China) respectively.

Samples of MCG and CG were collected from Huanren, Liaoning Province, China. All of the plants were identified as 15 and 5 years ginseng by Prof. Jincai Lu of Shenyang Pharmaceutical University. A voucher specimen (PT 20190316) was deposited in the herbarium of Shenyang Pharmaceutical University. The samples were dried at 45  $^{\circ}$ C to constant weight.

### 2.2 Extraction procedures

All the samples were pulverized into powder of over 100 mesh. Weighed 1.0 g dried ginseng powder and mixed with different volume of ethanol–water solution were put in the flash extractor for extracting different time in different times. After extracting, the extract was filtered, and then the content of total ginsenoside in the extract was tested. In this study, the optimal extraction conditions were established through RSM.

### 2.3 Selection of variables

The total ginsenoside content of the MCG extract extracted by flash extractor is influenced by various factors, such as liquid–solid ratio, ethanol concentration, time of extraction and extraction voltage.<sup>18</sup> In our preliminary experiment, we found that the extraction time also affected the total content of ginsenoside in MCG extracts. To select the appropriate range of the variables, single factor experiment was carried out on the extraction time (1–5 min), ethanol concentration (10–90%), extraction times (1–5) and liquid–solid ratio (10–50 mL g<sup>−1</sup>) separately. The total ginsenoside was extracted under different conditions with one factor changed at a time and the other three remain the same.

### 2.4 The design of Box–Behnken

RSM method was used to optimize the extraction method of ginsenosides in MCG. According to the single factor experiments, the experimental variables were ethanol concentration ( $X_1$ , %), extraction times ( $X_2$ , times) and liquid–solid ratio ( $X_3$ , mL g<sup>−1</sup>). Each variable was encoded at three levels (−1, 0, 1), a Box–Behnken response surface experimental design (BBD) was applied which leading to 17 experiments. The regression coefficient is obtained by fitting the experimental data to the second-order polynomial model. The best condition was defined as the factor to maximize the yield of the reaction.

### 2.5 Total ginsenoside content of MCG

Total ginsenoside content of MCG was determined by the vanillin method. The extract solution (100  $\mu$ L) was mixed with 200  $\mu$ L 5% vanillin in acetic acid and 800  $\mu$ L 70% perchloric acid. The mixture was incubated at 60  $^{\circ}$ C for 15 min, and then 5 mL acetic acid was added and incubated at room temperature (25  $^{\circ}$ C) for 30 min. Ginsenoside Re solution with different concentration were used as the quantification standard. UV-2800AH spectrophotometer (UNICO, Shanghai) was used to measure the absorbance of the mixture.

### 2.6 Extraction and isolation

To obtain the ginsenosides, the dried MCG samples (80 g) were extracted by flash extractor with the optimal extraction conditions. The extract (15 g) was fractionated by silica gel with a gradient of CH<sub>2</sub>Cl<sub>2</sub>–MeOH–EtOAc–H<sub>2</sub>O (2 : 2 : 4 : 1) to obtain seven fractions (Fr. A–G). We further isolated the components of fractions by semi preparative HPLC, MCI gels, and other chromatographic methods. The schematic representation of the extraction and isolation procedure is shown in Fig. 1.

### 2.7 HPLC-FT-ICR-MS analysis

The HPLC-FT-ICR-MS were carried out on an Agilent 1260 system with a FT-ICR-MS equipped with ESI interface and a 7.0 tesla magnet. The HPLC mobile phase consisted of water (A) and acetonitrile (B) at 0.8 mL min<sup>−1</sup> flow through an Agilent 5 HC-C<sub>18</sub> column (250  $\times$  4.6 mm). Mobile phase elution system was performed in linear gradient (28–35% B at 0–5 min, 35–50% B at 5–15 min, 50–68% B at 15–25 min, 68–90% B at 25–28 min, 90% B at 28–40 min). The column temperature was kept at 25  $^{\circ}$ C, the UV detection wavelength was 203 nm, and the injection amount was 10  $\mu$ L.



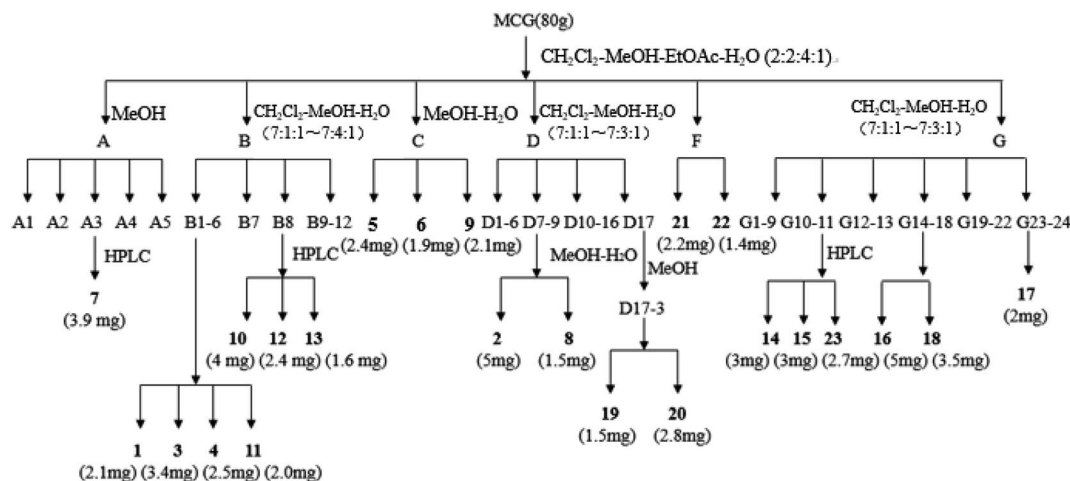


Fig. 1 MCG separation process.

The API source conditions were as follows: the ion source is ESI source; the capillary voltage is 4.0 kV, the end plate offset is  $-500$  V and the transfer capillary temperature is  $200$  °C. The source gas tune was as follows: the nebulizer gas pressure is  $1.0$  bar; the dry gas flow rate is  $1.2$  L  $\text{min}^{-1}$ ; the dry gas temperature is  $180$  °C. In the range of  $m/z$  value from  $150$  to  $2000$ , the full scan MS data are obtained, and the collision energy of the selected ion is initially set to  $10$  eV, then it is corrected according to the fragment. Under the soft ESI condition, most of the components showed negative ion mode in the full scan mass spectra. Calibrate the FT-ICR mass spectrometer in negative mode using NaTFA in accordance with the manufacturer's instructions. FT-MS control, Bruker Compass-Hystar and DataAnalysis Software (Bruker Daltonics, Bremen, Germany) were used to control the equipment, collect and analyze the data.

## 2.8 Simultaneous determination of individual ginsenosides

Analyses were performed on a CXTH-3000 HPLC system. The mobile phase consisted of water (A) and acetonitrile (B) at a flow rate of  $1.0$  mL  $\text{min}^{-1}$  through an Agilent 5 HC- $\text{C}_{18}$  column ( $250 \times 4.6$  mm). Mobile phase elution system was performed in a linear gradient (28–35% B at 0–5 min, 35–50% B at 5–15 min, 50–68% B at 15–25 min, 68–90% B at 25–28 min, 90% B at 28–40 min). The column temperature was kept at  $25$  °C, the UV detection wavelength was  $203$  nm, and the injection amount was  $20$   $\mu\text{L}$ .

## 2.9 Preparation of standard solutions and sample solutions

Compounds 1 to 23 were dissolved by HPLC grade methanol. 23 ginsenosides mixed stock solution were prepared by mixing 23 ginsenosides standard stock solution and diluting to scale with methanol (the concentrations of compounds 1 to 23 ( $\text{mg mL}^{-1}$ ): 0.01, 0.02, 0.02, 0.13, 0.10, 0.15, 0.18, 0.15, 0.18, 0.20, 0.19, 0.18, 0.15, 0.08, 0.13, 0.15, 0.15, 0.14, 0.20, 0.11, 0.10, 0.10, 0.14). All the solutions were stored at  $4$  °C until use.

Accurately  $0.5$  mg of MCG extract were weighed and dissolved in  $1$  mL of HPLC grade methanol, and then filtrated before use. A volume of  $10$   $\mu\text{L}$  was injected into the HPLC-FT-ICR-MS and HPLC system for further analysis.

## 2.10 Statistical analysis

All data were expressed as the mean  $\pm$  SD. Software Design Expert 8.05 was used to determine the linear, quadratic and interaction regression coefficients of RSM experiment.

# 3 Results and discussion

## 3.1 Extraction parameters

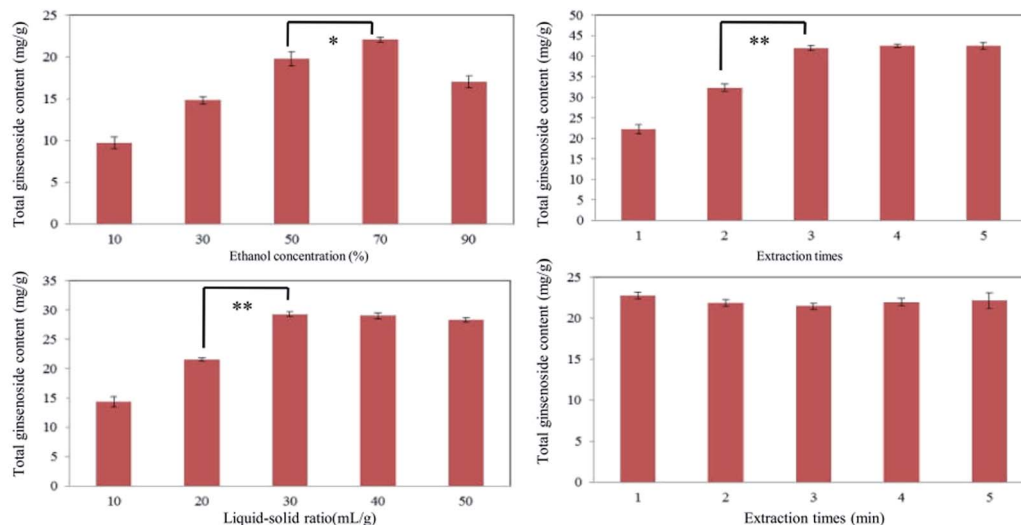
The total ginsenosides content of the ginseng extract extracted by FEM is influenced by various factors, including ethanol concentration, extraction time and liquid–solid ratio. In our preliminary study, extraction times were also affected by the total ginsenosides content in MCG extracts. The effects of extraction times, ethanol concentration, extraction time and liquid–solid ratio on the yield of total ginsenosides extract from MCG were preliminarily evaluated to determine the effective factors. As shown in Fig. 2A, excluding the extraction time, all parameters significant impacted the extraction rate of total ginsenosides in MCG. The total ginsenosides content was directly proportional to the liquid–solid ratio range from  $10$  to  $40$   $\text{mL g}^{-1}$ . The total ginsenosides content increased using 50% to 70% ethanol as the solvent, but did not significant differ from 70% to 90% ethanol. The extraction times also affected the extraction of total ginsenosides. Ultimately, the effective factors for latter response surface experiments were ethanol concentration (%),  $X_1$ , extraction times (times),  $X_2$  and liquid–solid ratio ( $\text{mL g}^{-1}$ ,  $X_3$ ) based on the experimental results, and their values were 60–80%, 3–5 times and 1 : 30–1:50, respectively.

## 3.2 Effects of the extraction variables on total ginsenosides content

In total, 17 experiments were performed for the analysis of variance for the fitted second-order regression model (Tables 1 and 2). According to the results, the model of experiment was adequate ( $p < 0.01$ ) and the lack of fit was not significant ( $p > 0.05$ ). The determination coefficient ( $R^2$ ) of the total ginsenosides was 0.9815, meaning 98.15% of the variations could be explained by the model. The model was highly significant and



A



B

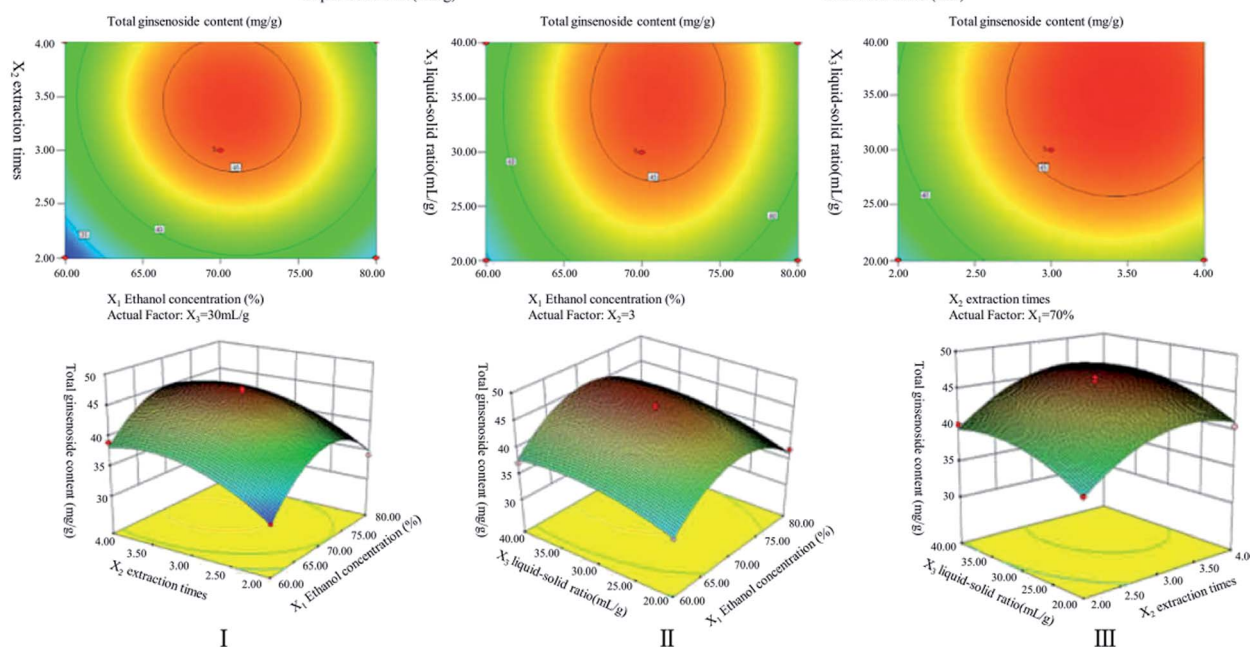


Fig. 2 Optimization of FEM process. (A) Effects of extraction ethanol concentration, extraction times, liquid–solid ratio and extraction time. (B) The 2D contour maps and 3D response surface of RSM showing the effect of extraction variables on total ginsenoside yields (I–III). \* $P < 0.05$ , \*\* $P < 0.01$ .

fitted the experimental data well. The positive linear effects of three independent variables on all response variables were significant ( $p < 0.05$ ), indicating that the model was highly significant and could fit the experimental data well. The positive linear effects of the three independent variables were significant ( $p < 0.05$ ) for all the response variables: the ethanol concentration ( $X_1$ ) ( $p < 0.05$ ), the extraction times ( $X_2$ ) and liquid–solid ratio ( $X_3$ ) ( $p < 0.01$ ) had significantly influenced the total ginsenoside content.

The second order polynomial model of ginsenoside was established by multiple regression analysis. According to the given coding factors, the mathematical regression model is as follows:

$$Y = 45.61 + 1.13X_1 + 2.74X_2 + 1.95X_3 - 0.49X_1X_2 + 0.31X_1X_3 + 0.16X_2X_3 - 6.29X_1^2 - 3.21X_2^2 - 1.94X_3^2.$$

### 3.3 Optimized conditions and experimental verification

The expected function method was used to optimize the maximum output of all three responses. Using the RSM generation model, the best experimental conditions for the maximum yield of all three reactions was 70.86% (ethanol content), 3.43 times (extraction times), and 35.29 : 1 mL g<sup>-1</sup> (liquid–solid ratio). The optimal levels of extraction included an ethanol content of 71%, extraction times of 3 and liquid–solid ratio of 36 : 1 mL g<sup>-1</sup>. To corroborate these results, five parallel assays were assessed under optimal levels. The average total





**Table 1** Regression coefficient, coefficient of determination ( $R^2$ ) and  $F$  value of the response surface quadratic models for total ginsenoside content<sup>a</sup>

| Source                  |          | Analytical results |
|-------------------------|----------|--------------------|
|                         |          | Total ginsenoside  |
| Intercept               |          | 45.61              |
| Linear                  | $X_1$    | 1.13**             |
|                         | $X_2$    | 2.74***            |
|                         | $X_3$    | 1.95***            |
| Quadratic               | $X_1^2$  | -6.29***           |
|                         | $X_2^2$  | -3.21***           |
|                         | $X_3^2$  | -1.94**            |
| Interactions            | $X_1X_2$ | -0.49              |
|                         | $X_1X_3$ | 0.31               |
|                         | $X_2X_3$ | 0.16               |
| $R^2$                   |          | 0.98               |
| Adj $R^2$               |          | 0.96               |
| $F$ Value (model)       |          | 41.36***           |
| $F$ Value (lack of fit) |          | 1.16               |

<sup>a</sup> \*\*\* indicates a statistically significant difference  $P < 0.001$ , \*\* indicates a significant statistical significance  $P < 0.01$ , \* indicates statistical significance  $P < 0.05$ .

ginsenosides content was  $45.01 \pm 0.1 \text{ mg g}^{-1}$ . The correlation between these results confirmed that the model fully reflected the expected optimization factors (Fig. 2B).

### 3.4 Phytochemical investigation

Twenty-three compounds (1–23) were obtained from the MCG. According to the structural of these compounds, the isolated compounds could be divided into two subtypes, including panaxadiol and panaxatriol. The structures of the compounds

were showed in Fig. 3A. Based on  $^{13}\text{C}$ -NMR data and comparison of known compound spectrum information, the compounds were determined as notoginsenoside  $R_1$  (1), 20-O-glu-ginsenoside  $R_f$  (2), ginsenoside  $R_f$  (3), ginsenoside  $R_e$  (4), ginsenoside  $R_{g_1}$  (5), 20(*S*)-notoginsenoside  $G$  (6), vinal-ginsenoside  $R_4$  (7), ginsenoside  $R_{b_1}$  (8), ginsenoside  $I_a$  (9), ginsenoside  $R_c$  (10), notoginsenoside  $F_z$  (11), ginsenoside  $R_{b_2}$  (12), ginsenoside  $R_{b_3}$  (13), 20(*S*)-ginsenoside  $R_{g_2}$  (14), 20(*R*)-ginsenoside  $R_{g_2}$  (15), 20(*R*)-ginsenoside  $R_{h_1}$  (16), ginsenoside  $F_1$  (17), notoginsenoside  $F_e$  (18), ginsenoside  $R_{d_2}$  (19), ginsenoside  $R_{g_6}$  (20), ginsenoside  $R_{k_3}$  (21), ginsenoside  $R_{g_3}$  (22) and 20(*S*)-ginsenoside  $R_{h_2}$  (23). The structure analysis and spectrums of all compounds are in ESI.†

Amongst the compounds, compound 1 (notoginsenoside  $R_1$ ) was regarded as the characteristic components of *Panax notoginseng*. Zhu<sup>20</sup> used UPLC-QTOF-MS/MS/MS and Unifi new information platform to rapidly analyze and identify the small and medium molecular components (molecular weight 100–1200 Da) of MCG and CG. Notoginsenoside  $R_1$  was detected in MCG and CG, but not isolated from MCG. This was the first study to isolate notoginsenoside  $R_1$  from MCG.

### 3.5 HPLC-FT-ICR-MS determination of individual ginsenoside

Compounds were analyzed using the by HPLC-FT-ICR-MS method for qualitative analysis. In total, 23 ginsenosides were identified from MCG. Gradient elutions performed to increase the resolution of HPLC analysis due to complexity of the components contained in MCG. Upon comparison of acetonitrile–water and methanol–water as mobile phases, the separation effect of the acetonitrile–water system performed to a higher level.

To determine the conditions for MS analysis, we tested both positive and negative ionization modes. The results showed that

**Table 2** The conditions and results of 17 experiments in response surface methodology experiments

| Run | Factors                         |                        |                                                 | Experimental values                              |
|-----|---------------------------------|------------------------|-------------------------------------------------|--------------------------------------------------|
|     | $X_1$ ethanol concentration (%) | $X_2$ extraction times | $X_3$ liquid–solid ratio ( $\text{mL g}^{-1}$ ) | Total ginsenoside content ( $\text{mg g}^{-1}$ ) |
| 1   | 80                              | 3                      | 20                                              | $36.96 \pm 0.34$                                 |
| 2   | 70                              | 4                      | 40                                              | $45.25 \pm 1.23$                                 |
| 3   | 60                              | 3                      | 40                                              | $37.19 \pm 0.56$                                 |
| 4   | 80                              | 4                      | 30                                              | $39.43 \pm 1.89$                                 |
| 5   | 80                              | 2                      | 30                                              | $34.23 \pm 0.67$                                 |
| 6   | 80                              | 3                      | 40                                              | $40.92 \pm 1.87$                                 |
| 7   | 60                              | 3                      | 20                                              | $34.49 \pm 0.65$                                 |
| 8   | 70                              | 3                      | 30                                              | $45.22 \pm 0.87$                                 |
| 9   | 60                              | 4                      | 30                                              | $38.98 \pm 0.45$                                 |
| 10  | 70                              | 3                      | 30                                              | $44.33 \pm 1.86$                                 |
| 11  | 70                              | 3                      | 30                                              | $46.20 \pm 2.01$                                 |
| 12  | 70                              | 3                      | 30                                              | $46.77 \pm 1.45$                                 |
| 13  | 70                              | 2                      | 40                                              | $40.15 \pm 1.09$                                 |
| 14  | 60                              | 2                      | 30                                              | $31.81 \pm 0.54$                                 |
| 15  | 70                              | 4                      | 20                                              | $40.45 \pm 1.65$                                 |
| 16  | 70                              | 2                      | 20                                              | $36.00 \pm 1.21$                                 |
| 17  | 70                              | 3                      | 30                                              | $45.55 \pm 1.69$                                 |



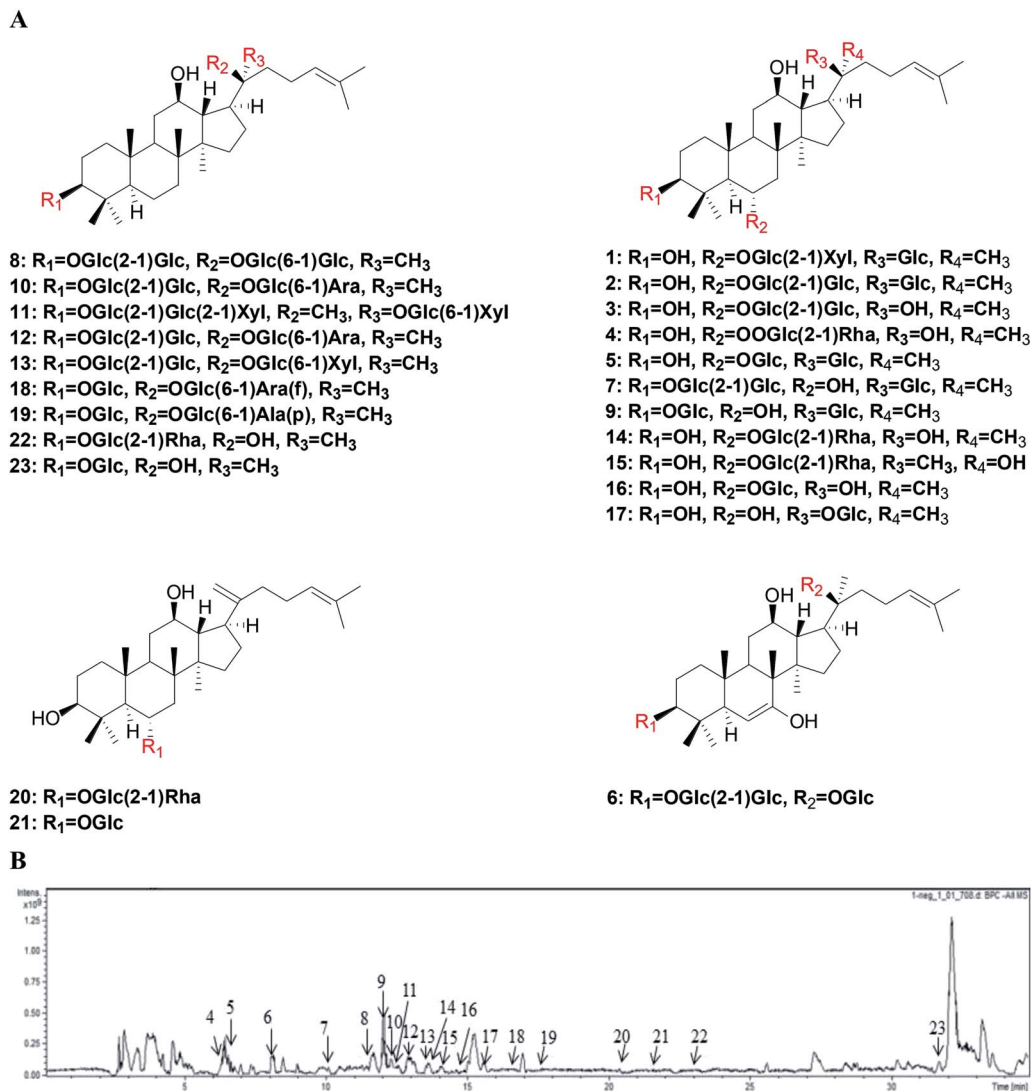


Fig. 3 Separation and HPLC-FT-ICR MS identification of MCG. (A) The structures of 23 compounds isolated from MCG. (B) The HPLC-FT-ICR MS chromatograms of MCG.

the negative ionization mode ( $[M - H]^-$  peak) provided more intense proton signals. The peak information for HPLC-FT-ICR-MS analysis is provided in Table 3 and the HPLC-FT-ICR-MS chromatograms of MCG along with the HPLC-FT-ICR-MS spectra are shown in Fig. 3B. The 23 tested compounds were assessed by HPLC from mixed stock solutions and mass spec analysis. A total of 20 samples in the reference substance showed a mass spectrum response. Upon comparison of the liquid phase and mass spectrometry information of the sample and reference substance, 20 ginsenosides were accurately identified, whilst 3 were undetected due to their low content.

### 3.6 HPLC analysis of individual ginsenoside

The results of linearity, precision, accuracy and stability experiments are shown in Table 4.  $Rg_1$  and  $Re$  in MCG are 0.342% (not less than 0.27% according to Chinese Pharmacopoeia), the content was 1.47 times of CG;  $Rb_1$  in MCG was 0.24% (not less than 0.18% according to Chinese

Pharmacopoeia), the content was 1.28 times of CG. The total content of all ginsenosides in MCG was twice that of CG. The content of rare ginsenosides  $Rg_3$ , 20(*R/S*)- $Rg_2$ ,  $Rk_3$ , 20(*S*)- $Rh_2$ , 20(*R*)- $Rh_1$ ,  $F_1$  in MCG were 2.5, 3.7, 1.7, 33, 24 and 5.5-fold higher than CG, respectively. The content of ginsenoside  $Rg_6$  in MCG was  $0.01 \text{ mg g}^{-1}$ , which was not detected in CG. Rare ginsenoside  $Rg_3$  is the main component of "Shenyi capsule", which is listed in China.  $Rh_2$  is the metabolite and transformation product of  $Rg_3$ .<sup>21</sup> Its anti-tumor activity is four-fold higher than that of  $Rg_3$ .<sup>22</sup> Rare ginsenosides  $Rg_2$ ,  $Rh_1$ ,  $Rk_3$  represent the transformation or metabolites of ginsenoside  $Re$ ,<sup>23,24</sup> have been touted as new 'anti-neoplastic pharmacophores', with improved bioavailability and potency compared to the prototype ginsenosides.<sup>25</sup>

## 4 Conclusion

In this study, FEM protocol was used to extract ginsenosides from MCG. In total, 23 ginsenosides were isolated from the



Table 3 The peak information of HPLC-FT-ICR MS analysis

| No. | Ginsenoside |            |       | $[M - H]^-$          | MCG   |            |       | Comp.                             |
|-----|-------------|------------|-------|----------------------|-------|------------|-------|-----------------------------------|
|     | $t_R$       | $m/z$      | Error |                      | $t_R$ | $m/z$      | Error |                                   |
| 1   | 6.28        | 945.54858  | -0.26 | $C_{48}H_{81}O_{18}$ | 6.34  | 945.54832  | -1.21 | Ginsenoside Re                    |
| 2   | 6.46        | 799.49041  | -1.70 | $C_{42}H_{71}O_{14}$ | 6.67  | 799.49041  | 1.18  | Ginsenoside Rg <sub>1</sub>       |
| 3   | 7.98        | 959.52758  | 3.51  | $C_{48}H_{79}O_{19}$ | 8.05  | 959.52758  | 2.62  | 20(S)-Notoginsenoside G           |
| 4   | 9.42        | 961.54323  | 1.80  | $C_{48}H_{81}O_{19}$ | 10.04 | 961.54323  | -4.00 | Vina-ginsenoside R <sub>4</sub>   |
| 5   | 11.53       | 1107.60114 | 4.22  | $C_{54}H_{91}O_{23}$ | 11.59 | 1107.60114 | 3.12  | Ginsenoside Rb <sub>1</sub>       |
| 6   | 12.09       | 799.49041  | 0.22  | $C_{42}H_{71}O_{14}$ | 12.01 | 799.49041  | -2.36 | Ginsenoside Ia                    |
| 7   | 12.33       | 1077.59058 | 2.25  | $C_{53}H_{89}O_{22}$ | 12.30 | 1077.59058 | 4.65  | Ginsenoside Rc                    |
| 8   | 12.64       | 1209.63284 | 0.22  | $C_{58}H_{97}O_{26}$ | 12.59 | 1209.63284 | 1.89  | Notoginsenoside Fz                |
| 9   | 12.94       | 1077.59058 | 3.98  | $C_{53}H_{89}O_{22}$ | 12.91 | 1077.59058 | 0.41  | Ginsenoside Rb <sub>2</sub>       |
| 10  | 13.56       | 1077.59180 | -1.09 | $C_{53}H_{89}O_{22}$ | 13.40 | 1077.59058 | -1.09 | Ginsenoside Rb <sub>3</sub>       |
| 11  | 13.94       | 783.49550  | 0.53  | $C_{42}H_{71}O_{13}$ | 13.79 | 783.49550  | -0.59 | 20(S)-Ginsenoside Rg <sub>2</sub> |
| 12  | 14.11       | 783.49550  | -1.03 | $C_{42}H_{71}O_{13}$ | 14.08 | 783.49550  | -2.09 | 20(R)-Ginsenoside Rg <sub>2</sub> |
| 13  | 14.56       | 637.43759  | -1.93 | $C_{36}H_{61}O_9$    | 14.63 | 637.43759  | -0.61 | 20(R)-Ginsenoside Rh <sub>1</sub> |
| 14  | 15.41       | 637.43759  | 0.62  | $C_{36}H_{61}O_9$    | 15.72 | 637.43759  | -1.51 | Ginsenoside F <sub>1</sub>        |
| 15  | 16.72       | 915.53775  | -0.03 | $C_{47}H_{79}O_{17}$ | 16.72 | 915.53775  | -2.37 | Notoginsenoside Fe                |
| 16  | 17.41       | 779.46420  | -1.41 | $C_{42}H_{67}O_{13}$ | 17.50 | 779.46420  | 2.51  | Ginsenoside Rd <sub>2</sub>       |
| 17  | 20.28       | 751.46928  | 0.19  | $C_{41}H_{67}O_{12}$ | 20.34 | 751.46928  | -1.37 | Ginsenoside Rg <sub>6</sub>       |
| 18  | 21.57       | 619.42702  | -0.54 | $C_{36}H_{59}O_8$    | 21.66 | 619.42702  | -1.42 | Ginsenoside Rk <sub>3</sub>       |
| 19  | 22.90       | 782.49550  | -0.91 | $C_{42}H_{71}O_{13}$ | 22.96 | 782.49550  | 0.18  | Ginsenoside Rg <sub>3</sub>       |
| 20  | 31.56       | 621.44267  | -1.08 | $C_{36}H_{61}O_8$    | 31.55 | 621.44267  | -1.60 | 20(S)-Ginsenoside Rh <sub>2</sub> |

Table 4 The methodology investigation results of simultaneous determination of ginsenosides

|           |                                     | Liner regression data              |                |                              | Precision RSD (%) |           |           | Content (mg g <sup>-1</sup> ) |             |                |
|-----------|-------------------------------------|------------------------------------|----------------|------------------------------|-------------------|-----------|-----------|-------------------------------|-------------|----------------|
| No. Comp. |                                     | Liner                              | R <sup>2</sup> | Range (mg mL <sup>-1</sup> ) | Intra-day         | Inter-day | Stability | RSD (%)                       | MCG         | Garden ginseng |
| 1         | Ginsenoside Re/Rg <sub>1</sub>      | Y = 5 × 10 <sup>6</sup> X + 11 560 | 0.9995         | 0.02–2.33                    | 2.28              | 1.44      | 2.54      |                               | 3.42 ± 0.19 | 2.33 ± 0.21    |
| 2         | 20(S)-Notoginsenoside G             | Y = 6 × 10 <sup>6</sup> X + 9935.1 | 0.9994         | 0.02–1.50                    | 2.16              | 2.30      | 1.52      |                               | 0.79 ± 0.01 | 0.10 ± 0.01    |
| 3         | Vina-ginsenoside R <sub>4</sub>     | Y = 7 × 10 <sup>6</sup> X + 1798.9 | 0.9999         | 0.02–1.82                    | 1.95              | 1.92      | 1.62      |                               | 0.38 ± 0.01 | 0.12 ± 0.07    |
| 4         | Ginsenoside Rb <sub>1</sub>         | Y = 7 × 10 <sup>6</sup> X + 18 577 | 0.9992         | 0.01–1.45                    | 2.21              | 1.42      | 2.46      |                               | 2.17 ± 0.12 | 1.69 ± 0.14    |
| 5         | Ginsenoside Ia                      | Y = 4 × 10 <sup>6</sup> X – 4337.3 | 0.9994         | 0.02–1.86                    | 1.04              | 2.77      | 2.52      |                               | 0.28 ± 0.02 | 0.02 ± 0.00    |
| 6         | Ginsenoside Rc                      | Y = 1 × 10 <sup>6</sup> X – 13 539 | 0.9991         | 0.02–1.96                    | 1.68              | 2.58      | 1.23      |                               | 0.28 ± 0.03 | 0.04 ± 0.01    |
| 7         | Notoginsenoside Fz                  | Y = 1 × 10 <sup>7</sup> X – 14 124 | 0.9991         | 0.02–1.85                    | 1.95              | 1.54      | 1.52      |                               | 0.04 ± 0.00 | 0.02 ± 0.00    |
| 8         | Ginsenoside Rb <sub>2</sub>         | Y = 2 × 10 <sup>6</sup> X – 2579.4 | 0.9997         | 0.02–1.77                    | 1.19              | 1.31      | 2.49      |                               | 0.05 ± 0.01 | 0.04 ± 0.01    |
| 9         | Ginsenoside Rb <sub>3</sub>         | Y = 2 × 10 <sup>6</sup> X – 2128.4 | 0.9998         | 0.01–1.48                    | 1.39              | 2.73      | 2.74      |                               | 1.27 ± 0.11 | 0.14 ± 0.03    |
| 10        | 20(R/S)-Ginsenoside Rg <sub>2</sub> | Y = 5 × 10 <sup>6</sup> X + 3572.7 | 0.9994         | 0.02–2.17                    | 2.13              | 2.29      | 1.64      |                               | 0.11 ± 0.02 | 0.03 ± 0.00    |
| 11        | 20(R)-Ginsenoside Rh <sub>1</sub>   | Y = 3 × 10 <sup>6</sup> X – 3519.3 | 0.9996         | 0.01–1.46                    | 1.05              | 1.19      | 2.40      |                               | 0.24 ± 0.06 | 0.01 ± 0.00    |
| 12        | Ginsenoside F <sub>1</sub>          | Y = 8 × 10 <sup>6</sup> X – 435.51 | 0.9996         | 0.01–1.46                    | 2.22              | 2.74      | 2.65      |                               | 0.11 ± 0.03 | 0.02 ± 0.00    |
| 13        | Notoginsenoside Fe                  | Y = 5 × 10 <sup>6</sup> X – 3287.5 | 0.9996         | 0.01–1.41                    | 1.44              | 2.84      | 1.79      |                               | 0.05 ± 0.01 | 0.03 ± 0.01    |
| 14        | Ginsenoside Rd <sub>2</sub>         | Y = 3 × 10 <sup>6</sup> X – 3709.4 | 0.9990         | 0.02–1.97                    | 1.60              | 1.47      | 2.09      |                               | 0.03 ± 0.00 | 0.01 ± 0.00    |
| 15        | Ginsenoside Rg <sub>6</sub>         | Y = 6 × 10 <sup>6</sup> X – 981.37 | 0.9999         | 0.01–1.07                    | 2.42              | 2.56      | 2.43      |                               | 0.01 ± 0.00 | —              |
| 16        | Ginsenoside Rk <sub>3</sub>         | Y = 1 × 10 <sup>7</sup> X – 6306.3 | 0.9998         | 0.01–1.02                    | 1.73              | 1.01      | 2.35      |                               | 0.05 ± 0.01 | 0.03 ± 0.00    |
| 17        | Ginsenoside Rg <sub>3</sub>         | Y = 8 × 10 <sup>6</sup> X + 3700.7 | 0.9994         | 0.01–1.04                    | 2.62              | 2.10      | 2.79      |                               | 0.05 ± 0.01 | 0.02 ± 0.00    |
| 18        | 20(S)-Ginsenoside Rh <sub>2</sub>   | Y = 6 × 10 <sup>6</sup> X – 4268.1 | 0.9999         | 0.01–1.36                    | 1.90              | 1.13      | 2.53      |                               | 0.33 ± 0.08 | 0.01 ± 0.00    |

MCG and were determined by HPLC-FT-ICR-MS. We isolated notoginsenoside R<sub>1</sub> for the first time in MCG. The simultaneous determination of 20 ginsenosides in MCG and CG by HPLC-UV showed that the ginsenoside content in MCG was significantly higher than that in CG. The content of rare ginsenosides Rg<sub>3</sub>, 20(R/S)-Rg<sub>2</sub>, Rk<sub>3</sub>, 20(S)-Rh<sub>2</sub>, 20(R)-Rh<sub>1</sub>, F<sub>1</sub> and Rg<sub>6</sub> in MCG were higher than those of CG, respectively. The FEM not only provided a simple, feasible, quick and efficient sample processing method for the quality analysis of MCG and CG, but

avoided the unstable or uncontrollable quality caused by the thermal transformation or decomposition of the prototype ginsenosides. The discovery of high content of rare ginsenosides in MCG provides characteristic detection indices for quality evaluation. This provides an important theoretical basis for its significant biological activities including anti-tumor and immune enhancement of MCG and its clinical medicinal and health care value.



## Conflicts of interest

There is no conflict of interests regarding the publication of this paper.

## Acknowledgements

The research was supported by National Natural Science Foundation of China (81903518 and 81703389), Doctoral Research Startup Fund Item of Liaoning Province (2019-BS-232), Liaoning (FGW) Engineering Technology Research Center for industrial chromatographic preparation of natural innovative drugs materials (2017-1007) and Technology Platform of Industrialization Chromatographic Preparation for Standard Extract of Traditional Chinese Medicine (2010ZX09401-304-105B).

## References

- 1 G. Y. Hou, J. Niu, F. R. Song, Z. Q. Liu and S. Y. Liu, *J. Chromatogr. B*, 2013, **923**–**924**, 1–7.
- 2 T. Li, W. Sun, X. Q. Dong, W. H. Yu, J. Y. Cai, Q. Yuan, L. Shan and T. Efferth, *Oncol. Lett.*, 2018, **16**, 4640–4648.
- 3 L. L. Jiao, X. Y. Zhang, B. Li, Z. Liu, M. Z. Wang and S. Y. Liu, *Int. J. Biol. Macromol.*, 2014, **65**, 229–233.
- 4 B. Zhao, X. Wang, H. Liu, C. Lv and J. Lu, *Int. J. Biol. Macromol.*, 2020, **150**, 737–745.
- 5 H. S. Lee, M. R. Kim, Y. Park, H. J. Park, Y. J. Chang, S. Y. Kim and H. J. Suh, *J. Med. Food*, 2012, **15**, 1015–1023.
- 6 S. Lee, K. Youn and M. Jun, *Food Funct.*, 2018, **9**, 4122–4134.
- 7 J. Xu, X. Wang, G. Su, J. Yue, Y. Sun, J. Q. Cao, X. S. Zhang and Y. Zhao, *J. Funct. Foods*, 2018, **46**, 567–578.
- 8 Y. Q. Zhao, L. J. Wu, C. L. Yuan, X. Li and J. Chen, *Chin. Chem. Lett.*, 1992, **3**, 887–888.
- 9 Y. Q. Zhao, W. Wang, E. Rayburn, D. L. Hill, H. Wang and R. W. Zhang, *Med. Chem.*, 2007, **3**, 51–60.
- 10 H. Fan, K. Li, F. Yao, L. W. Sun and Y. J. Liu, *BMC Plant Biol.*, 2019, **19**, 82.
- 11 D. Xiao, H. Yue, Y. Xiu, X. L. Sun, Y. B. Wang and S. Y. Liu, *J. Ginseng Res.*, 2015, **39**, 338–344.
- 12 *The Pharmacopoeia of the People's Republic of China*, 2015.
- 13 X. Huang, Y. Liu, Y. Zhang, S. P. Li, H. Yue, C. B. Chen and S. Y. Liu, *J. Ginseng Res.*, 2019, **43**, 27–37.
- 14 H. D. Song, R. J. Yang, W. Zhao, W. Katiyo, X. Hua and W. B. Zhang, *J. Agric. Food Chem.*, 2014, **62**, 3802–3812.
- 15 J. Liu, C. Liu, Y. H. Rong, L. Rong and G. L. Huang, *Solvent Extr. Res. Dev.*, 2012, **19**, 137–145.
- 16 Q. B. Song, X. Xia, C. M. Ji, D. F. Chen and Y. Lu, *J. Pharm. Biomed. Anal.*, 2019, **172**, 379–387.
- 17 L. Kan, L. Wang, Q. Ding, Y. Wu and J. Ouyang, *J. Oleo Sci.*, 2017, **66**, 345–352.
- 18 Y. P. Zhang, R. J. Yang and W. N. Zhang, *Food Chem.*, 2017, **219**, 48–53.
- 19 H. Suo, R. C. Tian, J. Li, S. T. Zhang, Y. Cui, L. X. Li and B. S. Sun, *Food Res. Int.*, 2019, **123**, 440–449.
- 20 H. Zhu, H. Lin, J. Tan, C. Wang, H. Wang, F. Wu, Q. Dong, Y. Lin, P. Li and J. Liu, *Molecules*, 2019, **224**, 33–54.
- 21 E. A. Bae, M. J. Han, M. K. Choo, S. Y. Park and D. H. Kim, *Biol. Pharm. Bull.*, 2002, **25**, 58–63.
- 22 J. H. Cheong, H. Kim, M. J. Hong, M. H. Yang, J. W. Kim, H. Yoo, H. Yang, J. H. Park, S. H. Sung, H. P. Kim and J. Kim, *Pharm. Bull.*, 2015, **38**, 102–108.
- 23 H. Y. Wang, C. Wang, S. C. Guo, Z. C. Chen, Z. T. Peng, R. Duan, T. T. X. Dong and K. W. K. Tsim, *J. Pharm. Biomed. Anal.*, 2019, **175**, 112779.
- 24 J. Han, J. Xia, L. Zhang, E. Cai, Y. Zhao, X. Fei, X. Jia, H. Yang and S. Liu, *J. Ginseng Res.*, 2019, **43**, 618–624.
- 25 T. Biswas, A. K. Mathur and A. Mathur, *Appl. Microbiol. Biotechnol.*, 2017, **101**, 4009–4032.

