Research Article

Study on the Spectrum-Effect Correlation of Anti-Inflammatory Active Extract of Sauropus spatulifolius Beille

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Sauropus spatulifolius Beille (S. spatulifolius) is a commonly used medicine of the Bourau and Yao nationalities. However, the composition of S. spatulifolius is complex, and simple chemical fingerprints cannot accurately evaluate the relationship between its composition and efficacy. In this study, high-performance liquid chromatography (HPLC) method was used to establish the fingerprint of the ethyl acetate extract of S. spatulifolius. Based on the evaluation of the similarity of chromatographic fingerprints of traditional Chinese medicine, combined with cluster analysis and principal component analysis (PCA), the common peaks of fingerprints were evaluated. The anti-inflammatory effect data were extracted through the dimethylbenzene-induced ear-swelling model in mice. The gray relational analysis (GRA) combined with partial least squares regression (PLSR) was used to study the spectrum-effect correlation of S. spatulifolius. As a result, the HPLC fingerprint of the ethyl acetate extract of S. spatulifolius was established, and 18 common peaks were identified. Except for S6, the other similarities are all above 0.915. The reference substance control method was used to identify two absorption peaks, namely, protocatechuic acid and caffeic acid. The cluster analysis results showed that 10 samples from different origins were grouped into four categories, which was consistent with the PCA results. Ethyl acetate extract of 10 batches of S. spatulifolius could significantly inhibit the ear swelling of mice (P < 0.01). Through GRA, the order of the contribution of each chemical component to the anti-inflammatory efficacy was obtained. The results of PLSR showed that the VIP values of peaks 3, 4, and 12 were greater than 1 and were positively correlated with anti-inflammatory activity. In this study, the HPLC fingerprint of the ethyl acetate extract of S. spatulifolius was established. Through the study of the spectrum-effect correlation, the anti-inflammatory active substance of the ethyl acetate extract of S. spatulifolius was obtained. The anti-inflammatory effect of S. spatulifolius was the result of the joint action of multiple ingredients. This research helps to quickly and accurately discover the active ingredient groups of traditional Chinese medicine and provides new ideas and methods for studying the effective substances of traditional Chinese medicine.

1. Introduction

Medicinal plants are found nearly throughout the world and have been used for therapeutic, religious, cosmetic, nutritional, and beautification purposes since ancient times, and the humanity of all civilizations and culture are familiar with their usages. They have been recognized for their human health benefits also. They have high content of nonnutritive, nutritive, and bioactive compounds, such as flavonoids, phenolics, anthocyanins, and phenolic acids, which have potent antioxidant, anticancer, and antimutagenic effects [1–4]. Sauropus spatulifolius Beille (S. spatulifolius) is a plant of the Euphorbiaceae. It can be picked all year round or collected in summer and autumn. It is better to pick the green old leaves and dry them in the sun [5]. S. spatulifolius is a small evergreen shrub that mainly grows in valleys,
mountain slopes, and moist and fertile jungles, distributed in Guangdong, Guangxi, Fujian, Hainan, and other regions [6]. The whole plant of *S. spatulifolius* can be used as medicine. In traditional medicine, the mature leaves of *S. spatulifolius* are generally used as medicine [7]. In clinical practice, *S. spatulifolius* is mainly used for clearing away heat and moisturizing lungs, resolving phlegm and relieving cough, improving pharynx, moistening intestines, and laxatives. *S. spatulifolius* mainly treats diseases such as lung heat, cough, spumt, constipation, dry mouth, sore throat, bronchitis, and bronchial asthma [5–8]. Modern pharmacological studies have shown that *S. spatulifolius* has anti-inflammatory and analgesic [9], antiallergic [10], antioxidation [11–13], antibacterial, antitussive, and expectorant effects [14,15].

Traditional Chinese medicine composition is complex, and its effect is usually the result of many components, many targets, and many ways. Traditional Chinese medicine fingerprint is a comprehensive and quantifiable identification method, which plays an important role in the identification and quality control of traditional Chinese medicine [16,17]. Although fingerprint is a widely accepted method for quality evaluation of traditional Chinese medicine in China, the simple chemical fingerprint cannot accurately evaluate the relationship between the composition and efficacy of traditional Chinese medicine. Therefore, it is necessary to change the traditional evaluation method based on chemical composition into one based on biological effect. The spectrum-effect correlation is a method to study the active components and quality control of traditional Chinese medicine. Through various mathematical model methods, the relation between fingerprint characteristic peak and drug effect is established to find out the substance that really has drug effect. This method makes up for the deficiency of separating the chemical composition from the medicinal effect. It provides a reliable method to elucidate the material basis of traditional Chinese medicine [18–20]. At present, the spectrum-effect relationship has been widely used in traditional Chinese medicine research, including the study of single medicinal substance of traditional Chinese medicine [21], medicinal substance of compound preparation [22, 23], compatibility of components [24], processing mechanism [25], and process optimization [26], providing more scientific basis for quality control of traditional Chinese medicine. It has made important contributions to its internationalization and globalization spread.

*S. spatulifolius* is a commonly used folk Zhuang Yao medicine. Now, it was included in Chinese pharmacopoeia [27]. Studies on quality control are mainly conducted by means of microscopic identification, spectral and chromatographic identification, heavy metal and pesticide inspection, and content determination, etc [28–33]. The fingerprint and spectrum-effect correlation of ethyl acetate extract of *S. spatulifolius* were not publicly reported. Therefore, high-performance liquid chromatography (HPLC) was used to establish the fingerprint effect of ethyl acetate extract of *S. spatulifolius*, and anti-inflammatory activity was used as a point of penetration. The correlation between common peaks and anti-inflammatory activity was studied by gray relational analysis (GRA) and partial least squares regression (PLSR) to further reveal the material basis of the anti-inflammatory effect of ethyl acetate extract of *S. spatulifolius*, in order to provide methods and basis for further research and quality control of *S. spatulifolius*.

2. Materials and Methods

2.1. Samples, Reagents, and Animals. The 10 batches of *S. spatulifolius* used in the experiment were all collected from Guangxi and Guangdong provinces. The plant was identified by Ma Life Deputy Chief Pharmacist (Deputy Chief Pharmacist of Guangxi Xixin Pharmaceutical Co., Ltd.; see Table 1 for details). Voucher specimens were preserved in the Department of Drug Analysis, Guangxi University of Chinese Medicine (No. 20180501). Methanol and acetonitrile were of chromatographic grade, purchased from Thermo Fisher Scientific Co., Ltd. (Massachusetts, USA); water was ultrapure; Tween-80 was of chemical grade, purchased from Tianjin Fuyu Fine Chemical Co., Ltd. (Tianjin, China, No: 20180928). Edible alcohol was of ordinary grade, purchased from Guangxi Haiying Alcohol Co., Ltd. (Nanning, China, No: 2016060701). Petroleum ether, ethyl acetate, n-butanol, glacial acetic acid, phosphoric acid, and xylene were all of analytical grades. Caffeic acid reference substance was purchased from China Institute for the Control of Pharmaceutical and Biological Products (Beijing, China, No: 110885-201703). The protocatechuc acid reference substance was purchased from China Institute for the Control of Pharmaceutical and Biological Products (Beijing, China, No: 111530-200303). Dexamethasone acetate tablets were purchased from Zhejiang Xianju Pharmaceutical Co., Ltd. (Zhejiang, China, No.: 170512). Sodium chloride injection was purchased from Huiyinbi Group Jiangxi East Asia Pharmaceutical Co., Ltd. (Jiangxi, China, No: 2018112215).

KM mice (SPF grade) were supplied by Hunan Slike Jingda Laboratory Animal Co., Ltd. (approval No: SCXK (Xiang) 2016-0002. Changsha, Hunan, China). The Animal Ethics Committee of Guangxi University of Chinese Medicine approved all animal protocols. The animal experiments were carried out according to the Guide for the Care and Use of Laboratory Animals.

2.2. Apparatus and Conditions. The following apparatus and conditions were used: Waters e2695 high-performance liquid chromatograph (Waters 2489UV detector; Waters Corporation, USA); Thermo Syncronis C18 column (4.6 mm × 250 mm, 5 μm); column temperature, 25°C; and detection wavelength, 260 nm. The flow rate was 0.8 mL·min⁻¹. The injection volume was 10 μL. The mobile phase was a mixture of methanol (A) and 0.1% glacial acetic acid (C). The gradient elution: 0–30 min, 15–18% A; 30–55 min, 18–30% A; 55–75 min, 30–35% A; 75–90 min, 35–38% A; 90–95 min, 38–15% A.

2.3. Preparation of Sample Solution. Preparation of extracts from different polar parts of *S. spatulifolius* is as follows: take 750 g dried leaves of *S. spatulifolius* and cut into fragments;
add seven times 75% ethanol to soak for 4 d. Percolated extraction until the color of the extract becomes pale, the ethanol solvent was recovered, and the concentrated extract liquid was evaporated to dryness in a water bath to obtain a 75% ethanol total extract. The total extract of 75% ethanol was dissolved with appropriate amount of distilled water and petroleum ether, ethyl acetate, and n-butanol solvents used in the extraction were recovered. The concentrated extract was dissolved with appropriate amount of distilled water and 75% ethanol total extract. The total extract of 75% ethanol was evaporated to dryness in a water bath to obtain a 75% ethanol total extract. The total extract of 75% ethanol was dissolved with appropriate amount of distilled water and petroleum ether, ethyl acetate, and n-butanol solvents used in the extraction were recovered. The concentrated extract liquid was evaporated to dryness in a water bath kettle and stored in the desiccator for subsequent experiments.

Samples solution for the HPLC fingerprinting analysis is prepared as follows: 0.1 g of ethyl acetate extract of S. spatulifolius was (equivalent to 4.77 g of crude drugs) precisely weighed; it was dissolved by ultrasound and transferred to a 2 mL volumetric flask. Methanol was added to the volumetric flask to scale and centrifuged 10 min at a speed of 13,000 r·min⁻¹. The supernatant was taken as the sample solution for HPLC analysis.

Mixed reference solution: appropriate amounts of caffeic acid and protocatechuic acid were precisely weighed and placed in 5 mL volumetric flasks and added methanol to scale, and then served as the reference stock solution of caffeic acid and protocatechuic acid. Appropriate amounts of caffeic acid reference stock solution and protocatechuic acid reference stock solution were precisely removed and placed in a volumetric flask of 2 mL, and added methanol to scale. Centrifuged 10 min at a speed of 13,000 r·min⁻¹. The supernatant was taken as the mixed reference solution (containing 159.96 μg/mL⁻¹ caffeic acid and 99.43 μg/mL⁻¹ protocatechuic acid).

\[
\text{degree of swelling} = \frac{\text{right ear piece weight} - \text{left ear piece weight}}{\text{left ear piece weight}}
\]

\[
\text{swelling inhibition rate} = \frac{(\text{average degree of swelling in the model group} - \text{average degree of swelling in the drug group})}{\text{average degree of swelling in the model group}} \times 100%.
\]

2.6. Data Handling. "Chinese traditional medicine chromatographic fingerprint similarity evaluation system (2012, 1 Edition)" was used to evaluate 10 batches of AIA data of ethyl acetate extract of S. spatulifolius. IBM SPSS Statistics 22.0 software was used to perform systematic clustering analysis on the original peak areas of 18 common peaks in the fingerprints of ethyl acetate extract of 10 batches of S. spatulifolius. Cluster analysis was performed using between-groups linkage, and the distance between samples was calculated using squared Euclidean distance. IBM SPSS Statistics 22.0 software was used to carry out principal component analysis (PCA) on the peak areas of 18 common peaks.
peaks in the ethyl acetate extract of 10 batches of *S. spatulifolius*, and the eigenvalues, variance contribution rate, and cumulative contribution rate of each component were obtained.

IBM SPSS Statistics 22.0 software was used for statistical analysis of animal experimental data. One-way analysis of variance was used for analysis, and *t* test was used for comparison between groups. The degree of swelling was expressed as *x ± s*. The difference between the model group (normal saline group) and each administration group was compared. *P* < 0.05 indicated that the difference was statistically significant, and *P* > 0.05 indicated that the difference was not statistically significant.

The inhibition rate of dimethylbenzene-induced ear swelling in mice by extracts of ethyl acetate extract of *S. spatulifolius* from different origins was used as the parent sequence, and the peak areas of 18 common peaks in the HPLC fingerprint were used as the subsequence. For calculation methods and steps, refer to [34, 35] for GRA. The peak area of the common peaks in the HPLC fingerprint was compared.

### Table 2: Experimental results of ear swelling induced by dimethylbenzene in mice (*n* = 8).

<table>
<thead>
<tr>
<th>Groups</th>
<th>Dosage of administration (g/kg)</th>
<th>Swelling degree (mg) (‾x ± s)</th>
<th>Inhibition ratio (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model group</td>
<td>—</td>
<td>4.9 ± 1.0</td>
<td>—</td>
</tr>
<tr>
<td>Positive group</td>
<td>0.003</td>
<td>1.6 ± 0.7**</td>
<td>67.35</td>
</tr>
<tr>
<td>Wuming, Nanning</td>
<td>3.4</td>
<td>2.4 ± 1.0**</td>
<td>51.02</td>
</tr>
<tr>
<td>Cen xi, wuzhou</td>
<td>3.4</td>
<td>2.7 ± 1.6**</td>
<td>44.90</td>
</tr>
<tr>
<td>Nanning qingxiu district</td>
<td>3.4</td>
<td>2.6 ± 1.0**</td>
<td>46.94</td>
</tr>
<tr>
<td>Zhaoping, Guangzhou</td>
<td>3.4</td>
<td>2.8 ± 1.0**</td>
<td>42.86</td>
</tr>
<tr>
<td>Nanning hengxian</td>
<td>3.4</td>
<td>2.5 ± 1.1**</td>
<td>48.95</td>
</tr>
</tbody>
</table>

Note. Compared with model group *P* < 0.05; **P** < 0.01.

### Table 3: Experimental results of ear swelling induced by dimethylbenzene in mice (*n* = 8).

<table>
<thead>
<tr>
<th>Groups</th>
<th>Dosage of administration (g/kg)</th>
<th>Swelling degree (mg) (‾x ± s)</th>
<th>Inhibition ratio (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model group</td>
<td>—</td>
<td>8.7 ± 1.7</td>
<td>—</td>
</tr>
<tr>
<td>Positive group</td>
<td>0.003</td>
<td>1.6 ± 0.6**</td>
<td>81.61%</td>
</tr>
<tr>
<td>Hepu, Beihai</td>
<td>3.4</td>
<td>4.5 ± 1.7**</td>
<td>48.28%</td>
</tr>
<tr>
<td>Fucheng, Beihai</td>
<td>3.4</td>
<td>4.3 ± 1.7**</td>
<td>50.57%</td>
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<tr>
<td>Shatian, Beihai</td>
<td>3.4</td>
<td>4.7 ± 1.6**</td>
<td>45.98%</td>
</tr>
<tr>
<td>Jinghai, Beihai</td>
<td>3.4</td>
<td>4.5 ± 1.7**</td>
<td>48.28%</td>
</tr>
<tr>
<td>Xinggang, Beihai</td>
<td>3.4</td>
<td>4.6 ± 1.4**</td>
<td>47.13%</td>
</tr>
</tbody>
</table>

Note. Compared with model group *P* < 0.05; **P** < 0.01.

![Figure 1: The overlay chromatograms of 10 batches of ethyl acetate extracts of *S. spatulifolius*.](image-url)
taken as the independent variable (X), the peak areas of the No. 1 peak to the No. 18 peak were recorded as x1–x18 in turn, and the inhibition rate of the mice ear swelling caused by dimethylbenzene was the dependent variable (Y). The data were imported into SIMCA-P software for PLSR.

3. Results

3.1. Anti-Inflammatory Effect. In this study, dimethylbenzene induced ear swelling in mice was used to evaluate extracts of ethyl acetate extract of *S. spatulifolius* anti-inflammatory activity. Due to the large number of mice, 10 batches of ethyl acetate extracts from *S. spatulifolius* were divided into two experiments. The previous study by our research group found that the high-dose ethyl acetate extract group (3.40 g/kg) had the best inhibitory effect on mice ear swelling. Therefore, the most effective dose of 3.40 g/kg was selected as the administration dose for mice. The results showed (Table 2 and Table 3) that compared with the model group, the ethyl acetate extract group of the S1–S10 samples significantly reduced the degree of mice ear swelling (*P < 0.05; **P < 0.01), and the ethyl acetate extracts of *S. spatulifolius* from different origins could inhibit the ear swelling of mice.

3.2. Fingerprint Analysis. In the process of establishing the fingerprint in this experiment, the chromatographic conditions were investigated, and it was found that when methanol–0.1% glacial acetic acid solution was used as the mobile phase, the detection wavelength was 260 nm, the column temperature was 25° C, and the flow rate was 0.8 mL/min, the separation effect and peak shape of each chromatographic peak were good, the baseline was stable, and the number of peaks was large. The method validation results of fingerprint established showed that the relative standard deviations of relative retention times for each chromatographic peak were good, the baseline was stable, and the sample was stable in 24 h.

Comparing the chromatograms of 10 batches of ethyl acetate extracts from *S. spatulifolius*, 18 common peaks were
identified, of which the 5th peak was protocatechuic acid and the 11th peak was caffeic acid. Protocatechuic acid has good resolution and peak shape and moderate retention time, so it was used as the reference peak. The overlay chromatograms of 10 batches of ethyl acetate extracts of *S. spatulifolius* is shown in Figure 1, the chromatogram of the mixed reference substance is shown in Figure 2, and the chromatogram of the ethyl acetate extract is shown in Figure 3. A total of 10 batches of ethyl acetate extracts from *S. spatulifolius* fingerprint similarity evaluation results are shown in Table 4. The results showed that, except for S6, which had a similarity of 0.879, the similarity of the other batches of samples were all greater than 0.9, indicating that the quality

<table>
<thead>
<tr>
<th>Peaks</th>
<th>Principal components 1</th>
<th>Principal components 2</th>
<th>Principal components 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.918</td>
<td>-0.206</td>
<td>0.101</td>
</tr>
<tr>
<td>2</td>
<td>0.624</td>
<td>-0.167</td>
<td>0.736</td>
</tr>
<tr>
<td>3</td>
<td>-0.180</td>
<td>0.720</td>
<td>-0.406</td>
</tr>
<tr>
<td>4</td>
<td>0.980</td>
<td>0.134</td>
<td>-0.024</td>
</tr>
<tr>
<td>5</td>
<td>0.230</td>
<td>0.166</td>
<td>0.940</td>
</tr>
<tr>
<td>6</td>
<td>0.940</td>
<td>-0.187</td>
<td>0.239</td>
</tr>
<tr>
<td>7</td>
<td>0.611</td>
<td>0.527</td>
<td>0.575</td>
</tr>
<tr>
<td>8</td>
<td>0.694</td>
<td>0.489</td>
<td>0.527</td>
</tr>
<tr>
<td>9</td>
<td>0.818</td>
<td>0.318</td>
<td>0.444</td>
</tr>
</tbody>
</table>

Table 6: Rotation component matrix.
of *S. spatulifolius* in each production area was basically stable, and the similarity of individual production areas was low, which may be related to the growth environment, harvesting season, cultivation method, processing method, and other factors.

The cluster analysis results are shown in Figure 4. The results showed that 10 batches of *S. spatulifolius* were grouped into four categories; the first category included S2, S7, S8, and S9; the second category included S1, S3, S5, and S6; the third category included S4; the fourth category included S10. The results showed that the sample classification was not affected by the origin, but may be related to the content of some chemical components.

PCA was a widely used statistical method, which was used to simplify data and quickly realize the visual pattern recognition of pattern or relationships. The principal component analysis results of 10 batches of ethyl acetate extracts from *S. spatulifolius* showed that with the characteristic value was greater than 1 as the principle of principal component extraction stated, three principal components could be extracted, and the 3D diagram of the principal components was shown in Figure 5. As shown in Table 5, the cumulative variance contribution rate of the first three principal components was 94.776%, which could comprehensively reflect the basic characteristics of its chemical components. It indicated that these three principal components represented most of the information contained in the 18 common peaks of the ethyl acetate extract of *S. spatulifolius*. From the rotation component matrix (Table 6), it was speculated that the reason for the difference in the quality of various batches of *S. spatulifolius* was the result of the combined effect of various chemical components. The variance contribution rate of the first principal component was 60.814%, which was the most “informative” component, and its information mainly came from chromatographic peaks 1, 2, 4, 6–9, 11, 12, 17, and 18; the variance contribution rate of the second principal component was 20.249%, and the information mainly came from chromatographic peaks 3, 13–15; the variance contribution rate of the third principal component was 13.713%, and the information mainly came from the 2, 5, 10 and 16 chromatographic peaks.

3.3. Spectrum-Effect Correlation Analysis. GRA is a quantitative analysis method in gray system theory, which is used to measure the gray relational degree between the target and the factors. The greater the gray correlation degree between
the two, the greater the correlation, and the smaller the gray correlation degree, the smaller the correlation [36]. In the establishment of the spectral-effect relationship, the GRA method mainly analyzes the correlation between the components and efficacy, but it cannot determine whether the characteristic peaks have a positive or negative effect on the efficacy. Compared with GRA, PLSR is a common technique and has been successfully applied to solve many practical problems, and this method is simple and accurate and can describe how each characteristic peaks contribute positively or negatively to the efficacy, and it gives the magnitude of the contribution, but the ability to analyze the correlation between composition and effect is poor. Therefore, the combined application of GRA and PLSR can draw on each other’s strengths to more accurately to analyze the spectral effect correlation of traditional Chinese medicine [7–12].

The GRA results were shown in Table 7. The correlation degree basically reached 0.8. Except for the No. 5 peak and No. 16 peak, the correlation degrees of the rest of the characteristic peaks were all greater than 0.8. It showed that the anti-inflammatory effect of ethyl acetate extract of S. spatulifolius was the result of the synergistic effect of multiple components. According to the degree of correlation, the order of contribution of each component to anti-inflammatory effect was determined as 8 > 13 > 14 > 7 > 2 > 15 > 18 > 9 > 10 > 1 > 6 > 12 > 4 > 17 = 11 > 3 > 5 > 16. PLSR obtained the partial regression coefficients (Figure 6) and VIP values (Figure 7) of 18 characteristic peaks (X) and inhibition rates (Y). Among the characteristic peaks of ethyl acetate extract of S. spatulifolius, nine peaks were positively correlated with the inflammation inhibition rate, and the remaining nine peaks were negatively correlated with the inflammation inhibition rate. The contribution of X to Y was determined according to the size of the VIP information. Generally speaking, the greater the VIP value, the greater the correlation between the variable and the drug efficacy. When VIP>1, this characteristic peak has a significant contribution to the inflammation inhibition rate. The results showed that the VIP values of peaks 5, 16, 4, 3, 2, and 12 were greater than 1. It showed that in the HPLC fingerprint of ethyl acetate extract of S. spatulifolius, its corresponding components played an important role in the inhibition rate of inflammation. The GRA and PLSR were combined to analyze, and the gray correlation degree>0.8, the partial least squares regression coefficient was positive, and the VIP value was greater than 1 for the chromatographic peak screening conditions. Finally, it was determined that the corresponding components of absorption peaks 3, 4, and 12 were the components in the ethyl acetate extract of S. spatulifolius that played an important role in the anti-inflammatory effect of mice.

4. Conclusions

In this study, the fingerprints of 10 batches of ethyl acetate extracts from S. spatulifolius were established, and 18 common peaks were identified. The evaluation results of the similarity of fingerprints showed that as excepted for S6, the other similarities were all above 0.915. Two peaks were identified by the reference method: protocatechuic acid absorption peak and caffeic acid absorption peak. The results of cluster analysis showed that samples from 10 different origins were clustered into four categories, which were consistent with the results of PCA. It showed that this method was feasible for analysis. In this study, GRA and PLSR were used to study the relationship between the common peaks of HPLC fingerprints of ethyl acetate extract of S. spatulifolius and the inhibition rate of inflammation. It was indicated that the corresponding components of No. 3, 4, and 12 absorption peaks in the ethyl acetate extract of S. spatulifolius were closely related components of the anti-inflammatory function of S. spatulifolius, and the spectrum-effect correlation between the ethyl acetate extract of S. spatulifolius and the anti-inflammatory activity was preliminarily expounded, which laid the foundation for the research on its pharmacodynamic substances. Therefore, chemical separation, LC-MS and other means could be used for further analysis to explore the substances represented by these three absorption peaks. The anti-inflammatory efficacy experiment was carried out on the three characteristic peak compounds separated and analyzed to further determine the anti-inflammatory efficacy of the compounds, which was of great significance for the determination of quality markers and quality control of S. spatulifolius medicinal materials.

Data Availability

The data used to support the findings of this study are included within the article.

Conflicts of Interest

The authors declare that there are no conflicts of interests.

Acknowledgments

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