Hindawi Evidence-Based Complementary and Alternative Medicine Volume 2018, Article ID 4689767, 9 pages https://doi.org/10.1155/2018/4689767



## Research Article

# Organoleptic Evaluation of Amomi Fructus and Its Further Background Verified via Morphological Measurement and GC Coupled with E-Nose

Dong Xu,¹ Yan Lin,¹ Rudolf Bauer,² Hui-Rong Chen,¹ Rui-Qi Yang,¹ Hui-Qin Zou,¹,¹ and Yong-Hong Yan,¹

<sup>1</sup>School of Chinese Materia Medica, Beijing University of Chinese Medicine, Beijing 100029, China <sup>2</sup>Institute of Pharmaceutical Science, University of Graz, 8010 Graz, Austria

Correspondence should be addressed to Hui-Qin Zou; zouhuiqin\_bucm@sina.cn and Yong-Hong Yan; lxdyyh@yeah.net

Received 21 November 2017; Revised 22 January 2018; Accepted 1 February 2018; Published 5 March 2018

Academic Editor: Wen-yi Kang

Copyright © 2018 Dong Xu et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The present study investigated the maneuverability and reasonability of sensory analysis, which has been applied in TCM identification for a long time. Ten assessors were trained and generated the human panel to carry out the organoleptic evaluation of twenty-five batches of Sha-Ren samples. Accordingly, samples were scored from 0 (lowest) to 10 (highest) for sensory attributes. Based on this, samples were divided into three classes: high class (Yang-Chun-Sha from Guang-Dong), moderate class (Yang-Chun-Sha samples from Yun-Nan and Guang-Xi), and low class (Lv-Qiao-Sha from marketplaces). For further background, three instrumental approaches were employed: morphological measurement with three indices (longitudinal diameter, transverse diameter, and 100-fruit weight), GC for determination of bornyl acetate contents, and E-nose for aromatic fingerprint. It is demonstrated in the results that GC and E-nose analyses were in great agreement with organoleptic evaluation. It gives insights into further studies on searching better morphological indicators and improving discriminant model of E-nose.

#### 1. Introduction

Amomi Fructus, called "Sha-Ren" in Chinese, has been one of the commonly used herbs in Traditional Chinese Medicine (TCM) for more than 1,300 years, mainly for the treatment of gastrointestinal diseases. Besides, its widespread application in food industry also gains much attentions, not only in China, but in other Southeast Asian countries, including Thailand, Vietnam, Burma, Indonesia, and so forth. Its yearly consumption reaches more than 3.1 million kg and maintains a constant increasing trend, which reflects strong market demands and potential commercial opportunities. However, due to confusable species of Sha-Ren from different producing areas, its price varies remarkably more than a hundredfold. Therefore, it is urgent and necessary to develop a rapid and reliable approach to distinguish Sha-Ren from different classes. On the other hand, the adulteration also needs cautiousness, especially in clinical use, because

incorrect or fake herbal medicines could result in low clinic effect or even poisoning. Many qualitative and quantitative methods have been employed in quality assessment of Sha-Ren from different species and habitats, for instance, GC-MS and IR analysis [1, 2]. However, those conventional physicochemical assays are mostly time-consuming, expensive, of high technical skills, and not so practical.

Macroscopic identification, as one of the five typical methods for TCM authentication [3–5], has been acknowledged to be a simple, convenient, and efficacious approach. During its procedure, morphological characteristics (shape, weight, length, etc.) and sensory properties (odor, taste, etc.) are carefully selected, observed, recorded, and analyzed, which help well-trained experts to do a quick and good job in marketing test of TCM [6–8]. Nevertheless it cannot not be neglected that the obvious drawbacks of this method are low repeatability and imprecision and depending on the experience of the estimators to a great extent. Therefore, this

leaves us with a serious of problems: Which morphological characteristics should be chosen to be valuable indices? What is the slight discrepancy among samples applying smelling, tasting, and touching senses? How to define the comparability of results from different experts, whom to trust, and whom not to? All in all, though this macroscopic identification seems to be a promising method for TCM quality evaluation, we firstly have to seek out the solutions to all the problems mentioned above.

The unique advantage of organoleptic analysis benefits from its capability of offering information perceived via human senses stimulating by a complex set of chemical compounds. For instance, odor is mainly perceived by the interactions of the volatiles with the olfactory epithelium in our nasal cavity [9]. Therefore, it could be a feasible way to introduce instrumental analysis as an additional supplementation, which helps to overcome the disadvantages of organoleptic analysis and also gives further background to demonstrate the material basis. Proper instrumental techniques permit detailed and objective information of human senses. Instead of vague odor from smelling, GC and electronic nose (E-nose) could provide a detailed and holistic odor profile, respectively. The former gives information concerning individual compounds and the latter concerning the origin, nondestructive, and holistic aromatic fingerprint. These two techniques have been employed in many aromatic researches and both got reliable results, separately or jointly [10-12]. However, the number of reports focused on combining these two instrumental techniques with morphological measurement and illustrating the basis of organoleptic evaluation from the material aspect is still limited, especially involving TCM.

In Pharmacopoeia of People's Republic of China [13], the dried fruits of three species from Zingiberaceae family are recorded as legal origins of Sha-Ren: (I) Amomum villosum Lour., (II) Amomum villosum Lour. var. xanthioides T. L. Wu et Senjen, and (III) Amomum longiligulare T. L. Wu. However, according to our previous investigations, the first (Yang-Chun-Sha in Chinese) and second (Lv-Qiao-Sha in Chinese) have been mainstream products whilst the third (Hai-Nan-Sha in Chinese) has not been easily pursued and collected in trading markets. Besides, back to the original habitat of the third, namely, Hai-Nan province, its plantation has been in a poor state for many years. On the contrary, the first one is well-known with its high quality and considered as genuine TCM from famous regions in Guang-Dong province. Even more noteworthy is that in 2005 Yang-Chun-Sha was nominated as one of 1992 China protected geographical indication products [14], among them 146 products locate themselves in Guang-Dong province, which means Yang-Chun-Sha representing high quality. As a result, Yang-Chun-Sha's price occupies the highest. Many researches have been carried on to find its unique indices and protect its correct usage [15, 16]. The contents of volatile constituents of Yang-Chun-Sha play a significant role in the authentication as well as its particular odor.

In the view of pros and cons of each technique, there are no universal and perfect methods for the simultaneous

analysis of every volatile compound and the particular odor, which is essential for the development of distinguishing Sha-Ren rapidly and reliably. To the best of our literature survey, no study on organoleptic analysis of Sha-Ren coupled with GC and E-nose has been reported to date. Therefore, this study herein aims (1) to establish a practical and precise organoleptic method for identification of Sha-Ren, (2) to replenish sensory odor with detailed information of volatile compounds and digitalized aromatic fingerprint from GC and E-nose, and (3) to offer high-priced Sha-Ren material background and shed light on its reasonability.

#### 2. Materials and Methods

2.1. Chemicals and Reagents. Bornyl acetate standard (batch number 110759-200604) was purchased from National Institutes for Food and Drug Control (Beijing, China). Absolute ethyl alcohol and other reagents used were all at chromatographic grade and from Dikma Technologies Co., Ltd. (Beijing, China).

2.2. Sha-Ren Samples. A total of 25 batches of Sha-Ren were collected: 19 batches of them were Yang-Chun-Sha from three habitats (Guang-Dong, Yun-Nan and Guang-Xi) and the rest 6 batches of them were Lv-Qiao-Sha from marketplaces. As mentioned above, the third species Hai-Nan-Sha were not obtained. All the samples have been authenticated via Professor Yong-Hong Yan from Department of Chinese Materia Medica of Beijing University of Chinese Medicine. They were packed in sealable plastic bags separately and stored at 4°C until analysis.

2.3. Organoleptic Evaluation. Based on observing, touching, and sniffing senses, organoleptic analysis was carried out by a human panel with ten assessors [17, 18], consisting of 10 trained postgraduate students of Beijing University of Chinese Medicine. The samples were evaluated by the panelists one by one, who were also placed in separate labs for unbiased analysis. Sha-Ren samples were taken out and presented at room temperature for at least two hours. They were scored from 0 (lowest) to 10 (highest) for sensory attributes of appearance, texture, and odor.

2.4. Morphological Measurement. Three objective indices were chosen to represent the morphological characteristics: longitudinal diameter (LD), transverse diameter (TD), and 100-fruit weight (100 FW). According to published reports and experienced experts, these three indices summarized the size of Sha-Ren and expressed the extent of oiliness, which are closely related to the appearance and texture of Sha-Ren. What is more, these three could be measured into figures, which helped us conduct further analysis.

Method of coning and quartering were employed to take 20 Sha-Ren fruits. Then LD and TD values were measured by vernier caliper (Measuring Instrument & Cutting Tool Factory, Beijing, China) and afterwards the mean was calculated.

Also using method of coning and quartering but to take 100 Sha-Ren fruits, then 100 FW value was measured by BS-124S electronic analytical balance (Sartorius, Germany) in triplicate and the means was calculated as well.

All values should not be utilized until RSD < 3%.

2.5. GC Analysis. According to previous reports, bornyl acetate is one of the main active volatile compounds [19, 20] in Sha-Ren and also one of the legal examination indicators in Chinese pharmacopoeia. Therefore, in this paper, bornyl acetate was chosen to be the index of detailed aromatic information and help supplement the results of sensory analysis.

Bornyl acetate standard was weighed precisely and dissolved in absolute ethyl alcohol to 11.90 mg/mL.

Around 1g of pulverized Sha-Ren samples was added to a conical flask with 25 mL absolute ethyl alcohol and extracted with ultrasonic at 40 kHz for 30 min. After cooling down to the room temperature and compensating weight, the processed samples were filtered and stored at 4°C until analysis.

The GC system included an Agilent 7890A instrument (California, America) with a flame ionization detector (FID): GC conditions: Agilent DB-1 column, (0.25  $\mu$ m, 0.25 mm × 30 m; Quartz Capillary); carrier gas: nitrogen; flow rate: 1.3 mL/min; the column oven temperature: 100°C;tThe injection temperature: 230°C; the FID temperature: 250°C; split ratio: 10:1; the injection volume: 1  $\mu$ L. The linearity of this method was assayed by analyzing standard solutions in the range of 0.1190–0.8333 mg/mL for bornyl acetate.

2.6. E-Nose Analysis.  $\alpha$ -Fox 3000 E-nose (Alpha MOS, Co., Ltd., France) with 12 metal-oxide gas (MOG) sensors was employed to obtain the digitalized aromatic fingerprint for the assessment of the odor strength. This E-nose instrument contains three main parts: an automatic sampling system, namely, HS-100 sampler, two chambers with six MOG sensors in each, and a data acquisition unit connected to a computer, which allows data storage and processing. The materials and properties of these 12 MOG sensors have been reported in our previous researches as well as the optimization method [21]. The whole 12 MOG sensors consist of (i) six LY-type sensors including LY2/LG, LY2/G, LY2/GH, LY2/AA, LY2/gCT, and LY2/gCTL (majorly sensitive to short chain volatile fatty acids and aldehydes); (ii) four P-type sensors including P10/1, P10/2, P40/1, and PA/2 (majorly sensitive to methane, propane, and other aliphatic nonpolar molecules), and (iii) two T-type sensors including T30/1 and T70/2 (majorly sensitive to polar alcoholic and chlorinated compounds), which allows E-nose perceived responses of most volatile compounds [22]. All 12 MOG sensors were named from S1 to S12.

Sha-Ren samples were firstly ground and sifted through a mesh of 850  $\mu$ m; then 0.4 g of powder was sealed into a glass and underwent predefined thermoincubation, which was vital to facilitate the generation and kinetic equilibrium of headspace volatile compounds inside the glass. Secondly the upper air containing particular volatile compounds was

injected into the  $\alpha$ -Fox 3000 E-nose system via the HS-100 autosampler applying pure air as the carrier gas at a flow rate of 150 mL/min. The incubation conditions were programmed at 45°C for 600 seconds with 250 r/s rotating speed. The temperature and volume of injection were set to 50°C and 500  $\mu$ L.

The volatile compounds in the upper air from the sealed glass interacted with the MOG sensors and resulted in changes of their resistance, positively or negatively, which were recorded in the computer. The relative change in their resistance ( $\Delta R/R_0$ ) value was monitored precisely per second for 200 s followed by the dormancy period of 600 s to ensure that the sensors must return to the base line and avoid their poisoning. The output  $\Delta R/R_0$  values were plotted against time. In that case, 12 MOG sensors lasting 200 seconds brought every sample a huge data array of 2400 values. Herein in our research, the absolute value of maximum sensor response was selected and analyzed to generate the holistic aromatic fingerprint of every sample. The test of each sample was repeated six times.

2.7. Statistical Analysis. Friedman test, one of nonparametric tests, was introduced to analyze the 25 related samples and compare the consistency of the organoleptic evaluation by 20 panelists. Then logistics regression analysis with ordinal variables was employed to find out the differences of sensory characteristics among the samples from different species and habitats.

For data processing of morphological measurement, hierarchical cluster analysis (HCA) and discriminant analysis (DA) were both utilized to see how many groups those all 25 batches samples were divided into and whether it was in the agreement with organoleptic evaluation.

In the past decades, E-nose has been used in many fields and well-known as a promising approach for fast and noninvasive detection [23, 24]. However, lack of unified and efficacy data analysis for its discriminative model has become the bottleneck to a great extent. Theoretically, it could be more than just one pattern recognition based on the rapid development of mathematical statistics. However, searching the optimum strategy becomes an inevitable problem. Our team figured out two solutions to improve the discriminative model and increase the distinguishing positive rate, concerning BestFirst + CfsSubsetEval (BC) feature extraction, and cascade classifier [21, 25, 26]. Therefore, as proven to be an efficient feature extraction method, BC was used to optimize the sensor with greater contributions and eliminate the tedious information of high-dimensional data from Enose. According to our previous achievements, three kinds of artificial neural networks (ANN), including Naïve-Bayes network (NBN), radial basis function network (RBF), and random forest (RF), were used for the discriminant models of E-nose original and optimized responses.

Simple correlation was applied to figure out the relationship among the experimental results of sensory and instrumental analyses, namely, organoleptic evaluation, GC, and E-nose. Afterwards, scatterplots and line charts were generated to see if there is any linear correlation. In the end,

Table 1: Descriptive statistics via	Friedman	test of 25	samples by 10
panelists.			

	N	Mean	Std. deviation	Minimum	Maximum
Sam. 1	10	9.50	.527	9	10
Sam. 2	10	8.90	.994	7	10
Sam. 3	10	9.40	.699	8	10
Sam. 4	10	9.50	.527	9	10
Sam. 5	10	8.10	.738	7	9
Sam. 6	10	9.00	.943	7	10
Sam. 7	10	8.80	.919	8	10
Sam. 8	10	9.10	.994	7	10
Sam. 9	10	5.40	.843	4	7
Sam. 10	10	5.10	.738	4	6
Sam. 11	10	4.30	.675	3	5
Sam. 12	10	5.10	.876	4	6
Sam. 13	10	4.50	.972	3	6
Sam. 14	10	5.00	1.054	3	6
Sam. 15	10	4.80	.919	3	6
Sam. 16	10	3.80	.919	2	5
Sam. 17	10	2.30	1.252	0	4
Sam. 18	10	2.50	.850	1	4
Sam. 19	10	2.80	1.135	0	4
Sam. 20	10	1.60	.843	0	3
Sam. 21	10	1.50	.707	1	3
Sam. 22	10	2.50	.707	1	3
Sam. 23	10	1.30	.949	0	3
Sam. 24	10	1.50	.972	0	3
Sam. 25	10	1.20	.919	0	3

principal components analysis (PCA) was used to classify the samples based on those three groups of data.

The statistical analyses were all performed applying SPSS version 22.0 (SPSS/IBM, Beijing, China; Licensing Certificate No. 20150408-LSBJ) and Weka software (free access at website: https://www.cs.waikato.ac.nz/ml/weka/).

#### 3. Results and Discussion

Odor is one of the key elements for identification of TCM. Since the pros and cons of sensory and instrumental approaches, organoleptic evaluation by 20 trained panelists was firstly carried out and then morphological measurement, GC, and E-nose were used to offer further background for further verification, which means digital and objective information of volatiles compounds and aromatic fingerprint.

3.1. Organoleptic Evaluation. The descriptive statistics including samples' number, mean value, and minimum/maximum were obtained via Friedman Test (Table 1). The test statistics were as follows: Chi-Square is 219.259, degree of freedom is 24, and Asymp. Sig. is 0.000, which is far smaller than 0.05. That reminds us that the organoleptic evaluation results of 10 panelists are different and can be quite useful for Sha-Ren's identification.

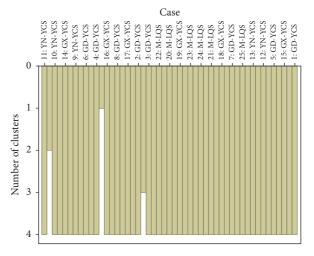


FIGURE 1: HCA vertical icicle diagram responses to samples based on morphological measurements with three indices (LD, TD, and 100 FW).

According to logistic regression analysis with ordinal variables, the probability of each kind of samples could be forecasted, which is how many scores they will earn through this human panel. For instance, the total number of Yang-Chun-Sha from Guang-Dong province is 80 (5 + 15 + 32 + 28) as shown in Table 2. Their predicted probabilities of being scored at 7, 8, 9, and 10 should be 0.06, 0.19, 0.40, and 0.35, which were calculated from 5/80, 15/80, 32/8, and 28/80, respectively. In other words, if samples scored at 8, 9, and 10 were logically assumed to have better quality, then more than 90% of Yang-Chun-Sha samples from Guang-Dong would be considered as superior products (0.19 + 0.40 + 0.35 = 0.94). In contrast, Yang-Chun-Sha samples from Yun-Nan and Guang-Xi should be of average quality, because their total ratios of scores 4, 5, and 6 are 0.84 and 0.5, respectively. And the samples of the lowest quality are Lv-Qiao-Sha collected from markets, whose probability of scores 0, 1, and 2 is 0.82.

To summarize the results of organoleptic evaluation by 10 panelists, all 25 batches of Sha-Ren samples were divided into three groups, namely, high class (Yang-Chun-Sha from Guang-Dong), moderate class (Yang-Chun-Sha samples from Yun-Nan and Guang-Xi), and low class (Lv-Qiao-Sha from marketplaces). It resembles the concept of TCM experts that Yang-Chun-Sha from Guang-Dong is the geoauthentic medicine of better quality and clinic effects.

3.2. Morphological Measurement. HCA applying furthest neighbor cluster method shows that all samples were classified into four groups (Figure 1). However, the results are confusable to a certain extent and not similar to organoleptic evaluation. It deserves to be noticed that all Lv-Qiao-Sha samples from markets were in the same group, though the classification of other samples was irregular.

To the next step, DA was used to find out whether these three morphological indices could help to get a satisfying identification of Sha-Ren samples. From the test of functions, the values of Wilks' Lambda are 0.295, 0.664, and 0.989,

TABLE 2: Cell information of logistic regression analysis with ordinal variables from 25 samples by 10 panelists.

Habitats         Species         0         1         2         3           GuangDong         Expected         0         2         2         2         2         1.139         4.479			Score					
YangChunSha         0         0         0           Observed         .000         .000         .000           Expected         .000         .000         .000           YangChunSha         0         0         0           Observed         .055         .247         1.149           Pearson Residual        235        499         -1.085           YangChunSha         2         2         10           Expected         .471         2.036         8.077           Expected         .727         .727           LvQiaoSha         7         21         21           Expected         7         21         21           Expected         8.280         19.830         20.631		3 4	5	9	7	∞	6	10
Observed         0         0         0           Expected         .000         .000         .000           Pearson Residual         .000         .000         .000           YangChunSha         0         0         0           Choserved         .055         .247         1.149           Pearson Residual        235        499         -1.085           YangChunSha         2         2         10           Expected         .471         2.036         8.077           Pearson Residual         2.235        025         .727           LvQiaoSha         7         21         21           Expected         7         21         21           Expected         8.280         19.830         20.631								
Expected       .000       .000       .000         Pearson Residual       .000       .000       .000         YangChunSha       0       0       0         Deserved       .055       .247       1.149         Pearson Residual      235      499       -1.085         YangChunSha       2       2       10         Doserved       .471       2.036       8.077         Pearson Residual       2.235      025       .727         LvQiaoSha       7       21       21         Expected       8.280       19.830       20.631			0	0	5	15	32	28
YangChunSha   000   000   000   000     YangChunSha   0   0   0   0     Expected   0.055   .247   1.149     Pearson Residual  235  499   -1.085     AngChunSha   2   2   10     Expected   .471   2.036   8.077     Expected   .225  025   .727     LvQiaoSha   7   21   21     Expected   8.280   19.830   20.631	000. 000.	000. 000	000.	000.	5.000	15.000	32.000	28.000
YangChunSha         0         0         0           Dbserved         0.055         .247         1.149           Pearson Residual        235        499         -1.085           YangChunSha         2         2         10           Expected         .471         2.036         8.077           Pearson Residual         2.235        025         .727           LvQiaoSha         7         21         21           Expected         8.280         19.830         20.631	000. 000.		000.	000.	000.	000.	000.	000
A colorerved 0 0 0 0 0 0 0 0 0 0 0 0 0 0 Expected 0.055 0.247 1.149  Pearson Residual -0.235 0.499 0.1.085  YangChunSha 2 2 10 0  Expected 4.71 2.036 8.077  Pearson Residual 2.235 0.025 7.27  LvQiaoSha 7 21 21  Expected 8.280 19.830 20.631								
Expected .055 .247 1.149 Pearson Residual .2354991.085  YangChunSha .2 .2 10 Expected .471 2.036 8.077 Pearson Residual 2.235025 .727  LvQiaoSha .7 .21 .21 Expected 8.280 19.830 20.631			19	12	1	0	0	0
Pearson Residual        235        499         -1.085           YangChunSha         2         2         10           Observed         .471         2.036         8.077           Pearson Residual         2.235        025         .727           LvQiaoSha         7         21         21           Expected         8.280         19.830         20.631	.247 1.149	479 11.597	17.566	14.056	.850	000	000.	000
YangChunSha       2       2       10         Observed       2       2       10         Expected       .471       2.036       8.077         Pearson Residual       2.235      025       .727         LvQiaoSha       7       21       21         Observed       7       21       21         Expected       8.280       19.830       20.631	499 -1.085		.425	647	.164	000.	000.	000
Ki         Observed Expected         2         2         10           Expected Expected Search         .471         2.036         8.077           Pearson Residual LyQiaoSha         7.235        025         .727           Observed Observed Search         7         21         21           Expected Search         8.280         19.830         20.631								
Expected .471 2.036 8.077 Pearson Residual 2.235025 .727  LvQiaoSha 7 21 21  Expected 8.280 19.830 20.631			10	9	0	0	0	0
Pearson Residual         2.235        025         .727           LvQiaoSha         7         21         21           Expected         8.280         19.830         20.631	2.036 8.077	.863 18.222	8.972	3.215	.145	000.	000.	000.
LvQiaoSha         7         21         21           Observed         7         21         21           Expected         8.280         19.830         20.631	025 .727		.372	1.597	381	000.	000.	000
Observed 7 21 21 Expected 8.280 19.830 20.631								
Expected 8.280 19.830 20.631	21		0	0	0	0	0	0
	19.830 20.631	.329 2.171	.583	.168	.007	000.	000.	000.
.100	.321 .100		767	411	085	000.	000.	000

respectively. Sig. values of them are all higher than 0.01. Therefore, this discriminant analysis could not be able to separate the samples efficaciously, which stays on the same side of HCA results.

3.3. Determination of Bornyl Acetate via GC. Bornyl acetate in extracted samples was identified by comparison of their retention time with that obtained from the chromatograms of standard substance. The content of bornyl acetate in each sample was calculated using standard curve of bornyl acetate and illustrated in Table 3. According to the level of bornyl acetate contents, all samples could clearly be divided into three groups. The first one contains eight batches of numbers 1, 2, 3, 5, 7, 12, 13, and 15, whose values are all exceeding 3. The second one would be composed of batch numbers 4, 6, 8, 10, and 14, whose values are between 1 and 2. And the third one consists of 12 batches, namely, numbers 9, 11, and 16-25. From these results we could tell that most of Yang-Chun-Sha samples are of higher amount of bornyl acetate and all Lv-Qiao-Sha samples gathered themselves together, with lower bornyl acetate, which agrees with the results of morphological measurement. Hence herein it is proven that Lv-Qiao-Sha samples are inferior-quality products, which means their price would be too high.

3.4. BC Feature Extraction and Discriminant Models of Aromatic Fingerprint from E-Nose. In the estimation of discriminant models using three different classifiers (NBN, RBF, and RF), two validation methods were applied and distinguishing positive rate was calculated, which were tenfold cross-validation in the training set and external test set validation in the test set.

Through BA feature extraction method performed in Weka software, six MOG sensors, namely, S2, S3, S6, S8, S10, and S12, were selected to generate the optimum data set. However, based on this discriminant model, only Yang-Chun-Sha from Guang-Dong could be distinguished from the other samples successfully, which leads to two categories based on E-nose responses. It is in similar situation of organoleptic evaluation results, where it sheds light on the concept again that Yang-Chun-Sha is the geoauthentic medicine.

As shown in Table 4, distinguishing positive rates of three classifiers were all increased after BC feature extraction. In a word, this optimum data set of six sensors are containing enough effective information extracted from aromatic fingerprint from E-nose, which enables it to afford the identification task by eliminating the redundant data.

3.5. Correlation among Sensory and Instrumental Analysis of Sha-Ren's Odor. For the simple correlation analysis, the organoleptic analysis showed positive correlation with GC and E-nose, with correlation index of 0.650 and 0.807, respectively. However, according to the scatterplots and line graphs, no obvious linear regression was found. But it is easy to see that the curves of organoleptic analysis and GC are in the same trend.

As for PCA, samples were placed in three zones (Figure 2). These results are in great consistency with the

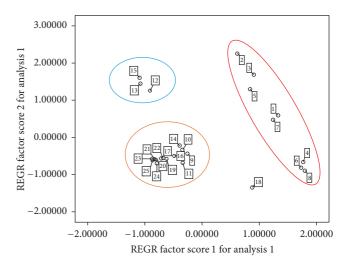


FIGURE 2: PCA score plot responses to Sha-Ren samples with PC1 and PC2.

results with organoleptic evaluation, though with subtle discrepancy. Furthermore, this model obviously provides much more details than E-nose applied alone, which could only differentiate the geoauthentic medicine from the others. That comes into a hypothesis: even E-nose is much more sensitive, but still there is some deeper information extracted by human panel, which have been searched out.

#### 4. Conclusions

Twenty-five batches of Sha-Ren samples were collected from Guang-Dong, Guang-Xi, and Yun-Nan and marketplaces, which are Yang-Chun-Sha (Amomum villosum Lour.) and Lv-Qiao-Sha (Amomum villosum Lour. var. xanthioides T. L. Wu et Senjen) from Zingiberaceae family. For rapid and reliable authentication of confusable and of great price variance samples, sensory analysis was introduced to development a practical method. Based on a human panel with 10 trained assessors, organoleptic evaluation divided experimental samples into three classes: high class (Yang-Chun-Sha from Guang-Dong), moderate class (Yang-Chun-Sha samples from Yun-Nan and Guang-Xi), and low class (Lv-Qiao-Sha from marketplaces). For providing further material background and objective verification, three instrumental approaches were employed: morphological measurement with three indices (longitudinal diameter, transverse diameter, and 100-fruit weight), GC for determination of bornyl acetate contents, and E-nose for aromatic fingerprint. It is demonstrated in the results that GC and E-nose analyses were in great agreement with organoleptic evaluation, whilst morphological measurement nevertheless showed irregular classification. It could be explained that those three indices were not perfect and at least they did not represent the whole effective information. Further researches are in progress in our lab to look for better indicators and also improve the distinguishing positive rate of E-nose discriminant model.

TABLE 3: Experimental results of Sha-Ren samples.

Batch number	Sample name	Species	Habitats	Score of human panel	Bornyl acetate content	S6-sensor response
(1)	GD-YCS	Yang-Chun-Sha	Guang-Dong	9.5	3.82	0.04
(2)	GD-YCS	Yang-Chun-Sha	Guang-Dong	8.9	6.64	0.04
(3)	GD-YCS	Yang-Chun-Sha	Guang-Dong	9.4	5.68	0.04
(4)	GD-YCS	Yang-Chun-Sha	Guang-Dong	9.5	1.74	0.04
(5)	GD-YCS	Yang-Chun-Sha	Guang-Dong	8.1	5.03	0.04
(9)	GD-YCS	Yang-Chun-Sha	Guang-Dong	9.0	1.52	0.04
(7)	GD-YCS	Yang-Chun-Sha	Guang-Dong	8.8	3.66	0.04
(8)	GD-YCS	Yang-Chun-Sha	Guang-Dong	9.1	1.35	0.04
(6)	YN-YCS	Yang-Chun-Sha	Yun-Nan	5.4	0.88	0.03
(10)	YN-YCS	Yang-Chun-Sha	Yun-Nan	5.1	1.06	0.03
(11)	YN-YCS	Yang-Chun-Sha	Yun-Nan	4.3	0.48	0.03
(12)	YN-YCS	Yang-Chun-Sha	Yun-Nan	5.1	3.71	0.03
(13)	YN-YCS	Yang-Chun-Sha	Yun-Nan	4.5	4.04	0.03
(14)	GX-YCS	Yang-Chun-Sha	Guang-Xi	5.0	1.24	0.03
(15)	GX-YCS	Yang-Chun-Sha	Guang-Xi	4.8	4.31	0.03
(16)	GX-YCS	Yang-Chun-Sha	Guang-Xi	3.8	0.80	0.03
(17)	GX-YCS	Yang-Chun-Sha	Guang-Xi	2.3	0.73	0.03
(18)	GX-YCS	Yang-Chun-Sha	Guang-Xi	2.5	0.71	0.04
(19)	GX-YCS	Yang-Chun-Sha	Guang-Xi	2.8	0.69	0.03
(20)	M-LQS	Lv-Qiao-Sha	Marketplaces	1.6	0.68	0.03
(21)	M-LQS	Lv-Qiao-Sha	Marketplaces	1.5	0.70	0.03
(22)	M-LQS	Lv-Qiao-Sha	Marketplaces	2.5	0.74	0.03
(23)	M-LQS	Lv-Qiao-Sha	Marketplaces	1.3	0.72	0.03
(24)	M-LQS	Lv-Qiao-Sha	Marketplaces	1.5	0.52	0.03
(25)	M-LQS	Lv-Qiao-Sha	Marketplaces	1.2	0.64	0.03

Cliassifer	Origin	al data set	Optimum data set	
Chassilei	Tenfold cross-validation	External test set validation	Tenfold cross validation	External test set validation
NBN	54	56	78	84
RBF	66	72	78	84
RF	64	76	78	84

TABLE 4: Distinguishing positive rates of three classifiers (NBN, RBF, and RF) of original and optimum data set.

#### **Conflicts of Interest**

The authors report no conflicts of interest related to this manuscript.

#### **Authors' Contributions**

Dong Xu and Yan Lin contributed equally to this project.

### Acknowledgments

This project was financially supported by the National Natural Science Foundation of China (Grants nos. 81403054, 81573542, and 81202682).

#### References

- [1] W. Kang, F. Zhang, Y. Su, and Y. Guo, "Application of gas chromatography-quadrupole-time-of-flight-mass spectrometry for post-target analysis of volatile compounds in Fructus Amomi," *European Journal of Mass Spectrometry*, vol. 19, no. 2, pp. 103–110, 2013.
- [2] C.-G. Cheng, Y.-M. Ruan, and B.-L. Li, "Studies on the identification of fructus amomi from its confusable varieties by Fourier transform infrared spectroscopy," *Guang Pu Xue Yu Guang Pu Fen Xi/Spectroscopy and Spectral Analysis*, vol. 24, no. 11, p. 1355, 2004.
- [3] Z. Zhao, Y. Hu, Z. Liang, J. P.-S. Yuen, Z. Jiang, and K. S.-Y. Leung, "Authentication is fundamental for standardization of Chinese medicines," *Planta Medica*, vol. 72, no. 10, pp. 865–874, 2006
- [4] Y. B. Zhang, P. C. Shaw, C. W. Sze, Z. T. Wang, and Y. Tong, "Molecular authentication of Chinese herbal materials," *Journal of Food and Drug Analysis*, vol. 15, no. 1, pp. 1–9, 2007.
- [5] Z. Zhao, P. Xiao, Y. Xiao, and J. P. S. Yuen, "Quality assurance of Chinese Herbal Medicines (CHMs)," *Journal of Food and Drug Analysis*, vol. 15, no. 4, pp. 337–346, 2007.
- [6] B. Plutowska and W. Wardencki, "Aromagrams Aromatic profiles in the appreciation of food quality," *Food Chemistry*, vol. 101, no. 2, pp. 845–872, 2007.
- [7] Y. Xiong, X. Xiao, X. Yang et al., "Quality control of Lonicera japonica stored for different months by electronic nose," *Journal* of *Pharmaceutical and Biomedical Analysis*, vol. 91, pp. 68–72, 2014
- [8] H.-Q. Zou, G. Lu, Y. Liu et al., "Is it possible to rapidly and noninvasively identify different plants from Asteraceae using electronic nose with multiple mathematical algorithms?" *Jour*nal of Food and Drug Analysis, vol. 23, no. 4, pp. 788–794, 2015.
- [9] A. Biolatto, G. Grigioni, M. Irurueta, A. M. Sancho, M. Taverna, and N. Pensel, "Seasonal variation in the odour characteristics of whole milk powder," *Food Chemistry*, vol. 103, no. 3, pp. 960– 967, 2007.

- [10] R. Baranauskiene, P. R. Venskutonis, A. Galdikas, D. Senuliene, and A. Šetkus, "Testing of microencapsulated flavours by electronic nose and SPME-GC," *Food Chemistry*, vol. 92, no. 1, pp. 45–54, 2005.
- [11] S. Cui, J. Wu, J. Wang, and X. Wang, "Discrimination of American ginseng and Asian ginseng using electronic nose and gas chromatography-mass spectrometry coupled with chemometrics," *Journal of Ginseng Research*, vol. 41, no. 1, pp. 85– 95, 2017.
- [12] Q. Li, X. Yu, L. Xu, and J.-M. Gao, "Novel method for the producing area identification of Zhongning Goji berries by electronic nose," *Food Chemistry*, vol. 221, pp. 1113–1119, 2017.
- [13] National Pharmacopoeia Committee, *Pharmacopoeia of People's Republic of China*, Beijing: China Medical Science Press, Beijing, China, 2015.
- [14] General Administration of Quality Supervision, "Inspection and Quarantine of the People's Republic of China. China protected geographical indication products," 2015, http://www.cgi.gov.cn/Products/Detail/1936.
- [15] C. Deng, A. Wang, S. Shen, D. Fu, J. Chen, and X. Zhang, "Rapid analysis of essential oil from Fructus Amomi by pressurized hot water extraction followed by solid-phase microextraction and gas chromatography-mass spectrometry," *Journal of Phar*maceutical and Biomedical Analysis, vol. 38, no. 2, pp. 326–331, 2005.
- [16] S. Shen, Y. Sha, C. Deng, D. Fu, J. Chen, and X. Zhang, "Comparison of solid-phase microextraction, supercritical fluid extraction, steam distillation, and solvent extraction techniques for analysis of volatile consituents in Fructus Amomi," *Journal of AOAC International*, vol. 88, no. 2, pp. 418–423, 2005.
- [17] R. Jónsdóttir, G. Ólafsdóttir, E. Chanie, and J.-E. Haugen, "Volatile compounds suitable for rapid detection as quality indicators of cold smoked salmon (Salmo salar)," *Food Chemistry*, vol. 109, no. 1, pp. 184–195, 2008.
- [18] X. Zhou, Y. Chong, Y. Ding, S. Gu, and L. Liu, "Determination of the effects of different washing processes on aroma characteristics in silver carp mince by MMSE-GC-MS, e-nose and sensory evaluation," *Food Chemistry*, vol. 207, pp. 205–213, 2016.
- [19] H. Ying, J. Liu, and Q. Du, "Analysis and determination of oestrogen-active compounds in fructus amomi by the combination of high-speed counter-current chromatography and high performance liquid chromatography," *Journal of Chromatography B*, vol. 958, pp. 36–42, 2014.
- [20] H. G. Choi, I.-G. Je, G. J. Kim et al., "Anti-allergic inflammatory activities of compounds of amomi fructus," *Natural Product Communications (NPC)*, vol. 10, no. 4, pp. 631-632, 2015.
- [21] H.-Q. Zou, S. Li, Y.-H. Huang et al., "Rapid identification of Asteraceae plants with improved RBF-ANN classification models based on MOS sensor E-nose," Evidence-Based Complementary and Alternative Medicine, vol. 2014, Article ID 425341, 6 pages, 2014.
- [22] R. Upadhyay, S. Sehwag, and H. N. Mishra, "Electronic nose guided determination of frying disposal time of sunflower oil

- using fuzzy logic analysis," *Food Chemistry*, vol. 221, pp. 379–385, 2017.
- [23] F. Röck, N. Barsan, and U. Weimar, "Electronic nose: current status and future trends," *Chemical Reviews*, vol. 108, no. 2, pp. 705–725, 2008.
- [24] S. Cui, L. Yang, J. Wang, and X. Wang, "Fabrication of a sensitive gas sensor based on PPy/TiO2 nanocomposites films by layerby-layer self-assembly and its application in food storage," *Sensors and Actuators B: Chemical*, vol. 233, pp. 337–346, 2016.
- [25] H. Lin, Y. Yan, T. Zhao et al., "Rapid discrimination of Apiaceae plants by electronic nose coupled with multivariate statistical analyses," *Journal of Pharmaceutical and Biomedical Analysis*, vol. 84, pp. 1–4, 2013.
- [26] L. Peng, H.-Q. Zou, R. Bauer et al., "Identification of chinese herbal medicines from zingiberaceae family using feature extraction and cascade classifier based on response signals from E-Nose," Evidence-Based Complementary and Alternative Medicine, vol. 2014, Article ID 963035, 7 pages, 2014.

















Submit your manuscripts at www.hindawi.com























