

Original Research Article

Comparative studies on the toxicological, anti-inflammatory and analgesic properties of three sources of Xuedan in mice and their rapid identification by electronic tongue

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Abstract

Purpose: To compare the toxicological, anti-inflammatory and analgesic properties of three sources of Xuedan, viz, *Hemsleya omeiensis* (HO), *Hemsleya gigantea* (HG) and *Hemsleya dolichocarpa* (HD) in mice, and to study their rapid identification based on electronic tongue (E-tongue).

Methods: After 7 days of administration, the median lethal doses (LD_{50}) of the three xuedan decoctions in mice were determined. In addition, the anti-inflammatory and analgesic effects of the three xuedans were evaluated in mice using xylene-induced ear edema and acetic acid-induced pain. Furthermore, E-tongue technology was used to identify HO, HG and HD. Principal component analysis (PCA) and discriminant factor analysis (DFA) were used to analyze the data acquired by E-tongue.

Results: The median lethal dose (LD_{50}) values of *H. omeiensis*, *H. gigantea* and *H. dolichocarpa* were 32.3, 17.4 and 13.7g/kg, respectively. Compared with normal control group, the anti-inflammatory effects of Xuedan were obvious in xylene-induced ear edema ($p < 0.05$), and pain sensation was significantly inhibited in acetic acid-induced writhing test ($p < 0.05$). Furthermore, E-tongue technology effectively identified HO, HG and HD.

Conclusion: *H. omeiensis* exhibits the highest LD_{50} value and best analgesic effect among the three sources of xuedan. E-tongue technology is effective and rapid in identifying HO, HG and HD.

Keywords: Xuedan, *Hemsleya omeiensis*, *Hemsleya gigantea*, *Hemsleya dolichocarpa*, Anti-inflammation, Analgesia, Electronic tongue

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INTRODUCTION

Xuedan is derived from *Hemsleya* (*Cucurbitaceae*) which is distributed in subtropical and temperate regions of Asia,

especially in China [1]. In Chinese Traditional medicinal theory system, *xuedan* is used for clearing heat and removing toxins. *Xuedan* exerts antibacterial and anti-inflammatory effects. Consequently, *xuedan* is used for treating

various diseases such as hepatitis, enteritis, bronchitis and coronary heart disease [2]. In China, *xuedan* comes from three varieties of *Hemsleya* viz *Hemsleya omeiensis*, *Hemsleya gigantea* and *Hemsleya dolichocarpa*. Studies have revealed that there are wide differences in compositions and in bioactivities among the three sources of *xuedan*, especially with respect to their bitter taste. However, no studies have been carried out on effective and rapid strategy for discriminating between the three sources of *xuedan*. The present study was carried out to compare the toxicology and pharmacology of the three sources of *xuedan*, and to develop a rapid method based on electronic tongue for their identification.

EXPERIMENTAL

Plant materials

The three varieties of *xuedan* used in this study were provided by Sichuan Fuzheng Pharmaceutical Co., Ltd., and identified by Professor Xian-Ming Lu (College of Pharmacy, Chengdu University of Traditional Chinese Medicine, Chengdu, China).

Animal studies

A total of 200 mice (18 - 22 g) were obtained from the Chengdu *DaShuo* Experimental Animal Co., Ltd. (Chengdu, China). All animals received humane care according to the Declaration of Helsinki [3], and fed in accordance with standard laboratory protocols. All experimental protocols followed the principles and guidelines recommended by the Animal Care and Use Committee of Chengdu Municipal Hospital of Traditional Chinese Medicine, Chengdu, China [approval no. SCXK (chuan) 2015-3].

Acute toxicity studies

Acute toxicity studies were carried out in accordance with the procedure reported in previous studies, but with minor modifications [4-6]. First, the complete lethal dose (LD_{100}) and zero (0), i.e., lethal dose-0 (LD_0) were determined through oral administration of the test samples. The results showed that the LD_{100} and LD_0 values for *H. omeiensis* were 48 and 24 g/kg, respectively; and LD_{100} and LD_0 values for *H. gigantea* and *H. dolichocarpa* were 24 and 6 g/kg, respectively. Based on these results, mice were randomly divided into five groups ($n = 8$) for the testing of *Xuedan* samples.

The mice were fasted overnight but were permitted free access to water, and were

administered the designed dosages (Table 1) orally (*intragastrically*). The mice were observed for signs of poisoning or death for 14 days. Mortality was calculated from the number of mouse deaths in each group, and LD_{50} was calculated using the weighted regression probability unit method (Bliss method) with LD_{50} calculation version 2.0 software. The 95 % confidence interval was calculated using the improved method of Käber [6].

Anti-inflammatory studies

Mice were randomly divided into four groups ($n = 10$): control group, *H. omeiensis* group, *H. gigantea* group and *H. dolichocarpa* group. The test samples were administered orally at a dose of 0.6 g/kg (equivalent to daily human adult dose) for 7 continuous days. Then, 30 min after the last administration, 20 μ L of xylene was uniformly applied to both sides of the left auricle of each mouse, while the right ear was used as a control. After 30 min, the mice were euthanized. The ears were cut along the baseline of the auricle. The circular ear was punched in the same part of the left and right ears with an 8 mm diameter puncher [7]. The ears were weighed with an electronic balance, and the weight difference of the two ears ($W_l - W_r$) was used as swelling index (E_s), calculated as in Eq 1.

$$E_s = W_l - W_r \dots\dots\dots (1)$$

Inhibition of swelling (P_i) was calculated using Eq 2.

$$P_i (\%) = \{(E_{sc} - E_{st})/E_{sc}\}100 \dots\dots\dots (2)$$

where E_{sc} and E_{st} are the swelling index of control and test groups, respectively.

Analgesic studies

Mice were randomly divided into four groups ($n = 10$): control group, *H. omeiensis* group, *H. gigantea* group and *H. dolichocarpa* group. The mice in each group were orally administered corresponding test samples at a dose of 0.6 g/kg/day (equivalent to the daily adult human dose) continuously for 7 days. Then, one hour after the last administration, each mouse was treated with 0.6 % acetic acid (0.1 mL/kg) by intraperitoneal injection. The number of writhes (N_w) (abdominal contraction and extension of hind limbs) within 15 min after acetic acid injection in each group of mice was recorded, and the percentage inhibition (P_i) was calculated as in Eq 3 [8].

$$H (\%) = \{(N_{wc} - N_{wt})/N_{wc}\}100 \dots\dots\dots (3)$$

where Nwc and Nwt are the number of writhes of control and test groups, respectively.

Electronic tongue test

The α Astree E-tongue, used to investigate the taste of the three varieties of *Xuedan*, had six cross-sensitive potential sensors (ZZ, AB, GA, CA, DA and JE), Alphasoft data acquisition system, signal processing system and pattern recognition software (Alpha M.O.S, Version 2012.45). The method [8] measures E-tongue under conditions that ensure that the acquired data are reliable and stable. Each test sample powder (1.0 g) was extracted with 200 mL boiled water in a conical flask for 3 h. Thereafter, it was filtered through a 0.22 μ L microporous filter membrane, and 25 mL of the filtrate was placed in the auto-sampler tray of the E-tongue apparatus for analysis. Each sample was analyzed for 120 sec in triplicate. The data were recorded using Astree II software.

Statistical analysis

Data are expressed as mean \pm standard deviation (SD), and were analyzed using SPSS 21.0 software with single factor analysis of variance between groups. Values of $p < 0.05$

were considered statistically significant. All E-tongue data were analyzed using Alpha M.O.S. statistical software. Principal component analysis (PCA) and discriminant factor analysis (DFA) were used for discrimination analysis of the samples.

RESULTS

Acute toxicity

As can be seen from the Table 1, LD₅₀ values of the *H. omeiensis*, *H. gigantea*, *H. dolichocarpa* were 32.3, 17.4 and 13.7 g/kg, respectively. All samples showed moderate toxicity, but the *H. omeiensis* showed the lowest toxicity in mice, when compared to the other two sources of *xuedan* (32.3 g/kg vs. 17.4 and 13.7 g/kg).

Anti-inflammatory effects

The results of anti-inflammatory studies of the three *xuedan* sources are shown in Table 2. Compared with the normal group, the three aqueous extracts of samples significantly inhibited xylene-induced ear swelling in mice ($p < 0.01$). There was no significant difference in percent inhibition of swelling between the three samples.

Table 1: LD₅₀ values of the three sources of *xuedan* (n = 8)

Group	Dose (g/kg)	Mortality (%)	LD ₅₀ (g·(kg·day) ⁻¹)	95 % confidence limit (g·(kg·day) ⁻¹)
<i>H. omeiensis</i>	20.48	0	32.3	28.8-36.1
	25.6	0		
	32	25		
	40	87.5		
	50	100		
<i>H. gigantea</i>	10.24	0	17.4	15.6-19.4
	12.8	0		
	16	25		
	20	87.5		
	25	100		
<i>H. dolichocarpa</i>	10.24	0	13.7	11.8-16.0
	12.8	62.5		
	16	62.5		
	20	87.5		
	25	100		

Table 2: Effect of the three sources of *xuedan* on ear edema induced by xylene in mice

Group	Dosage (g/kg)	Swelling (%)	Inhibition (%)
Normal	--	76.43 \pm 6.05	--
<i>H. omeiensis</i>	0.6	54.85 \pm 7.17**	23.93
<i>H. gigantea</i>	0.6	62.88 \pm 6.08**	25.36
<i>H. dolichocarpa</i>	0.6	59.43 \pm 7.26**	27.58

Data are expressed as mean \pm SD (n=10); * $p < 0.05$, ** $p < 0.01$, compared with normal

Analgesic activity

Compared with the normal group, the three varieties of *xuedan* significantly reduced the number of acetic acid-induced writhes in mice ($p < 0.05$), and the effect of HO was the most significant ($p < 0.01$). There were significant differences in the percentage inhibition of writhes among HO, HG and HD ($p < 0.05$ (Table 3).

E-tongue results

Typical sensor responses recorded by E-tongue apparatus are shown in Figure 1. An average value was acquired based on the stable sensor responses of E-tongue analyses from 100 to 120 sec, and was used as the output data. The repeatability of the E-tongue results using this method of study is shown in Table 4.

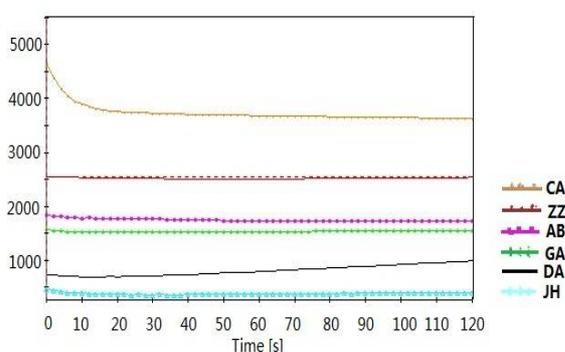


Figure 1: Typical sensor responses of E-tongue during measurement

In this study, 9 batches of samples were collected for each source of *xuedan*. Each sample was measured and analyzed three times. From the signals acquired through E-tongue measurements, 81 data points were used for further PCA and DFA. The raw data was projected onto the PC axis, and each sample will a set on the PCA three-dimensional map. The distance between sample sets is usually used to

indicate similarities in taste. The three-dimensional map of PCA is shown in Figure 2, and the three principal components were PC1 (82.83 %), PC2 (16.21%), and PC3 (0.66 %). The total degree of contribution of the three principal components was 99 %.

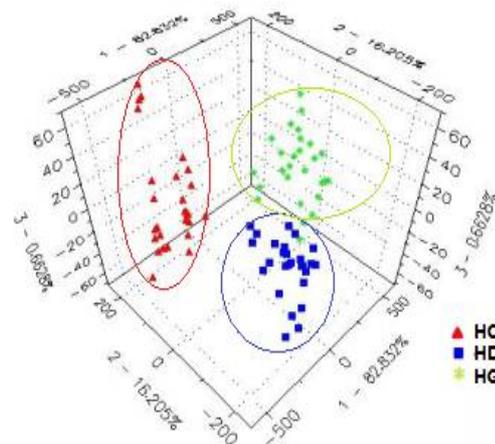


Figure 2: PCA score plots of E-tongue measurements of three varieties of *Hemsleya chinensis* (n = 9). Each sample was measured and analyzed three times. HO, HD and HG refer to *H. omeiensis*, *H. Gigantha*, *H. dolichocarpa*, respectively

Similar to PCA, a total 81 data points were projected onto the DFA score plots. As can be seen in Figure 3, the two principal components for E-tongue were DF1 = 92.36 % and DF2 = 7.64 %, and their total contribution was 100 %. The DFA model effectively classified the three sources *xuedan*. In addition, DFA is often used to establish a sample database for qualitative discrimination between unknown samples and determination of grouping information of unknown samples. This model can be used to identify samples from unknown base sources of *xuedan*.

Table 3: Effects of the three sources of *xuedan* on number of writhes in mice

Group	Dose (g/kg)	Number of writhes	Inhibition (%)
Normal	--	37.3 ± 6.86	--
<i>H. omeiensis</i>	0.6	19.2 ± 5.53**	48.53
<i>H. Gigantha</i>	0.6	28.2 ± 6.05 ⁺	26.66
<i>H. dolichocarpa</i>	0.6	30.6 ± 5.54 ⁺	17.96

Data are expressed as mean ± SD (n=10); * $p < 0.05$, ** $p < 0.01$, compared with normal

Table 4: Repeatability of E-tongue determination method (n = 6)

Sensor	CA	ZZ	AB	GA	DA	JE
RSD (%)	1.24	0.61	0.19	0.30	0.50	0.28

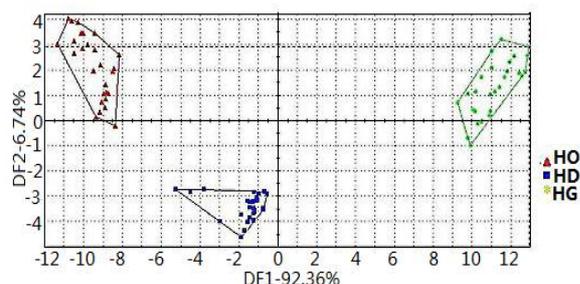


Figure 3: DFA score plots of E-tongue measurements of three varieties of *Hemsleya chinensis* ($n = 9$). Each sample was measured and analyzed three times. HO, HD and HG represent *H. omeiensis*, *H. Gigantha*, *H. dolichocarpa*, respectively

DISCUSSION

It is well known that traditional Chinese medicines (TCMs) have been applied in the treatment of various diseases for thousands years, and studies have demonstrated their pharmacological activities [10]. It has been shown that the main active phytochemical components of the genus *Hemsleya* are triterpenoid saponins which possess significant antitumor, antibacterial and anti-inflammatory properties [1,11]. Some scholars have suggested that hemslecin A is a major antibacterial agent in the genus *Hemsleya* genus [12]. It has been directly used as an effective agent for treating inflammation and bacterial infection-related disorders. Subsequent studies revealed that hemslecins A and B exhibit potent anti-HIV-1 activities [13-16]. Therefore, *Hemsleya* genus is a promising plant resource for discovery of drugs for treating infection and inflammation. However, some TCMs are not derived from one plant but lots of plants, and the use of original sources of the plants would be beneficial for guaranteeing the therapeutic effects of the TCMs.

Xuedan is widely distributed in southwest areas of China, in particular Sichuan. It is a popular herbal medicine in China, and it is derived from three varieties: *H. omeiensis*, *H. gigantha* and *H. dolichocarpa*. It has various pharmacological effects such as anti-inflammatory and analgesic properties, but there are some differences among the three sources of the drug. In the present study, *H. omeiensis* had the lowest toxicity, whereas its analgesic effect was significantly better those of the other two. This is an important experimental evidence for the clinical use of *xuedan*. *Xuedan* is characterized by light smell and an extremely bitter taste [2].

However, based on the feedback from the doctors in clinics and the users, *H. omeiensis* differs from the other two sources of *xuedan*

which have moderate bitter taste instead of extreme bitter taste. In addition, it produces better bioactive effects, relative to the other two sources. In view of these desirable qualities, a proposal to distinguish *H. omeiensis* from the other two sources of *xuedan* was designed in the present study. The bitter taste is the key point for quick and effective discrimination of the 3 sources of *xuedan*.

Electronic tongue technology is an objective quantitative evaluation method for sensory indicators. By collecting objective and accurate data, the results of the taste characteristics of samples are obtained, and the differences in the taste of different samples are scientifically reflected. It is increasingly used in the Chinese medicine industry. These results suggest that E-tongue combined with PCA can effectively discriminate between the three varieties of *Xuedan*, especially *H. omeiensis* which differs in taste from the other two sources of *Xuedan*.

DFA is a classification technique that optimizes differentiation by recombining sensor data and is another classical analysis model in addition to PCA. It can be used to establish a sample database for qualitative discrimination between samples from unknown base sources of *xuedan*. This study has demonstrated that electronic tongue technology, when combined with PCA and DFA can effectively distinguish among *Xuedans* from the three varieties of *Hemsleya*.

CONCLUSION

Hemsleya omeiensis possesses the highest LD_{50} value and best analgesic effect among the three varieties of *Hemsleya*. In addition, E-tongue, when combined with PCA and DFA, can effectively and rapidly discriminate between the three sources of *xuedan*.

DECLARATIONS

Acknowledgement

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Conflict of interest

No conflict of interest is associated with this work.

Contribution of authors

The authors declare that this work was done by the authors named in this article and all liabilities pertaining to claims relating to the content of this article will be borne by them. All authors read and approved the manuscript for publication.

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