Bioactivity guided isolation of apigenin from Stachys lavandulifolia Vahl. in mice with anxiolytic effects

Mohammad Rabbani¹, Seyed-Ebrahim Sajjadi², Masoumeh Karimi-Firouzjaei¹, and Mustafa Ghanadian¹*¹

¹Department of Pharmacology, Isfahan University of Medical Sciences, Isfahan, Iran
²Department of Pharmacognosy, School of Pharmacy and Pharmaceutical Sciences, Isfahan University of Medical Sciences, Isfahan, Iran
³Isfahan Pharmaceutical Sciences Research Centre, School of Pharmacy and Pharmaceutical Sciences, Isfahan University of Medical Sciences, Isfahan, Iran

*Corresponding author: Mustafa Ghanadian, Tel: +98-3117922553, Email: ghannadian@gmail.com

Article History:
Received: 7 October 2017
Accepted: 4 February 2018

Article Type: Original Article

ABSTRACT

Introduction: Stachys lavandulifolia is traditionally used for the treatment of anxiety. Our previous study showed that the ethyl acetate fraction of the plant had substantial anxiolytic action in mice. The present study was aimed to investigate the main constituent responsible for the observed effects.

Methods: Ethyl acetate extract was fractionated using column chromatography. Yielded fractions (FR1-4) at 50 mg/kg, and diazepam at 0.5 mg/kg were tested on the elevated plus-maze (EPM). Bioactive fraction was subjected to more purification on repeated chromatography columns. The isolated compound was identified based on ¹H-NMR, ¹³C-NMR, and ESI-Mass spectra.

Results: In the time spent in open arm, Fr1, and Fr3 did not show any significant effect on mice behavior, Fr2 marginally decreased the percentage of spent time by -8.4%, and Fr4 significant increased in the time spent in the open arms by 15.7%. In the open arm entry number Fr1, and Fr2 did not show any significant effect on mice behavior, Fr3 marginally increased the percentage of open arm entries by 7.9 %, and Fr4 significantly increased the open arm entry by 18.2%. The bioactive fraction (Fr.4) was subjected to more purification. Phytochemical analysis of Fr4 lead to the identification of 4',5,7-trihydroxyflavon (apigenin).

Conclusion: Semi-polar sub-fraction of S. lavandulifolia showed anxiolytic effects by increased time spent and the entry numbers in the open arms comparable to diazepam in the EPM model in mice. Bioactivity-guided isolation leads to the characterization of apigenin with flavone structure as its active constituent. Hence, it might be introduced as a new anxiolytic agent.

Implication for health policy/practice/research/medical education:
In this research bioactivity guided technique was successively used for separation of active fraction of Stachys lavandulifolia. Apigenin was the main component identified in the most active fraction suggested to be partly responsible of S. lavandulifolia reported effects on anxiety in traditional medicine. Hence, it might be used as antianxiety medication.


Introduction
Anxiety disorders are among the most prevalent of all mental disorders that exist in many forms and have a huge impact on the quality of life. Anxiety is a basic symptom of many psychiatric disorders and almost inevitable component of many medical and surgical conditions. Common treatment includes serotonin specific reuptake inhibitors (SSRIs), benzodiazepines, antidepressants and mono-amine oxidase inhibitors (MAOIs) which may cause important side effects like sedation, addiction, tachycardia, insomnia, decreased libido and ineffectiveness with their long-term use (1). To overcome the side effects of the synthetic drugs, researches have focused on medicinal plants like Valeriana officinalis (2), Matricaria recutita (3), Passiflora incarnata (4), Nepeta persica (5), Tilia americana (6), Tilia europaea (7) and Stachys lavandulifolia (8). Among them S. lavandulifolia Vahl. is an herbaceous wild plant native to Iran (9) which
is used in Iranian folk medicine as a mild sedative tea for reducing anxiety and for treatment of gastrointestinal disorders (10). In previous studies by the same authors on hydroalcohol total extract, ethyl acetate, petroleum ether, n-butanol and aqueous partitions of this plant, ethyl acetate partition showed suitable effects on anxiety. 

EPM test in mice (11). Therefore, in the current study a possible anxiolytic property of subfractions of ethyl acetate extract of S. lavandulifolia was evaluated and the bioactive fraction was subjected to purification to find the main constituent responsible for observed effects. For anxiety test, the elevated plus-maze was done which one of the most extensively used model for the investigation of drug effects on anxiety-related behavior in mice. It is based on the aversion and fear of mice for open spaces and is done on an elevated plus-maze with two closed and two opened arms. Anxiolytics increase the entry numbers and time spent on the open arms (12). The present study was aimed to investigate the anxiolytic effects of sub-fractions of ethyl acetate extract of S. lavandulifolia on elevated plus-maze (EPM) model of anxiety and investigating the main constituent responsible for the observed effects. 

Materials and Methods

Drugs and solutions

Solidified ethyl acetate extract or fractions (Fr.1–Fr.4) were weighed and prepared as 50 mg/mL stock solution in water using a trace amount of Tween 80. Diazepam (Sobhan Pharmaceutical Co., Iran) as standard drug was prepared as 0.5 mg/mL stock solution in distilled water and a trace amount of Tween 80. Vehicle (distilled water and a trace amount of Tween 80) was used as control. Unless stated, all the chemicals were purchased from Merck Company (Germany).

Plant material

The aerial parts of S. lavandulifolia were collected in April 2013 from Sharekord (Iran). The plant was identified and the voucher specimen (No: 1113) was deposited at the Herbarium unit of the Faculty of Pharmacy, Isfahan University of Medical Sciences, Isfahan, Iran.

Extraction and preliminary fractionation

The air dried plant material powder (1 kg) was macerated with 6 L ethyl acetate (72 h × 4 times) at room temperature and concentrated under reduced pressure (25.4 g). Ethyl acetate extract was applied on MPLC over RP-18 silica-gel cartridge (20 × 4 cm) using MeOH:H2O (7:3) as a solvent to remove chlorophylls and fats. Defatted fraction was fractionated by chromatography on an open glass column (7 X 80 cm) using normal silica gel (63-200 μm, 400 g) as adsorbent and mixtures of hexane: acetone as solvent to yield four fractions (Fr1, 90: 10; Fr2, 80: 20; Fr3, 70 30; Fr4, 50:50). The fractions were concentrated to dryness and stored at 0°C until use.

Results

Elevated plus-maze test

The EPM test has been described in details elsewhere (11,2). Briefly, the apparatus comprised of two open arms (35 × 5 cm) and two closed arms (30 × 5 × 15 cm) that extended from a common central platform (5 × 5 cm). The floor and the walls of each arm were wooden and painted black. The entire maze was elevated to a height of 60 cm above floor level as validated and described by Rabbani et al (5). Testing was conducted in a quiet room that was illuminated only by a dim light. Mice were given a single sc dose of the sub-fraction of extract 30 minutes before their placement on the EPM. To begin a test session, mice were placed on the open arm facing the open arm of the maze. An entry into an arm was defined as the animal placing all four paws over the line marking that area. The number of entries and the time spent in the open and closed arms and the number of entries in the middle arm was recorded during a 5-minute test period. The percentage of open arm entries (100 × open/total entries) was calculated for each animal. Between each trial, the elevated plus-maze was wiped clean with an alcohol (70%) and dried with napkin.

Isolation of bioactive fraction

Antianxiety effects of the resulted fractions: Fr.1 to Fr.4 were compared in vivo on EPM model in mice. According to the EPM pharmacological results, the bioactive fraction (Fr.4) was subjected to more purification. Based on primary identification by thin layer chromatography (TLC) analyses using cerium sulfate (1%) and natural product reagent which was applied on a SC6 polyamide column (2 X 20) using CHCl3:MeOH in a stepwise gradient manner (2→20 %) as solvent. The major sub-fraction which was eluted by CHCl3:MeOH (88:12), was purified on sephadex-LH 20 gel chromatography (2 X 60) using methanol as solvent and yielded compounds 1 (56 mg) as pure compound.

Statistical analysis

Statistical analysis was performed using one-way ANOVA with post hoc Duncan test. P < 0.05 was considered significant. All data were expressed as mean ± S.D.

Animals

Male Syrian mice weighing 25–35 g were housed in a cage with controlled room temperature at 22–25°C. Food and water were available. All experiments were carried out between 09:00 and 13:00. Each mouse received a single subcutaneous injection of drug or vehicle and was tested once in the EPM.

Animal (EPM) model of anxiety and investigating the main constituent responsible for observed effects. 

http://www.herbmedpharmacol.com
in vivo on EPM model in mice.

**Elevated plus-maze**

In the time spent in open arm in EPM test in comparison to control group, diazepam group significantly increased the percentage of spent time on open arms by 39.1% \( (P<0.05) \), ethyl acetate extract, Fr1, and Fr3 did not show significant effect on mice behavior, Fr2 marginally decreased the percentage of spent time by -8.4% \( (P<0.1) \), and Fr4 had significant increase in the time spent in the open arms by 15.7% \( (P<0.05; \text{ Figure 1A}) \).

In the open arm entry number in EPM test in comparison to control group, diazepam group was marginally significant by 6.9% \( (P<0.1) \), ethyl acetate extract, Fr1, and Fr2 did not show any significant effect on mice behavior, Fr3 marginally increased the percentage of open arm entries by 7.9% \( (P<0.1) \), and Fr4 significantly increased the open arm entry by 18.2% \( (P<0.01) \) (Figure 1B).

The summarized data in Table 1 represent the occupancy time as well as the entry numbers into the closed arms. None of the tested fractions or diazepam showed significant change versus control \( (P<0.05, \text{ Table 1}) \).

**Analysis of bioactive fraction**

In between group analysis, the most bioactive fraction (Fr4) with more time spent and entry number percentages were selected and subjected to purification methods using polyamide and sephadex gel chromatography. Finally, it yielded a pale yellowish solid with positive reaction to natural product reagent (1% methanolic diphenylboric acid-ethylamino ester) and FeCl₃ test. Its UV spectrum showed two absorptions maxima at 340.7 and 265.5 nm which is characteristic for flavones. Its ¹H-NMR resonances at δₓ (400 MHz, pyridine-d₆): 6.11 (1H, d, \( J = 2 \) Hz, H-6), 6.37 (1H, d, \( J = 2 \) Hz, H-8), 6.50 (1H, s, H-3), 6.84 (2H, d, \( J = 8 \) Hz, H-5,3’), and 7.76 (2H, d, \( J = 8.5 \) Hz, H-5,3’) indicated that isolated compound was a flavone derivative. Its negative ESI MS \( \text{m/z} \) 269 [M-1] and comparison of ¹H- and ¹³C-NMR data \( \delta \), (100 MHz, pyridine-d₆): 163.8, 104.4, 183.2, 106.5, 163.2, 100.5, 166.4, 95.3, 159.0, 122.8, 129.4, 117.3, 163.3, 117.3, 122.8] with literature and co-TLC with authenticated standards determined it as 4’,5,7-trihydroxyflavon known as apigenin (13). Phytochemical analysis of other fractions (Fr1-Fr3, and Fr5) was also performed in another research work which its data were published elsewhere (14).

**Discussion**

In our previous studies we showed that different extracts (petroleum ether, ethyl acetate, butanol and water) of *S. lavandulifolia* had various degrees of anxiolytic properties (11). From these extracts, ethyl acetate proved to be more suitable for further studies as it had fewer sedative properties with acceptable anxiolytic effects. In this study, in continuing our previous studies, the ethyl acetate extract was obtained and fractionated by column chromatography. Yielded fractions (Fr1-Fr4) at 50 mg/kg were tested again by elevated plus-maze test from which Fr4 significantly increased time spent and the entry numbers in the open arms. Analysis of the bioactive fraction, Fr4, characterized its major component as apigenin suggested

**Table 1.** Behavioral parameters recorded in the plus-maze from mice treated with various fractions of *S. lavandulifolia*

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Closed arm times (mean ± SEM)</th>
<th>Closed arm entries (mean ± SEM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>59.9 ± 8.4</td>
<td>71.8 ± 10.8</td>
</tr>
<tr>
<td>Diazepam (0.5 mg/kg)</td>
<td>20.8 ± 3.3</td>
<td>64.9 ± 15.2</td>
</tr>
<tr>
<td>Ethyl acetate extraction</td>
<td>60.2 ± 10.3</td>
<td>72.6 ± 18.4</td>
</tr>
<tr>
<td>Fr1 (50 mg/kg)</td>
<td>59.5 ± 9.3</td>
<td>63.6 ± 12.2</td>
</tr>
<tr>
<td>Fr2 (50 mg/kg)</td>
<td>68.3 ± 11.2</td>
<td>67.3 ± 9.8</td>
</tr>
<tr>
<td>Fr3 (50 mg/kg)</td>
<td>62.6 ± 9.7</td>
<td>63.9 ± 10.3</td>
</tr>
<tr>
<td>Fr4 (50 mg/kg)</td>
<td>44.2 ± 5.9</td>
<td>53.6 ± 7.8</td>
</tr>
</tbody>
</table>

Effects of diazepam, ethyl acetate extract and ethyl acetate extract sub-fractions (Fr1-4) of *S. lavandulifolia* on the percentage of time spent in the closed arms and the closed arm entries of the elevated plus-maze during a 5-minute test in mice. Fractions (each 50 mg/kg), diazepam (0.5 mg/kg), and control (water and trace amount of tween 80), were injected single subcutaneous 30 minutes prior to test. Data are presented as mean values (±SEM) from a group of six mice. * \( (P<0.05) \) compared with vehicle-treated control.
Isolation of apigenin from Stachys lavandulifolia with anxiolytic effects

Conflict of interests
The authors have no conflicts of interest.

Ethical considerations
The study was approved by the Ethical Committee of Isfahan University of Medical Sciences, Isfahan, Iran (393148). Ethical issues (including plagiarism, misconduct, data fabrication, falsification, double publication or submission, redundancy) have been completely observed by the authors.

Funding/Support
This project has been supported by Isfahan Pharmaceutical Sciences Research Center, Isfahan University of Medical Sciences (Grant No. 393148).

References
11. Rabbani M, Sajjadi SE, Jalali A. Hydroalcohol extract and fractions of Stachys lavandulifolia Vahl: effects on...


