Comparative Study of Anxiolytic-like Effect of a Novel Herbal Treatment with Diazepam on Balb/c mice

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Abstract

Background: The most widely prescribed medications for anxiety disorders are the Benzodiazepines; however, they have prominent side effects. Thus, interest in alternative medicine that affects the mind is growing. In Morocco folk medicine, the use of Mercurialis annua is commonly recommended for relief of anxiety, although no scientific information supporting this use is available and therefore we undertook the study to evaluate the anxiolytic potential the methanolic extract of Ma by using a battery of appropriate rodent test models.

Methods: The purpose of this study was to characterize the putative anxiolytic-like effects of methanolic extract prepared from the aerial parts of Ma using the Elevated Plus Maze (EPM) and Open Field tests (OF) in mice. The methanolic extract of Ma was administered orally to Balb/c mice, at graded doses; 1H prior to behavioural assessments.

Results: In the elevated plus-maze test, methanolic extract of Ma at 100 mg/kg showed an anxiolytic effect by increasing the time spent on and the number of entries into the open arms of the EPM comparable to standard anxiolytic drug, diazepam. In an open field test methanolic extract of Ma (100 mg/kg) increased the central area crossing, the time spent and number of rearing in the center of arena.

Conclusions: These observations suggest that Mercurialis annua might possess significant anxiolytic potential.

Keywords: Anxiety; Balb/c mice; Elevated plus maze; Open field; Methanolic extract of Mercurialis annua, Morocco.

Introduction

Anxiety disorders are among the most common psychiatric disorders that affect all age groups in the general population. In Morocco, anxiety disorders affect 9% of the population according to the national survey on mental health conducted by the Ministry of Health[1].

Anxiety disorders include Agoraphobia, Specific Phobia, Social Anxiety Disorder (Social Phobia), Panic Attack, Separation Anxiety Disorder, and Selective Mutism[2] and this trouble became a very important area of research interest in psychopharmacology. It’s well known that changes and control of emotion are related to dysregulation and neurochemical abnormalities in a certain specific pathway or neurotransmitter that is theoretically implicated in anxiety.

In a broader view of anxiety, there are at least five neurotransmitters that are “perturbed”, namely GABA, nor-adrenaline, serotonin, Corticotropic-Releasing Hormone(CRH), and cholecystokinin[3,4]. The interaction between these neurotransmitters is carefully orchestrated in the brain and changes in one neurotransmitter system elicit changes in another. All of these neurotransmitters have become important targets for therapeutic agents either already marketed or in development[5].

Abbreviations: Ma: Mercurialis annua, MeOH extract: methanolic extract, OF: Open field test, EPM: Elevated plus maze
Since the introduction of benzodiazepines in the 1960s, they have been the most commonly employed medicinal treatment for anxiety, these classical benzodiazepines act via the benzodiazepine receptors which are present on the GABA\textsubscript{A} pentameric complex\cite{6,7}. Current drug therapies for anxiety are relatively safe drugs for a short-term treatment of anxiety despite their drug dependence potential and side effects\cite{8,9}. Therefore, the development of other anxiolytic drugs without such adverse effects is important for the treatment of anxiety disorders and the study of medicinal plants could provide new therapeutic options.

There are numerous plants that have been used for their tranquilizing properties in Moroccan traditional medicine. Mercurialis annua L (Euphorbiaceae) is a wind-pollinated annual plant that occupies ruderal and roadside habitats throughout central and western Europe and around the Mediterranean Basin\cite{10}. The species is a winter annual in the Mediterranean region, and has long been known to have tranquilizing effects among the Moroccan people\cite{11,12}. Ethnobotanical reports attribute purgative, diuretic or antisyphilitic effects to the dried plant. Despite the wide spread use of Ma as an anxiolytic, there are no pharmacological data to support such effect, therefore, we explored the anxiolytic-like effects of Methanolic extract of Ma by using the elevated plus maze and open field tests, classic experimental models.

Materials and Methods

Animals
Balb/c mice of either sex (20 - 30 g) were employed in the present study. Animals were procured from the animal experimental center of Mohammed V. University, Medecine and pharmacy Faculty, Rabat. Animals were provided normal diet and water ad libitum and were maintained in a room with controlled temperature of 20 - 25 °C, and lighting (light/dark 12:12 hour) in polypropylene cages. The animals were acclimatized to the laboratory condition before experiments at least 1h. The animals were kept fasted 2h before drug administration, the Open field and the elevated plus maze were performed between 2 pm and 6 pm.

Plant Material
The aerial part of Mercurialis annua (Ma) was collected from the north of Morocco near the town of Wazzan (Jaaouna el Basra), with assistance of a traditional medical practitioner. The plant was authenticated by botanists of scientific institute Pr. M. Ibn Tatou and Pr. Halim Khammar. A voucher specimen (N° RAB78984) was deposited in the Herbarium of Botany Department of the Scientific Institute of Rabat.

Preparation of the Methanolic Extract
The aerial part was dried at room temperature and crushed. 700 g of plant material was extracted with six litre of methanol by maceration at room temperature (25 °C) over period of 48 hours. Methanol containing the extract was then filtered through Whatman paper and the solvent was vacuum-distilled at 60 °C in a rotary evaporator. The remaining extract was finally dried by desiccator. Final extract was a dark green paste, with 21.22 % dry weight. The residue was dissolved in water for final suitable concentrations.

Drugs and Chemicals
The methanolic extract of Ma was suspended in distilled water. Diazepam was diluted with saline to the required concentration before use. It is well known that benzodiazepines act as anxiolytics at low doses and that they induce sedation and muscle relaxant effects at higher doses\cite{14}. Therefore, we used diazepam (1 mg/kg; ip) as a positive control for anxiolytic-like effects.

Treatment Schedule
Mice were divided into five (5) groups (n = 6). Three groups treated orally (p. o.) with graded doses of methanolic extract of Ma (100, 200 and 400 mg/kg). One group received normal saline by the same route to serve as control and remaining group received diazepam (1 mg/kg i.p.). All drugs were freshly prepared before each experiment. The doses of extracts were calculated to administer 0.25 ml of the suspension of extracts to the mice of 20 g. The anxiolytic activity was examined by using the EPM and OF test. The trial was carried out 1H after the treatments.

Acute Toxicity Study
The present section of our study has been under taken to study the adverse or hazardous effects of methanolic extract from Ma, accordingly to determine the LD50, to establish the safety of methanolic extract of Ma in Balb/c mice as per OECD guidelines 423.

In acute toxicity studies, single dose of drug given in large quantity to determine immediate toxic effect. Acute toxicity studies are commonly used to determine LD50 of drug or chemicals and natural products. The procedure was followed as per OECD 423 guidelines\cite{15} (OECD/OCDE, 2002). The extract was administered orally at a dose of 2000 mg/kg body weight. Mice were kept under observed for 14 days to register possible mortality; their weights were registered and study their behavioral neurological toxicity.

Behavioral Paradigms
Elevated plus maze
The elevated plus maze (EPM) was first proposed as an animal model of anxiety by\cite{16}. Was first evaluated for rats\cite{17} and
later adapted for mice\cite{18}. Constructed of black–colored wooden planks consisting of two close arms, 50 cm x 10 cm x 40 cm, and two open arms, 50 cm x 10 cm, connected to a central platform (10 cm x 10 cm). Covered with a removable lid, such that the open or closed arms were opposite to each other\cite{17}. The maze was elevated to a height of 50 cm above the floor. During the experiment each mouse was placed in the central compartment facing one of the open arms. The number of entries and the time spent in open arms were recorded for 5 min. An entry was counted when all four paws of the mouse entered an open or closed arm. An increase in open arms entries and increase in time spent in open arms were interpreted as an index of potential anxiolytic activity. The number of closed arm entries was used to assess motor activity\cite{19}. All test sessions were taped by using a video camera.

**Open field test**

The open field apparatus was an opaque plexiglass cage (72 X 72 -cm) with walls 35 -cm in height where the floor was divided with white lines by 16 squares (18 X 18 -cm) of identical dimension. A digital video camera was installed above the cage to record the activity of the mice. The entire room, except the OF was kept dark during the experiment, in this study was evaluated both the general motor activity of the mice to discard hypoactivity, hyperactivity, or no changes associated with treatments, which could interfere with the behavioral activity of the mice in the elevated plus maze also the observed parameters were as follows: (a) total number of crossings, (b) central area crossings, (c) time spent in central area (d) central area rearing\cite{20,21}. An animal was placed in the OF and allowed to freely explore the apparatus for 5 min.

**Ethics approval**

The study was conducted in accordance with the accepted principles outlined in the “Guide for the Care and Use of Laboratory Animals” prepared by the National Academy of Sciences and published by the National Institutes of Health and all efforts were made to minimize animal suffering and the number of animals used. Ethics approval was obtained from the central laboratory of animal of Faculty of medicine and pharmacy from the Mohammed V University of Rabat.

**Statistical Analysis**

All the results were expressed as mean ± SEM. All statistical analysis was done using one way Analysis of Variance (ANOVA) followed by the Turkey’s post hoc test. P < 0.05 was considered as significant when compared to their respective control group.

**Results**

**Acute Toxicity Study**

Following oral administration methanolic extract of Mercurialis annua at a dose of 2000 mg/kg, P.O., animals were observed for signs of toxicity such as convulsions, hypothermia, hyperactivity, and grooming continuously for 2 h and for mortality up to 24 h after administration of the doses. The study of acute oral toxicity showed that oral administration of a single dose of 2000 mg/kg of methanol extract of Ma to mice caused no deaths within 14 days the study. No significant changes were observed in body weight and wellness parameters used for evaluation of toxicity.

Individual body weights of animals were recorded before the administration of drug on 1st day of the study and thereafter for 14 days of the experiment. Changes in the weight of individual animals were calculated and compared with that of the control animals.

It was noted a change in body weight of treated animals by extract, body weight of these animals underwent a regression in the first days of observation, and after 5 days we noticed that the reweight gradually increased.

**Figure 1:** Body weight in control (WC) and treated (WT) within 14 days.

LD50 Value: As per observations and calculations from Acute Oral Toxicity (OECD Guidelines 423), the LD50 value of Methanolic Extract of Ma was found to be more than 2000 mg/kg body weight.
Elevated Plus Maze

One way ANOVA indicated a significant difference among the groups, mice treated with graded doses of Ma (100, 200, 400 mg/kg) and Diazepam show significant activity when compared to control but group which receive 100 mg/kg of Ma spent more time on the open arm than other groups.

The analysis of the number of entries into open arms showed a significant effect among treatments **p < 0.01, ***p < 0.001, the post-hoc test showed that the total number of entries into the open arms was augmented (**p < 0.001) in the group treated with 100 mg/kg of Ma or 1 mg/kg of diazepam, when compared with others groups.

Open Field
Total crossing:

In the open field test, administration of Ma induced no changes in the total number of crossings. Diazepam showed a slight decrease of locomotor activity but this result is not significant.
Central area crossing:
The number of central area crossings was increased by Ma (100 mg/kg) (*p < 0.05 vs. vehicle) and diazepam (1 mg/kg) (**p < 0.001).

Figure 5: Effect of methanolic extract of Mercurialis annua 100, 200, 400 mg/kg on central area crossing.

Time spent in central area: Time spent in the central area was increased by Ma (100 mg/kg) (**p < 0.001 vs. control) and diazepam (1 mg/kg) (**p < 0.01 vs. control). The group treated with 100 mg/kg of methanolic extract of Ma, spent more time in the arena than other groups.

Figure 6: Effect of methanolic extract of Mercurialis annua 100, 200, 400 mg/kg on time spent on central area.

Central area rearing: Rearing behavior in central area was increased by Ma (100 mg/kg) (*p < 0.05 vs. control) and diazepam (1 mg/kg) (**p < 0.001 vs. control).

Figure 7: Effect of methanolic extract of Mercurialis annua 100, 200, 400 mg/kg on central area rearing.
The OF is capable to induce anxiety in mice and has been widely used to assess the behavioral effects of anxiety [24]. The OF with dose at 100 mg/kg.

In mice, the anxiolytic effects of the drug saturated at higher doses of 400 mg/kg as they did not show maximum activity compared to the respective control groups (P < 0.001 and P < 0.001, respectively). Figure 2. The values of the group treated with Ma at 100 mg/kg body weight were higher than that of the treatment with higher doses (200, 400 mg/kg) and diazepam 1 mg/kg.

In conclusion, the normalcy and insignificant changes in wellness parameters and body weights in the test of toxicity reveals the safety of methanolic extract of Ma at a dose of 2000 mg/kg body weight. Our results are promising since Ma might be considered as an alternative for the treatment of anxiety disorders to other medications currently used. Our findings also favor the position that conventional EPM and OF measurements are sufficient and reliable for detecting the anxiolytic effects of Ma extract. The phytoconstituent(s) responsible for the observed central effects has to be isolated and identified in future studies. Future study should be focused on the neurobiological mechanisms of action and possible interactions of Ma extract with classical neurotransmitters and neuromodulators responsible of pathology of anxiety.

Conflicts of Interest: All authors assert that none has any commercial or financial involvements that might present an appearance of a conflict of interest in connection with the submitted manuscript.
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