Original Article

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Effect of Matricaria recutita Hydroalcoholic Extract on Anxiety Behavior in Mice by Hole-Board Test

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| Article information | Abstract |
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| Article history: Received: 30 Aug 2012 Accepted: 20 Oct 2012 Available online: 12 Mar 2013 ZJRMS 2014; 16(3): 21-24 Keywords: Matricaria recutita Anxiety Mouse | Background: An anxiolytic effect of chamomile has been shown in various studies. In the previous study was indicated that the Iranian specious of chamomile, <i>Matricaria recutita</i> (<i>M. recutita</i>) hydro alcoholic extract acts sex dependent in the elevated plus maze. It showed anxiolytic effect in the presence and absence of male mice gonads but not in female mice. In this study we examined the anxiety model dependent of <i>M. recutita</i> in another unconditioned anxiety model, hole-board test, because there are various model for evaluating anxiety with specific properties. <i>Materials and Methods:</i> Adult male and female of N-MARI mice (N=120) were prepared and each sex divided into 5 groups (each group consist of 12 animals): control |
| *Corresponding author at: Department of Biology, Faculty of Sciences, Shahid Chamran University, Ahvaz, Iran E-mail: m.kesmati@scu.ac.ir | group, saline and 3 experimental groups that received different doses (10, 30, and 50 mg/kg, intraperitoneally) of <i>M. recutita</i> hydro alcoholic extract. Hole-board instrument was used to anxiety measurement, and delay time, the devour number and maintained time in the holes, as anxiety indices in this device, were evaluated. <i>Results:</i> There were not any significant differences between anxiety indices in control and saline groups in both sexes. <i>M. recutita</i> extract (10, 30 and 50 mg/kg via i.p.) reduced significantly an anxiety in both male and female mice and an anxiolytic effects of 30 mg/kg than the other doses were considerably higher. <i>Conclusion:</i> It seems an anxiolytic effect of <i>M. recutita</i> is independent to anxiety model and the similarity effect at male and female mice in this model emphasizes the validity of the model. |
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Introduction

A nxiety is a speared unpleasant sense and usually is imprecise care that is with one or many unpleasant bodily sense [1]. This disorder is as numbered century widespread disorders that everybody experienced it and freedom of it sometime is not possible. Multiple neural and hormonal centers play role in anxiety outbreak [2]. In this field GABA (Gamma Amino Butyric Acid), serotonin, dopamine and central areas, like amygdala and hippocampus involve in anxiety regulation [3-6]. Understand of intermediated mechanism in anxiety and finding suitable methods for treatment has been research topic for many investigators. In this way multiple chemical and plant drugs for treatment of anxiety have introduced [7, 8].

Side effect of anxiolytic drugs like benzodiazepines and barbiturates lead to usage of plant drugs be attractive for investigators [9]. Plant drugs have lower cost and side effect also there is another component with effective component in them that in more cases potentiate therapeutic effects and reduces side effects of them [10]. chamomile has been attractive for investigators that is a very aromatic and growing wild in champs and beside of pathways [11]. The flowers of this plant are used in industry and medicine. This plant from the past has been high usage in many clinical disorders as antiinflammatory, anti-spasm, analgesic drug and etc. [12, 13].

Studies show that has been attention to this plant for anti-anxiety effects and difference methods confirm antianxiety effects of it [12-16]. Yamada et al. shown that inhalation of Commomile extract gas in ovarectomized mice, lead to reduce plasma ACTH and reduce anxiety in them [17]. Also Della-Loggia et al. shown oral usage of chamomile in mice has anxiolytic effects [18]. In spite of above investigations, in previous study at this laboratory has been shown that hydro alcoholic extract of *M. recutita* flower in elevated plus maze test at the male and female mice significantly change anxiety indexes. In male mice with and without gonads had anxiolytic effect while in female didn't show this effect. So suggested that anxiolytic effect of *M. recutita* has been sex dependent [19].

Also in HPLC technique revealed that Iranian M. *recutita* has phytoestrogenic components such as Apigenin and Chrysin. And probably by the effect of these components has been observed difference in antianxiety effect of it [20]. By attention to exist of many kind of anxiety measurement models that every measure special aspect for answer to this ambiguity that the antianxiety effect of *M. recutita* is related to the model of evaluation of anxiety or not? In this study the effect of this plant drug investigate in another test. For this aim in this study hole-board apparatus as another unconditional model to make and measurement of anxiety was used [21-23]. This model in evaluation of anti-anxiety effect of *M. recutita* has been lowering noteworthy by investigators. So in this study with new insight, by hole-board apparatus, we investigate and compare the effect of hydro-alcoholic extract of *M. recutita* in adult male and female mice.

Materials and Methods

In this experimental study were used adult male and female N-MARI mice (3 months) weighting 32±2 g for male and 28±2 g for female that were purchased from Razi Institute of Hesarak of Kraj. All animals in this study kept in 22±2°C with wet percentage 30-50 and light/dark system 12 hours dark and 12 hours light with air ventilation. Animals randomly divided in special cages (the number of animal in each group was 12) and were fed with tap water and animal food from Dam Pars, Tehran, Iran. During the test and before it tried to appropriate quiet and out of stress place for animals. All procedures were carried out in accordance with institutional guidelines for animal care and use that accepted by Medical Science University of Ahvaz.

M. recutita flowers from Gol-Daru company (Isfahan, Iran) prepared and confirm by expert. In this investigation soak method was used to preparation of hydro alcoholic extract of *M. recutita* flower. For this reason evaporate branch with flowers was powdered by electrical instrument. In another stage 20 g of powder was get and put in 200 ml ethanol 70% (hydro alcoholic solvent) and mouth of vessel was banned. Prepared mixure kept for 48 hours in laboratory (shaked every 12 hours for few minutes) and then vessel components cleared by Wathman paper and funnel and after kept liquid in oven with 40°C temperature, dried powder of plant extract prepared and was used to preparation doses of it.

In this study to evaluation of anxiety hole-board apparatus was used. This apparatus is an unconditional model for evaluation of anxiety and make of flat square disk $(35\times35 \text{ cm})$ that has 16 regulate holes (4×4) . Diagonal of each hole was 3 cm and apparatus elevated 50 cm above the floor on the chair [23-25]. To evaluation the anxiety animal first sit in the center of apparatus and its behavior evaluated during 5 minutes. In this apparatus each animal was used just for one and there was no any prior instruction and learning. After each test apparatus cleaned by cotton for next test. In this study three anxiety indexes from this apparatus including latency time: time that animal for the first time plunged its head in one of the holes, head-dip in (count of the animal inserts the head into the hole in 5 min), head-dipping (duration of headdip in seconds).

In this method if latency time being higher, this show anxiety reducing also if head entries in holes and time spent were higher these show anxiety increased and inverse of these cases show anxiety reduced [23-25]. In this experiments two class containing 60 adult male and female mice were used that divided randomly into 5 groups containing 12 animals. Every of this class were containing: control, group receiving saline and 10, 30, 50 mg/kg of hydro alcoholic extract of *M. recutita*. Animals 30 minutes after i.p. injection of saline or hydro alcoholic extract of *M. recutita* placed in the center of hole-board apparatus and anxiety indexes were evaluated during 5 minutes for each of them [25-29].

Data analyzed by SPSS-16 and one way ANOVA with post hock LSD. In all experiment p < 0.05 used as significant level and bars show mean \pm SD.

Results

Statistical comparison between control and saline groups in anxiety indexes (latency time, head-dipping and headdip in the hole-board test show that saline injection hasn't any effect and there is no any difference between male and female (Fig. 1, 2, 3). Also figure 1, 2, 3 show that between saline group and

Also figure 1, 2, 3 show that between saline group and groups that receiving different doses of hydro alcoholic extract of *M. recutita* in both male and female mice there is significant increasing in latency time while number of head-dipping and head-dip in significantly reduced. All of these results showing *M. recutita* extract reduced the anxiety in male and female mice.

Statistical analysis between both of male and female in equal doses didn't show significant differences so we did not presented them in this section but seems anxiolytic effect of *M. recutita* on female mice partially was more effective.

Many studies confirmed that *Matricaria chamomilla* has anxiolytic effect [11, 16]. Della-Loggia et al. shown that oral usage of chamomile in mice can make anxiolytic effects [18]. Also Yamada et al. shown that inhalation of chamomile gas in ovariectomized female mice, lead to reduce blood plasma ACTH and reduce stress and anxiety in them [17]. In a clinical study identified that Apigenin isolated from chamomile is effective in some of drugs withdrawal syndrome including anxiety [30, 31]. Palladini et al. shown that Apigenin isolated from chamomile has anxiolytic effects and can act as stimulator neurotransmitter in central nervous system [32]. In addition in another similar study identified that Apigenin and Chrysin flavonoids extracted from chamomile show anxiolytic and benzodiazepines like effects [16, 20, 33].

Studies show that phytoestrogens, like flavonoids isolated from chamomile, can be responsible for anxiolytic effects of it in both of male and female mice and oral administration of them in mice lead to make anxiolytic effects in lower doses [18, 26]. In this study identified that hydro alcoholic extract of *M. recutita* has anxiolytic affects in male and female mice. Thought difference between male and female in equal doses was not significant so didn't present in this study but anxiolytic effects of *M. recutita* in female according to the level of significant partially seems is more effective.

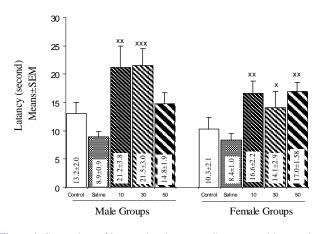


Figure 1. Comparison of latency time between saline group with control and groups receiving 10, 30, 50 mg/kg of hydro-alcoholic extract of Matricaria recutita in hole-board apparatus. Significant level for every male and female group has been shown in compared to saline group of same sex. The number of animal in each group was 12. ($\times = p < 0.05$, $\times \times = p < 0.01$, $\times \times \times = p < 0.001$)

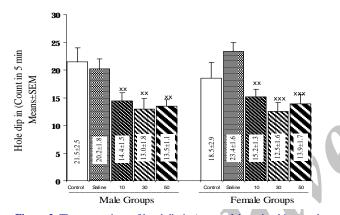


Figure 2. The comparison of head dip in (count of the animal inserts the head into the hole in 5 min) between saline group with control and groups receiving 10, 30, 50 mg/kg of hydro-alcoholic extract of Matricaria recutita has in hole-board apparatus. Significant level for every male and female group has been shown in compared to saline group of same sex. The number of animal in each group was 12. ($\times = p < 0.01$, $\times \times = p < 0.001$)

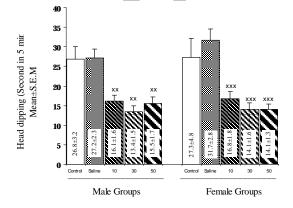


Figure 3. The comparison of head-dipping (duration of head-dip in seconds), between saline group with control and groups receiving 10, 30, 50 mg/kg of hydro-alcoholic juice of *Matricaria recutita* has in holeboard apparatus. Significant level for every male and female group has been shown in compared to saline group of same sex. The number of animal in each group was 12. ($\times = p < 0.01$, $\times \times = p < 0.001$)

Today it has been demonstrated that flavonoids materials isolated from botanic drugs like: Apigenin that with high affinity bind to benzodiazepines site of GABA-A receptors [34]. Studies show that Apigenin by binding to these receptors in central nervous system, play its sedative and anxiolytic effect [35]. So it possible that powerful effect of *M. recutita* in reduce anxiety is due to this material. In addition identified that Apigenin as flavonoid material has its best anxiolytic effect at dose of 2 mg/kg in elevated plus maze test [26], and it has more similarity with our finding, actually maximum anxiolytic effect of *M. recutita* is at dose of 30 mg/kg in animals.

Main point in this study in contrary with our pervious finding, about the effect of M. recutita extract on mice in elevated plus maze test, is the effect of this extract on female mice anxiety in hole-board test [19]. This difference probably can be related with different aspects of anxiety measuring and strong and weak points of these tests. For example it has been identified that elevated plus maze test as unconditional model of anxiety in rodent because it need animal travels on the surface of open arm of apparatus make unusual anxiety in animals and so the validity of some results of it can have poor confidence in compared to others tests [36]. On the other hand Borrin et al. said that of two evaluated parameters anxiety and locomotor activity, in male genus, anxiety, and in female genus, locomotor activity, affected in anxiety test when usage of effective materials [37]. So the reason of didn't change in anxiety behaviors in female in elevated plus maze test can be due to different in the kind of utilized test.

So according to the findings from this study can conclude that the *M. recutita* probably with containing flavonoids has anxiolytic effect in male and female mice in hole-board test and affecting central nervous system through the change in GABA-A receptors activity or/and stress hormones and makes this plant drug good candidate to produce anti-anxiety drug. Also because of similarity in anxiolytic effect of this plant drug in both of male and female mice, this model to measurement of anxiety has higher validity in compared to elevated plus maze test to evaluation of anxiety and investigation of the effects of drugs.

Acknowledgements

The authors wish to express their gratitude to the Research Council of Shahid Chamran University for their financial supports (Grant No. 90/02/18672).

Authors' Contributions

The first author had role in design and manuscript writhing, the second one in experimental work and the third one in manuscript edition.

Conflict of Interest

The authors declare no conflict of interest.

Funding/Support

Shahid Chamran University, Ahvaz.

References

- 1. Sylvers P, Lilienfeld SO, La Prairie JL. Differences between trait fear and trait anxiety: Implications for psychopathology. Clin Psychol Rev 2011; 31(1): 122-137.
- Ohman A. Fear and anxiety: Evolutionary, cognitive, and clinical perspectives. In: Lewis M, Haviland-Jones JM, Feldman-Barrett L. Handbook of emotions. 3rd ed. New York: Guilford Press; 2010: 573-593.
- Ali BH, Al-Qarawi AA. An evaluation of drug used in the control of stressful stimuli in domestic animals. Acta Vet Brno 2002; 71: 205-216.
- Altshuler LL, Hendrick V, Cohen LS. An update on mood and anxiety disorders during pregnancy and the postpartum period. Prim Care Companion J Clin Psychiatry 2000; 2(6): 217-222.
- 5. Clement Y, Chapouthier G. Biological bases of anxiety. Neurosci Biobehav Rev 1998; 22(5): 623-633.
- 6. Nutt DJ, Bell CJ, Malizia AL. Brain mechanisms of social anxiety disorder. J Clin Psychiatry 1998; 59(17): 4-9.
- Hatano VY, Torricelli AS, Giassi AC, et al. Anxiolytic effects of repeated treatment with an essential oil from Lippia alba and (R)-(-)-carvone in the elevated T-maze. Braz J Med Biol Res 2012; 45(3): 238-43.
- Karbalay-Doust S, Dehghani F, Panjehshahin MR, et al. Effects of hydroalcoholic extract of Matricaria chamomilla on serum testosterone and estradiol levels, spermatozoon quality, and tail length in rat. Iran J Med Sci 2010; 35(2): 122-128.
- Griffiths RR, Johnson, MW. Relative abuse liability of hypnotic drugs: A conceptual framework and algorithm for differentiating among compounds. J Clin Psychiatry 2005; 66(9): 31-41.
- Kennedy DO, Little W, Haskell CF and Scholey AB. Anxiolytic effects of a combination of Melissa officinalis and Valeriana officinalis during laboratory induced stress. Phytother Res 2006; 20(2): 96-102.
- 11. Cemek M, Kaga S, Simsek N, et al. Antihyperglycemic and antioxidative potential of Matricaria chamomilla L. in streptozotocin-induced diabetic rats. J Nat Med 2008; 62(3): 284-293.
- 12. Gardiner P. Chamomile (Matricaria recutita, Anthemis nobilis). Available from: http://www.mcp.edu/herbal/ default.htm.
- 13. Nemecz G. Herbal pharmacy: Chamomile. U.S. Pharmacist 2000; 23: 115-123.
- Avallone R, Zanoli P, Puia G, et al. Pharmacological profile of apigenin, a flavonoid isolated from Matricaria chamomilla. Biochem Pharmacol 2000; 59(11): 1387-1394.
- Johnston GAR. Dietary chemicals and brain function. J Proc Royal Soc N S W 2003; 135: 57-71.
- Zanoli P, Avallone R, Baraldi M. Behavioral characterization of the flavonoid apigenin and chrysin. Fitoterapia 2000; 71(1): 117-123.
- Yamada K, Miura T, Mimaki Y and Sashida Y. Effect of inhalation of chamomile oil vapour on plasma ACTH level in ovariectomized rat under restriction stress. Biol Pharm Bull 1996; 19(9): 1244-1246.
- Della-Loggia R, Tubaro A, Redaelli C. [Evaluation of the activity on the mouse CNS of several plant extracts and a combination of them] Italian [Abstract]. Riv Neurol 1981; 51(5): 297-310.
- 19. Pourmehdi-Rad G, Kesmati M. Comparison of anxiolytic effect of matricaria recutita in male and female mice in the

presence and absence of gonads. Zah J Res Med Sci 2009; 11(2): 19-28.

- 20. Kesmati M, Zande-Moghadam A, Hoshmand-Nia A and Abasizadeh Z. Comparison between of Matricaria recutita aqueous and hydrolacholic extract on morphine withdrawal signs in the presence and absence of tamoxifen. Iran J Med Arom Plants 2009; 25(2): 170-181.
- Dhawan K, Kumar S, Sharma A. Anti-anxiety studies on extracts of Passiflora incarnata Linneaus. Ethnopharmacol 2001; 78(2-3): 165-170.
- 22. Romanova D, Grancai D, Jozova B, et al. Determination of apigenin in rat plasma by high-performance liquid chromatography. J Chromatogr A 2000; 870(2): 463-467.
- Silva AL, Elisabetsky E. Interference of propylene glycol with the hole-board test. Med Bio Res 2001; 34(4): 545-547.
- 24. Flint J. Animal models of anxiety and their molecular dissection. Semin Cell Dev Biol 2003; 14(1): 37-42.
- 25. Monnier C, Lalonde R. Elevated plus maze and hole-board exploration in lurcher mutant mice. Brain Res 1995; 702(1-2): 169-172.
- 26. Lund TD, Lephart ED. Dietary soy phytoestrogens produce anxiolytic effects in the elevated plus maze. Brain Res 2001; 913(2): 180-184.
- 27. Long SF. Preventing and treating insomina. Drug Topics 2000; 144(13); 49-57.
- 28. Aikey JL, Nyby JG, Anmuth DM and James PJ. Testosterone rapidly reduces anxiety in male house mice (Mus musculus). Horm Behav 2002; 42(4): 448-460.
- Belzung C, Griebel G. Measuring normal and pathological anxiety-like behaviour in mice. Behav Brain Res 2001; 125(1-2): 141-149.
- Breinholt V, Hossaini A, Svendsen GW, et al. Estrogenic activity of flavonoid in mice, the importance of strogen receptor distribution, metabolism, bioavailability. Food Chem Toxicol 2000; 38(7): 555-564.
- 31. Rasmussen P. A role for phytotherapy in the treatment of benzodiazepine and opiate drug withdrawal. Herb Med 1997; 3(1): 11-21.
- Paladini AC, Marder M, Viola H, et al. Flavonoids and the central nervous system: From forgotten factors to potent anxiolytic compounds. J Pharm Pharmacol 1999; 51(5): 519-526.
- Wolfman C, Viola H, Paladini A, et al. Possible anxiolytic effects of chrysin, a central benzodiazepine receptor ligand isolated from Passiflora coerulea. Pharmacol Biochem Behav 1994; 47(1): 1-4.
- Losi G, Puia G, Garzon G, et al. Apigenin modulates GABAergic and glutamatergic transmission in cultured cortical neurons. Eur J Pharmacol 2004; 502(1-2): 41-46.
- 35. Campbell EL, Chebib M, Johnston GA. The dietary flavonoids apigenin and (-)-epigallocatechin gallate enhance the positive modulation by diazepam of the activation by GABA of recombinant GABA (A) receptors. Biochem Pharmacol 2004; 68(8): 1631-8.
- 36. Shepherd JK, Grewal SS, Fletcher A, et al. Behavioural and pharmacological characterization of the elevated "zero-maze" as an animal model of anxiety. Psychopharmacology (Berl) 1994; 116(1): 56-64.
- Bourin M, Petit-Demouliere B, Dhonnchadha BN and Hascoet M. Animal models of anxiety in mice. Fundam Clin Pharmacol 2007; 21(6): 567-74.

Please cite this article as: Kesmati M, Izadi L, Mard-Soltani M. Effect of matricaria recutita hydroalcoholic extract on anxiety behavior in mice by hole-board test. Zahedan J Res Med Sci (ZJRMS) 2014; 16(3): 21-24.