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Effect of *Punica granatum* on behavior in rats

Azra Riaz and Rafeeq Alam Khan*

Department of Pharmacology, Faculty of Pharmacy, University of Karachi, Pakistan

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The importance of diet in depression is now increasingly recognized, since balanced diet not only reduces the risk of various chronic diseases but also promotes normal development and functioning of all systems including brain. Hence the present study was designed to observe the effect of three doses of *Punica granatum* 2, 5 and 8 ml/kg on the behavioral changes in rats, particularly anxiolytic and antidepressant activity. During 15 days study, experimental paradigm for anxiety and depression were assessed twice using open field, elevated plus maze and forced swimming test. In open field, distance travelled, number of centre entries and number of rearing's were increased. In elevated plus maze, number of open arm entries were increased and in forced swimming test, there was decrease in duration of immobility and increase in duration of climbing. All these changes were significant at moderate and high doses of *P. granatum*. Thus it may be suggested that *P. granatum* possesses anxiolytic and antidepressant activity at moderate and high doses.

Key words: *Punica granatum*, anxiolytic, antidepressant.

INTRODUCTION

Anxiety is the most common of all mental disorders. The persistent anxiety is manifested by symptoms from at least three out of four categories that is, motor tension, characterized by muscle aches restlessness; autonomic hyperactivity, characterized by sweating, dizziness, increased heart rate; apprehensive expectation, characterized by fear and vigilance characterized by difficulty in concentration (Brown et al., 2001; Fricchione, 2004). In recent years, use of herbal remedies and dietary supplements has increased to treat mild to moderate anxiety disorders (Saeed et al., 2007). Open field test and elevated plus maze are best experimental methods for the measurement of anxiety (Herskin and Jensen, 2002; Mansouri et al., 2014).

Depression could be defined as a multifaceted condition

of psychosomatic, neuroendocrine and somatic symptoms difficult to reproduce in animals (Petit-Demouliere et al., 2005), but since research in humans is limited, animal models of depression have been developed. Many symptoms of depression cannot be easily measured in laboratory rodents for example, feelings of worthlessness, suicide tendency etc., but some behavioral tests have shown to be very effective in evaluating depressive symptoms and are classically used to predict the antidepressant effect of new medications. They also provide theoretical useful models to study neurobiological and genetic mechanisms underlying depressive behavioral changes.

The existence of numerous behavioral tests to measure depression in rodents reflects the heterogeneity of

*Corresponding author. E-mail: rkhan1959@gmail.com.

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depression like symptoms. Hence forced swimming test is the classical paradigms used to evaluate behavioral misery. Hopelessness, reported as a common trait of depression in humans, is mimicked in rodents by the paradigm of learned helplessness. Thus forced swimming test is considered to be useful for investigating depressive state in human since the behavioral immobility of animals during forced swimming has been reported to reproduce some aspects of human depression (Porsolt et al., 1977, 1978; Wilner, 1984; Yoshimura and Yamakawa, 2000).

Force swim test (FST) was than modified by different investigators (Petit-Demouliere et al., 2005; Carbajal et al., 2009), while it has also been reported that regional skin temperature changes and prolonged immobility like symptoms observed in ovariectomized mice are very identical to depressive hot-flush symptoms in women (Mori-Okamoto et al., 2004). Thus force swim test is a currently used popular model for depression due to low cost and reliability (Petit-Demouliere et al., 2005).

Pomegranate (*Punica granatum L.*, Punicaceae) is a globally consumed fruit and contains high level of phytochemicals, including polyphenols (punicalagin, ellagic acid, anthocyanins) and flavonoids (quercetin, kaempferol). *P. granatum* also contains testosterone, estrogen and estrogen like compounds (Gil et al., 2000; Langley, 2000; Kohno et al., 2004; Lansky et al., 2007; Hong et al., 2008; Wang et al., 2010; Fischer et al., 2011). These compounds have variety of activities for example, ellagic acid has anxiolytic activity (Girish et al., 2013). Progesterone has been shown to regulate anterior pituitary level of follicle stimulating hormone (Conner et al., 1999). Testosterone has been shown to have anti-depressant like activity (Buddenberg et al., 2009). Polyphenols have hypoglycemic and hypolipidemic effects (Yin et al., 2011). Flavonoids have anti-inflammatory activity (Kim et al., 2004; Garcia-Lafuente et al., 2009). However studies regarding exploratory behavior and antidepressant-like action are very rare. Thus present study was designed to evaluate the effects of orally administered *P. granatum* juice on reduced immobility during forced swim test in rats and on anxiety and motor action using elevated plus-maze and open field procedures.

MATERIALS AND METHODS

Animals

Study was carried out on adult male Wister rats with mean body weight of 220 ± 10 g. Animals were kept under controlled condition of temperature $23 \pm 2^\circ\text{C}$ and humidity 50 to 60% throughout the experiment in a 12/12 h light and dark cycle with free access to rat chow and tap water. Five rats were housed in each plastic cage measuring $81 \times 46 \times 41$ cm. The use of animals during the study was in accordance with the National Institute of Health (NIH) guide for the care and use of Laboratory Animals (National Research Council, 1996) and approved by the Board of Advance Studies and Research University of Karachi.

Punica granatum Juice

The *P. granatum* (Pomegranate) was purchased from local market, identified by center of plant conservation, University of Karachi. The voucher specimen no P.G 11-12 was deposited in Department of Pharmacognosy, University of Karachi. The fruits were peeled and squeezed to yield fresh juice which was than filtered and administered through oral route in three doses that is, 2, 5 and 8 ml/kg, respectively according to body weight.

Drug treatment

All rats were divided into six groups each comprising of ten animals. One group served as control and received sterile water, three groups were treated with *P. granatum* at 2, 5 and 8 ml/kg and two other groups received standard drugs imipramine and diazepam in the dose of 25 and 3 mg/kg, respectively according to body weight (Carbajal et al., 2009). Sterile water, juices and standard drugs were administered on once daily basis for 15 days through oral route.

Open field test

Open field test (OFT) is used to assess locomotion and anxiety related behavior (Prut and Belzung, 2003) and is also the measure of exploration (Walsh and Cummins, 1976). The open field was made up of Plexi glass, 75 cm long, 75 cm wide and 40 cm high in the shape of quadratic box. Black lines were drawn on floor, which divides the floor into twenty-five (15×15 cm) squares. A central square (30×30 cm) was drawn around the middle centre squares to count number of central entries of animal. Cross section length of squares was 21 cm. The open field was located in test room, illuminated by 60 watt light bulb. All tests were recorded by camera (SONY HANDYCAM-DCR-DVD755/E) from the top view of the field. *P. granatum* juice, sterile water and diazepam were administered daily between 8.0 to 10.0 am orally for 15 days. OFT for all animal groups was performed twice on 8th and 15th day, just 1 h after administration of drugs.

Procedure

All rats received individual application and were handled by their tail. Time of exploration of field by the animal was 30 min, after which rats were returned in their home cages and arena was cleaned with 70% ethanol solution. Animals were exposed to the apparatus twice that is, 8th and 15th day of drug administration by placing into the center of the open field maze. Variables measured were, (1) distance traveled by the animal in the field by line crossing with all four paws, (2) total number of entries in central square (30×30 cm), when animal crossed one of the central square lines with all four paws, (3) total number rearing-frequency with which animal stood or rising up on hind legs with the forelegs against the wall of maze or in the air, (4) duration of rearing's - time spent by animal in rising up or standing position.

Elevated plus-maze

Elevated plus-maze (EPM) was used to assess the anxiety related behavior (Hogg, 1996) of *P. granatum*. Animals used in OFT underwent testing in EPM just 30 min after open field test. The EPM consisted of two open arms and two close arms placed opposite to each other, arranged around (10×10 cm) central platform. Open arms were 50 cm long and 10 cm wide. While close arms were 50 cm long, 10 cm wide and 38 cm high with open roof.

Table 1. Effect of *P. granatum* and diazepam on behavior of rats in open field.

Group	Days	Parameter			
		No. of rearing's	Duration of rearing's (s)	No. of centre entries	Distance travelled (cm)
Control (Sterile water)	8	50.0±5.41	200.1±47.26	4.9±0.86	2137.9±151.87
	15	51.4±5.58	207.1±48.46	5.6±0.97	2329.8±187.51
<i>P. granatum</i> 2 ml/kg	8	53.4±5.05	203.3±46.74	4.7±0.84	2190.0±147.83
	15	54.5±5.09	210.4±47.43	5.2±0.84	2646.9±317.49
<i>P. granatum</i> 5 ml/kg	8	62.9±2.12*	210.0±47.54	7.2±0.69*	2786.2±382.00*
	15	63.7±2.18*	213.8±47.73	7.7±0.65*	3231.0±334.96*
<i>P. granatum</i> 8 ml/kg	8	63.4±2.40*	219.4±50.74	7.7±0.66*	2762.8±164.00*
	15	64.1±2.34*	220.4±50.06	8.2±0.53*	3225.3±350.31*
Diazepam	8	62.6±2.95*	235.9±48.67	7.6±0.45*	1852.3±264.28
	15	63.1±2.92*	236.5±48.72	8.1±0.37*	1983.8±264.43

n=10, Values are mean ± SEM. *P ≤ 0.05 significant as compared to control.

Procedure

Animals behavior in all test was recorded by means of a video camera (SONY HANDYCAM-DCR-DVD755/E) mounted 100 cm above the maze to have the top view. Animals were placed into the central platform, facing one of the open arms. During the test, rats were allowed to move freely in maze and explore the environment for 5 min. Variables measured were, (1) spent time in open arms, (2) spent time in close arms, (3) number of entries, when animal entered with all four paws into the open or closed arms.

Forced swimming test

Forced swimming test (FST) is used to assess depressive-like behavior. Test was done twice following administration of *P. granatum* juice, imipramine and sterile water to respective groups on 15th day 1 h just after the EPM, marked as day-1 (pre-test phase), and at 16th day, twenty-four hours after the 1st FST, marked as day-2 (test-phase).

Procedure

FST was performed in Plexiglass cylinder 46 cm tall and 20 cm wide. The cylinder had been filled with water at 25°C up to 20 ± 2 cm from bottom. The forced swim apparatus was situated in a sound isolated and dimly illuminated room. On the day-1, rats were placed individually into the water for a 15 min period. After the completion of pre-test phase, animals were removed from the water and then placed under a light heating lamp in a plastic cage for about 15 to 20 min to make them dry. Same experimental conditions of FST were applied to animals after twenty four hours that is, placed into the water for 15 min, water was replaced after each animal testing. All tests were recorded by camera (SONY HANDYCAM-DCR-DVD755/E) from the front view of the whole cylinder. Variables measured were, (1) duration of immobility that is, time at which animal showed lack of motion, except only those movement necessary to keep his head above water, (2) duration of

swimming that is, time at which animal showed vigorous movements with fore paws in water, (3) duration of climbing that is, time at which animal showed vigorous movements with fore paws and hind paw along the walls of cylinder (Buddenberg et al., 2009).

Statistical analysis

All data was expressed as the mean ± standard error of mean (SEM), data was analyzed using superior performance statistical software (SPSS) version 20. ANOVA followed by post hoc was performed for comparisons of values with control. Values of p ≤ 0.05 were considered statistically significant and p ≤ 0.005 highly significant.

RESULT

Table 1 shows the effect of *P. granatum* on behavior of rats in open field. There was significant increase in No. of rearing's, No. of centre entries and distance travelled at 5 and 8 ml/kg than control groups on respective days. While no significant change was observed in any parameters at 2 ml/kg than control. Animals received diazepam showed significant increase in number of rearing's and central entries on both 8th and 15th day than as control. Table 2 shows the effect of *P. granatum* on behavior of rats in elevated plus maze. There was significant increase in number of entries in open arm but no significant changes were observed in time spent on open arm, close arm and no. of entries in close arm at 5 and 8 ml/kg both on 8th and 15th day than control. However no significant changes were observed in any parameters at 2 ml/kg than control. Animals received diazepam showed significant increase in number of entries in open arm and

Table 2. Effect of *P. granatum* and diazepam on behavior of rats in elevated plus maze.

Groups	Days	Parameter			
		No. of entries in open arm	Time spend in open arm (s)	No. of entries in close arm	Time spend in close arm (s)
Control	8	3.5±0.70	125.0±23.0	6.5±0.76	175.0±23.0
	15	4.1±0.73	140.0±24.8	5.7±0.84	160.0±24.8
<i>P. granatum</i> 2 ml/kg	8	5.0±0.81	122.0±20.48	5.6±1.13	178.0±20.48
	15	5.1±0.79	138.0±22.64	5.1±1.03	162.0±22.64
<i>P. granatum</i> 5 ml/kg	8	6.2±1.04*	101.0±22.03	6.1±1.10	199.0±22.03
	15	7.1±0.90*	132.0±26.02	6.8±0.91	168.0±26.02
<i>P. granatum</i> 8 ml/kg	8	6.7±1.08*	114.0±21.86	6.2±1.08	186.0±21.86
	15	7.0±1.01*	126.5±25.10	6.5±1.14	173.5±25.10
Diazepam	8	6.6±0.66*	157.0±17.9	3.2±0.55*	143.0±17.9
	15	7.0±0.85*	210.0±16.3*	2.4±0.60*	90.0±16.3*

n=10, Values are mean ± SEM. *P ≤ 0.05 significantly different as compared to control.

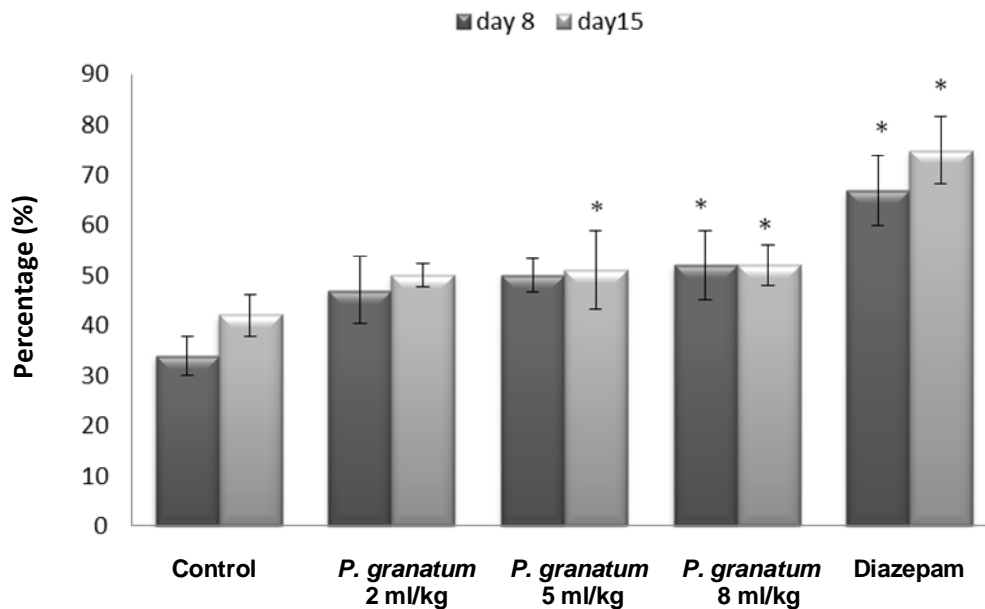


Figure 1. Percentage of entries into the open arm in Elevated Plus Maze. mean ± S.E.M% of open arm entries (percent increase from control). *P ≤ 0.05 significantly different as compared to control.

and time spent in open arm, while there was significant decrease in number of entries in close arm and time spent in closed arm than control.

Figure 1 shows percentages of entries in to the open arm by control, *P. granatum* at all three doses and diazepam in elevated plus maze. Percentages at 8th day were 34, 47, 50, 52, 67% and at 15th day percentages were 42, 50, 51, 52 and 74% of control, 2, 5, 8 ml/kg and diazepam, respectively. Change in percentage in 8 ml/kg

and diazepam group at both days and 5 ml/kg at day 15 were significantly increased as compare to control. Figure 2 shows percentages of time spent in the open arm by control, *P. granatum* at all three doses and diazepam in elevated plus maze. Percentages at 8th day were 42, 41, 34, 38, 52% and at 15th day 47, 46, 44, 42, 70% of control, 2, 5, 8 ml/kg and diazepam, respectively. Change in percentage in diazepam group at both days was significantly increased as compare to control.

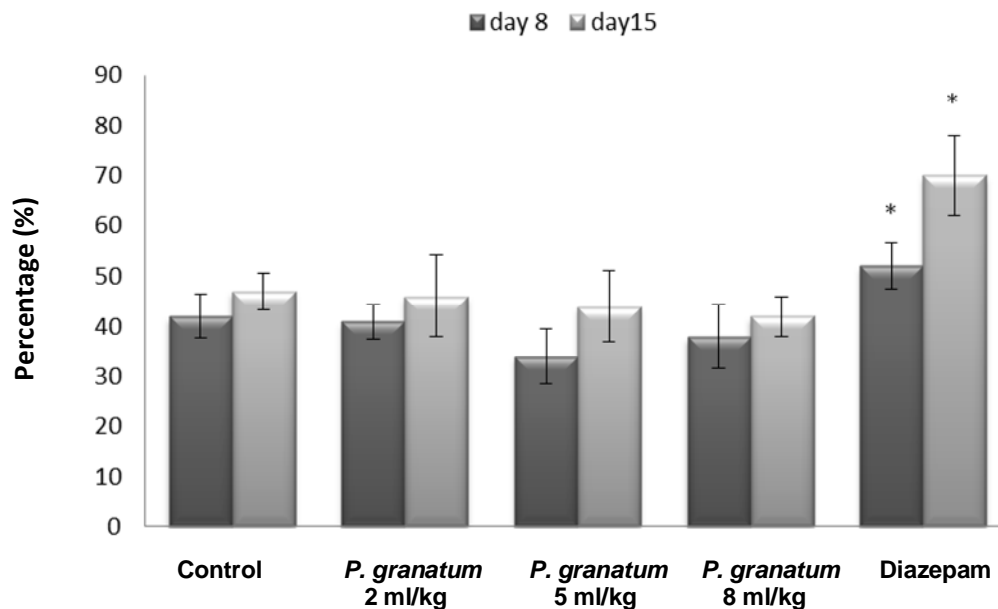


Figure 2. Percentage of time spent in the open arm in elevated plus maze. Mean \pm SEM% of time in open arm (percent increase from control). * $P \leq 0.05$ significantly different as compared to control.

Table 3. Effect of *P. granatum* and imipramine on behavior of rats in forced swimming test

Groups	Day	Parameters		
		Immobility duration (s)	Climbing duration (s)	Swimming duration (s)
Control	1	151.5 \pm 13.39	67.4 \pm 8.83	681.1 \pm 12.56
	2	160.5 \pm 21.60	66.4 \pm 6.43	673.1 \pm 18.20
<i>P. granatum</i> 2 ml/kg	1	116.3 \pm 03.69	82.6 \pm 3.09	701.1 \pm 02.69
	2	122.3 \pm 04.91	84.6 \pm 4.26	693.1 \pm 05.58
<i>P. granatum</i> 5 ml/kg	1	105.1 \pm 16.32*	97.4 \pm 10.58*	697.5 \pm 17.24
	2	108.1 \pm 16.89*	96.4 \pm 12.45*	695.5 \pm 18.34
<i>P. granatum</i> 8 ml/kg	1	105.5 \pm 03.60*	98.40 \pm 06.58*	696.1 \pm 08.86
	2	109.5 \pm 04.77*	93.7 \pm 07.00*	696.8 \pm 09.97
Imipramine	1	89.1 \pm 06.88**	71.0 \pm 08.66	739.9 \pm 14.02*
	2	87.7 \pm 08.91**	68.1 \pm 03.01	744.2 \pm 47.17*

n=10, Values are mean \pm SEM. * $P \leq 0.05$ significantly different as compared to control. ** $P \leq 0.005$ highly significant as compared to control.

Table 3 shows the effect of *P. granatum* on behavior of rats in forced swimming test. There were significant decrease in duration of immobility and significant increase in duration of climbing at 5 and 8 ml/kg during first and second exposure to forced swimming than control. While highly significant decrease in duration of immobility and significant increase in duration of swimming were observed by imipramine during first and second exposure of forced swimming test than control. However no significant changes in duration of immobility, duration of climbing and duration of swimming were observed at 2 ml/kg on both exposures as compare to

control.

Figure 3 shows percentage decrease in immobility duration by control, *P. granatum* at all three doses and imipramine in force swimming test. Percentages during first exposure were 17, 14, 11, 11, 10% and during the second exposure percentages were 18, 15, 12, 13, 10% of control, 2, 5, 8 ml/kg and imipramine, respectively. Decrease in percentage in 5, 8 ml/kg and imipramine group at both exposure were significant as compare to control. Figure 4 shows percentage increases in climbing duration by control, *P. granatum* at all three doses and imipramine in force swimming test. Percentage during first

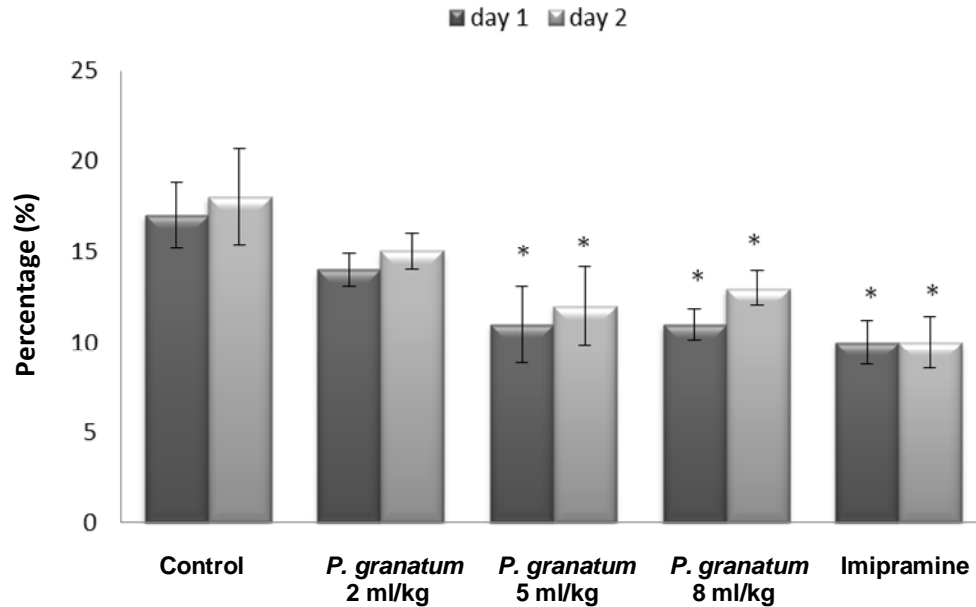


Figure 3. Effect of *P. granatum* and imipramine on immobility inhibition in FST. Mean \pm SEM, inhibition immobility (percent decrease from control).
* $P \leq 0.05$ significantly different as compared to control.

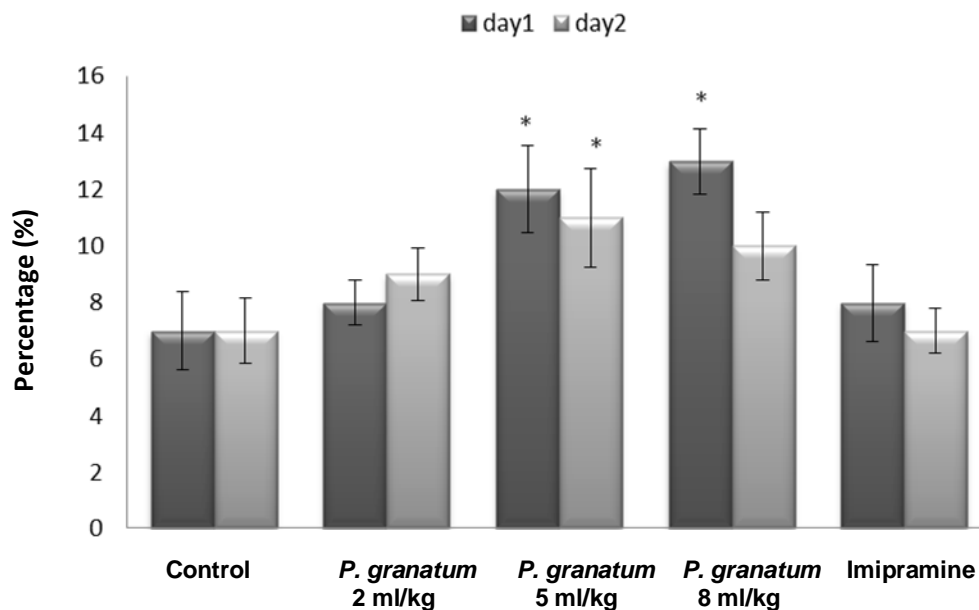


Figure 4. Effect of *P. granatum* and imipramine on percentage of climbing duration. Mean \pm SEM, increase climbing duration (percent increase from control).
* $P \leq 0.05$ significantly different as compared to control

first exposure were 7, 8, 12, 13, 8% during first exposure and 7, 9, 11, 10, 7% during the second exposure of control, 2, 5, 8 ml/kg and imipramine, respectively. Increase in percentage in 5 and 8 ml/kg group at both exposure were significant as compared to control. Figure 5 shows percentage increases in swimming duration

by control, *P. granatum* at all three doses and imipramine in force swimming test. Percentages during first exposure were 76, 78, 77, 76, 82 and 75, 74, 76, 77 and 83% during the second exposure by control, 2, 5, 8 ml/kg and imipramine, respectively. Increase in percentage in imipramine at both exposures was significant as compared

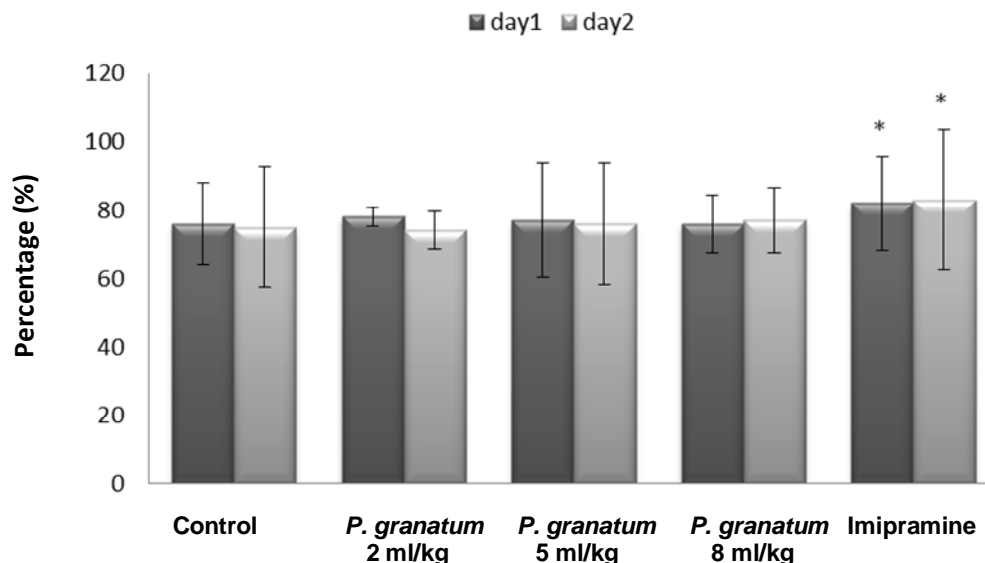


Figure 5. Effect of *P. granatum* and Imipramine on percentage increase in swimming duration Mean \pm SEM, increase swimming (percent increase from control). *P \leq 0.05 significantly different as compared to control

to control.

DISCUSSION

In present study *P. granatum* was tested for its spontaneous motor activity, anti-anxiety and anti-depressant actions in healthy male rats. These behavioral changes were studied using open field test, elevated plus maze and forced swimming tests. *P. granatum* was administered in three different doses 2, 5 and 8 ml/kg. Turk et al. (2008) used *P. granatum* juice in the dose of 1 ml and Patel et al. (2008) assessed safety of the juice. Punicalagin, major water soluble ellegatannins of *P. granatum*, found in its juice vary from 1500 to 1900 mg/L and its daily intake of 6 oz/60 kg individual weight may range from 4.4 to 5.6 mg/kg (Patel et al., 2008).

Open field test (OFT) and elevated plus maze (EPM) were used to evaluate anxiolytic effect, as behavior animal models (Mansouri et al., 2014). In OFT the 5 and 8 ml/kg showed significant increase in number of centre entries, number of rearing's and distance traveled both on 8th and 15th day as compared to control. The behavior of increase in number of centre entries, number of rearing's at 5 and 8 ml/kg were comparable to that of diazepam treated group. These observed effects of *P. granatum* may be due to its increase spontaneous, locomotor and anxiolytic activity, since the number of central square entries is the measure of exploration behavior and anxiety. A high frequency/duration of these behaviors indicates high exploratory behavior and low anxiety levels (Walsh and Cummins, 1976; Brown et al., 1999; Bailey and Crawley, 2009). The spontaneous

activity was measured by distance traveled and rearing's. The rearing is an index of locomotor activity (Alves et al., 2005), while increase in distance traveled is an indication of increase in motor activity as well as stimulation of central nervous system (Kennett et al., 1987; Czech, 2002). *P. granatum* significantly increased the number of centre entries, rearing's and distance travelled. This stimulation in motor activity can be interpreted in term of central nervous system (CNS) stimulant effect.

In EPM *P. granatum* showed increase in total entries at all three doses; however significant increase in number of entries in open at 5 and 8 ml/kg both on 8th and 15th day was observed. This increase in number of entries to open arm were quite similar to standard drug diazepam. Comparison between parameter "no. of entries in open arm" with "no. of entries in close arm" 5, 8 ml/kg and diazepam group on both 8th and 15th day showed more no. of entries in open arm as compared to close arm. While comparison between the parameter "time spend in open arm" with "time spend in close arm" only diazepam treated group showed more time spent in open arm as compared to the close arm. All mentioned significant changes at 5 and 8 ml/kg are comparable to diazepam group. Thus it could be suggested that the increase are likely be due to general stimulatory activity. Hence, anti-anxiety behavior can be determined by measuring spontaneous motor activity that is, total or open arm entries (Pellow and Sandra, 1986; Budzynska et al., 2013; Mansouri et al., 2014).

In EPM, data of percentages of total activity gives us a picture of differences in locomotor activity between control, treated and diazepam groups. Among treated group 5 and 8 ml/kg showed highest locomotor activity

that could be compared with diazepam group. Increases in locomotor behaviors are the measurement of CNS stimulant activity and decreased anxiety-like behavior. All these behavioral changes of *P. granatum* can be taken in terms of CNS stimulant and anxiolytic effect. Since there is no way to confidently differentiate a drug effect as either stimulant or anxiolytic due to a similar phenotype, and there certainly can be an overlap in terms of underlying mechanisms. Hence increase in distance traveled is an indication of increase in motor activity as well as stimulation of central nervous system (Kennett et al., 1987; Czech, 2002) and number of rearing's is an index of locomotor activity (Alves et al., 2005). Thus, forced swimming test (FST) is considered to be useful for investigating depressive state, since the behavioral immobility of animals during forced swimming has been reported to reproduce some aspects of human depression (Porsolt et al., 1977, 1978; Wilner, 1984; Yoshimura and Yamakawa, 2000). Duration of immobility is used to quantify overall anti-depressant or depressive like effect of the drugs (Taiwo et al., 2012). Present study also characterized the effect of the *P. granatum* on rats in FST following 15 day treatment. To obtain full antidepressant effect, rats were subjected to forced swimming at the end of dosing period that is, 15th and 16th day, and was considered as day 1 and day 2, respectively (Buddenberg et al., 2009). Hence it is critical to perform FST on repeated administration in rat model (Mora et al., 2005).

Result of the present study reveals that 5 and 8 ml/kg treated group significantly reduce the duration of immobility, which is comparable to that of imipramine. 5 and 8 ml/kg treated group also showed significant increase in duration of climbing, as a compensatory mechanism of reduction in immobility, whereas imipramine showed significant increase in duration of swimming. Decrease in immobility induced by imipramine like drugs is generally accompanied by increase in swimming, whereas climbing duration was not affected by these drugs (Barros and Ferigolo, 1998). Comparison between parameter "immobility duration" with "climbing duration" 2 ml/kg treated group immobility duration is much greater than climbing duration, whereas in case of 5, 8 ml/kg and imipramine treated group on both day 1 and 2 showed less differences. Percentages of reduction in immobility at 5 and 8 ml/kg were also quite similar to imipramine. This comparison reveals similar response of these two doses of *P. granatum* with that of the response of standard drug imipramine. Hence 5 and 8 ml/kg revealed a trend, indicating an antidepressant like effect.

Comparison of "day 1" with "day 2" *P. granatum* treated rats showed reduction in mobility and increase in climbing behavior at day 1, the first FST exposure as compare to day 2 second FST exposure. This possibly may be due to the reduced flavonoids level in body, since flavonoids absorbed from the gastrointestinal tract and excreted either unchanged or as metabolites in the urine or feces

(Cook and Samman, 1996; Sousa et al., 1996). Result of the present study showed significant decrease in the duration of immobility in FST, increase in the distance travelled, number of central entries in OFT and number of entries in open arm in EPM by *P. granatum* in dose dependent manner, suggesting the need to work on higher dose, to obtain desired effects. The antidepressant action might be due to the presence of flavonoid (Ohlsson et al., 2010). Major flavonoid found in *P. granatum* is ellagic acid, since ellagic acid is reported to produce anxiolytic action (Girish et al., 2013). These results of *P. granatum* may be due to presence of testosterone in it (Kim et al., 2004). Since there is evidence that testosterone could attenuate immobility in healthy rats and act as antidepressant by increasing central dopaminergic and 5-hydroxytryptaminergic metabolism (Buddenberg et al., 2009). Another study suggests that anti-depressant like effect might be due to the estrogen which is an important component of *P. granatum* (Kim et al., 2002). Hence, role of estrogen or estrogen like compound as antidepressant have been well documented (Mori-Okamoto et al., 2004).

From the results of the present study, it may be concluded that *P. granatum* is most effective in moderate and high doses for its CNS stimulant, anxiolytic and antidepressant effects. However further studies on more different doses are required to reveal the role of essential substances in all their observed effects.

Conflict of interest

Authors declare that they have no competing interests.

REFERENCES

- Alves R, de Carvalho Barbosa JG, Benedito Campana MA (2005). High and low rearing subgroups of rats selected in the open field differ in the activity of K⁺-stimulated p-nitrophenylphosphatase in the hippocampus. *Brain Res.* 1058(1-2):178-182.
- Barros HM, Ferigolo M (1998). Ethnopharmacology of imipramine in the forced-swimming test: gender differences. *Neurosci. Biobehav. Rev.* 23(2):279-286.
- Bailey KR, Crawley JN (2009). Anxiety-Related Behaviors in Mice. In: Buccafusco JJ (ed). *Methods of Behavior Analysis in Neuroscience*. 2nd edition. Boca Raton (FL): CRC Press. Chapter 5.
- Brown RE, Corey SC, Moore AK (1999). Differences in measure of exploration and fear in mhc-congenic C57BL/6J and B6-H-2K Mice. *Behav. Gen.* 26:263-271.
- Brown TA, O'Leary TA, Barlow DH (2001). Generalized anxiety Disorder. Chapter 4. *Clinical Handbook of Psychological Disorders*, Third Edition: A Step-by-Step Treatment Manual. pp. 154-208.
- Buddenberg TE, Komorowski M, Ruocob LA, de Souza Silva MA, Topic B (2009). Attenuating effects of testosterone on depressive-like behavior in the forced swim test in healthy male rats. *Brain Res. Bull.* 79:182-186.
- Budzynska B, Boguszewska-Czubarra A, Kruk-Slomka M, Skalicka-Wozniak K, Michalak A, Musik I, Biala G, Glowinski K (2013). Effects of imperatorin on nicotine induced anxiety and memory related responses and oxidative stress in mice. *Physiol. Behav.* 122:46-55.
- Carbajal D, Ravelo Y, Molina V, Mas R, Arruzazabala ML (2009). D-004, a lipid extract from royal palm fruit, exhibits antidepressant

- effects in the forced swim test and the tail suspension test in mice. *Pharmacology, Biochemistry and Behavior* 92:465–468
- Conner JL, Wade MF, Edwards DP, Mahesh VB (1999). Progesterone and regulation of the follicle-stimulating hormone (FSH- β) gene. *Steroids* 64:592–597.
- Cook NC, Samman S (1996). Flavonoids—Chemistry, metabolism, cardio protective effects and dietary sources. *J. Nutritional Biochem.* 7(2):66–76.
- Czech DA (2002). A simple integrated circuit device for measuring distances traveled and determining speed in open-field environments. *Pharmacol. Biochem. Behav.* 72(1-2):73–75.
- Fischer UA, Carle R, Kammerer DR (2011). Identification and quantification of phenolic compounds from pomegranate (*Punica granatum* L.) peel, mesocarp, aril and differently produced juices by HPLC-DAD–ESI/MSn. *Food Chem.* 127:807–821.
- Fricchione G (2004). Generalized anxiety disorder. *N. Engl. J. Med.* 351:675–82.
- Garcia-Lafuente A, Guillamon E, Villares A, Rostagno MA, Martinez JA (2009). Flavonoids as anti-inflammatory agents: implications in cancer and cardiovascular disease. *Inflamm. Res.* 58:537–552
- Gil MI, Tomas-Barberan FA, Hess-Pierce B, Holcroft DM, Kader AA (2000). Antioxidant Activity of Pomegranate Juice and its Relationship with Phenolic Composition and Processing. *J. Agric. Food Chem.* 48:4581–4589
- Girish C, Raj V, Arya J, Balakrishnan S (2013). Involvement of the GABAergic system in the anxiolytic-like effect of the flavonoid ellagic acid in mice. *Eur. J. Pharmacol.* 710(1-3):49–58.
- Herskin MS, Jensen KH (2002). Effects of open field testing and associated handling vs. handling alone on the adrenocortical reactivity of piglets around weaning. *Anim. Sci.* 74:485–491.
- Hong MY, Seeram NP, Heber D (2008). Pomegranate polyphenols down-regulate expression of androgen-synthesizing genes in human prostate cancer cells overexpressing the androgen receptor. *J. Nutritional Biochem.* 19:848–855.
- Hogg S (1996). A review of the validity and variability of the elevated plus-maze as an animal model of anxiety. *Pharmacol. Biochem. Behav.* 54(1):21–30.
- Kennett GA, Dourish CT, Curzon G (1987). Antidepressant-like action of 5-HT_{1A} agonists and conventional antidepressants in an animal model of depression. *Eur. J. Pharmacol.* 134(3):265–274.
- Kim HP, Son KH, Chang HW, Kang SS (2004). Anti-inflammatory Plant Flavonoids and Cellular Action Mechanisms. *J. Pharmacol. Sci.* 96:229–245.
- Kim ND, Mehta R, Yu W, Neeman I, Livney T, Amichay A, Poirier D, Nicholls P, Kirby A, Jiang W, Mansel R, Ramachandran C, Rabi T, Kaplan B, Lansky E (2002). Chemopreventive and adjuvant therapeutic potential of pomegranate (*Punica granatum*) for human breast cancer. *Breast Cancer Res. Treat.* 71(3):203–17.
- Kohn H, Suzuki R, Yasui Y, Hosokawa M, Miyashita K, Tanaka T (2004). Pomegranate seed oil rich in conjugated linolenic acid suppresses chemically induced colon carcinogenesis in rats. *Cancer Sci.* (95)6:481–486.
- Langley P (2000). Why a pomegranate? *BMJ* 321:1153–1154.
- Lansky PE, Newman RA (2007). *Punica granatum* (pomegranate) and its potential for prevention and treatment of inflammation and cancer. *J. Ethno Pharmacol.* 109:177–206.
- Mansouri MT, Soltani M, Naghizadeh B, Farbood Y, Mashak A, Sarkaki A (2014). A possible mechanism for the anxiolytic-like effect of gallic acid in the rat elevated plus maze. *Pharmacol. Biochem. Behav.* 117:40–46.
- Mora S, Diaz-Veliz G, Lungenstrass H, Garcia-Gonzalez M, Coto-Morales T, Poletti C, De Lima TCM, Herrera-Ruiz M, Tortoriello J (2005). Central nervous system activity of the hydroalcoholic extract of *Casimiroa edulis* in rats and mice. *J. Ethnopharmacol.* 97(2):191–197.
- Mori-Okamoto J, Otawara-Hamamoto Y, Yamato H, Yoshimura H (2004). Pomegranate extract improves a depressive state and bone properties in menopausal syndrome model ovariectomized mice. *J. Ethnopharmacol.* 92:93–101.
- Ohlsson A, Ulleras E, Cedergreen N, Oskarsson A (2010). Mixture effects of dietary flavonoids on steroid hormone synthesis in the human adrenocortical H295R cell line. *Food Chem. Toxicol.* 48(11):3194–3200.
- Patel C, Dadhaniya P, Hingorani L, Soni MG (2008). Safety assessment of pomegranate fruit extract: Acute and subchronic toxicity studies. *Food Chem. Toxicol.* 46:2728–2735.
- Pellow S, Sandra FE (1986). Anxiolytic and anxiogenic drug effects on exploratory activity in an elevated plus-maze: A novel test of anxiety in the rat. *Pharmacol. Biochem. Behav.* 24(3):525–529.
- Petit-Demouliere B, Chenu F, Bourin M (2005). Forced swimming test in mice: a review of antidepressant activity. *Psychopharmacology* 177:245–255.
- Porsolt RD, Anton G, Blavet N, Jalfre M (1978). Behavioral despair in rats: a new model sensitive to antidepressant treatments. *Eur. J. Pharmacol.* 47(4):379–91.
- Porsolt RD, Le Pichon M, Jalfre M (1977). Depression: a new animal model sensitive to antidepressant treatments. *Nature* 266:730–2.
- Prut L, Belzung (2003). The open field as a paradigm to measure the effects of drugs on anxiety-like behaviors: a review. *Eur. J. Pharmacol.* 463(1-3):3–33.
- Saeed SA, Bloch RM, Antonacci DJ (2007). Herbal and Dietary Supplements for Treatment of Anxiety Disorders. *Am. Fam. Physician* 76(4):549–556.
- Sousa MC, Braga RC, Cintra BAS, de Oliveira V, Andrade CH (1996). In silico metabolism studies of dietary flavonoids by CYP1A2 and CYP2C9. *Food Res. Int.* 50(1):102–110.
- Turk G, Sonmez M, Aydin M, Yuce A, Gur S, Yudsell M, Aksu HE, Aksoy H (2008). Effects of pomegranate juice consumption on sperm quality, spermatogenic cell density, antioxidant activity and testosterone level in male rats. *Clin. Nutr.* 27:289–296.
- Taiwo Adefunmilayo E, Leite Franco B, Lucena Greice M, Barros M, Silveira D, Silva Monica V, Ferreira Vania M (2012). Anxiolytic and antidepressant-like effects of *Melissa officinalis* (lemon balm) extract in rats: Influence of administration and gender. *Indian J. Pharmacol.* 44(2):189–192.
- Walsh RN, Cummins RA (1976). The open-field test: a critical review. *Psychol. Bull.* 83:482–504.
- Wang R, Ding Y, Liu R, Xiang L, Du L (2010). Pomegranate: Constituents, Bioactivities and Pharmacokinetics. *Fruits, Vegetables and Cereal Sci. Biotechnol.* 4(2):77–87.
- Wilner P (1984). The validity of animal models of depression. *Psychopharmacology* 83:1–16.
- Yin P, Zhao S, Chen S, Liu J, Shi L, Wang X, Liu Y (2011). Hypoglycemic and hypolipidemic effects of Polyphenols from Burs of *Castanea mollissima* Blume. *Molecules* 16:9764–9774.
- Yoshimura H, Yamakawa K (2000). Animal models for behavioral disorder in females. *Brain Sci.* 22:49–54.