

# Study Effect of *Satureja bachtiarica* Alcoholic Extract on Some Components of Complement System and IgM in Rat Serum

Meysam Khodadadi<sup>1</sup>, Hamed Soleyman Dehkordi<sup>1,2</sup>, Hamid Iranpour Mobarakeh<sup>1</sup>,  
Faham Khamesipour<sup>2\*</sup>, Mohsen Jafarian Dehkordi<sup>3</sup>

<sup>1</sup>Faculty of Veterinary Medicine, Shahrekord Branch, Islamic Azad University, Shahrekord, Iran

<sup>2</sup>Young Researchers and Elite Club, Shahrekord Branch, Islamic Azad University, Shahrekord, Iran

<sup>3</sup>Department of Pathology, Faculty of Veterinary Medicine, Islamic Azad University, Shahrekord Branch, Shahrekord, Iran

**Abstract** This plant is antioxidant, Carminative, Astringent, Fungicide, mucolytic, antibacterial, strong stomach tonic and Anthelmintic. With regard to valuable properties of *Satureja bachtiarica* plant, we decided to investigate effects of *Satureja bachtiarica* alcoholic extract on some components of complement system (C<sub>1</sub>inhibitor, CH50, C<sub>3</sub> and C<sub>4</sub>) and serum IgM in rat. In this study *Satureja bachtiarica* extract at concentrations of 200 and 400mg/kg was given to animals in two treatment groups for 30 days and the third group as whiteness received no compound. In this study after measuring components of complement system and IgM a relative increase was observed in the amount of complement system components and IgM compared to the control group.

**Keywords** *Satureja bachtiarica* L., Complement system, Serum, IgM, Rats

## 1. Introduction

*Satureja* is a plant of dicotyledons class and lamiaceae family, herbaceous, annual and its height reaches up to 10-30 cm. Stems are distributed with a grayish-green appearance and leaves are long, narrow, linear and sharp with their surfaces having numerous fine points containing essence. This plant is distributed in west Iran (Azerbaijan) and north Iran (Firouzkouh) regions (Rechinger, 1982). *Satureja* has tannin, fatty substances, different sugars and essence about 20% and its color is yellow or bright brown (Lawrence, 1981). Also this plant has compounds named carvacrol and parasimen. According to Lawrence book, the value of *Satureja* essence is due to carvacrol amount and phenolic aromatic substances as fresh that is reminder of *Zataria multiflora* and *Origanum majorana* (Lawrence, 1978; Iranpour Mobarakeh et al., 2014). Its essence is an effective anti-diuretic that is due to presence of carvacrol at concentration of 35-40% in wild plant and 65% in cultivable plants (Soodi et al., 2012; Iranpour Mobarakeh et al., 2014). Leaves contain uric acid and there is 272 calories, 9 g water, 6.7g protein, 5.9 g fat, 68.7 g carbohydrates, 140 mg phosphorus 2.32 mg calcium and 37.9 mg iron in each

100g of dried *Satureja* (Duke, 1981). Extract of this plant is gargled to decrease tonsils hyperemia (Abdollahi et al., 2003; Zareii et al., 2014). In cases of neuralgia and rheumatic pains, this plant is also used in therapeutic baths to reduce pain (Iranpour Mobarakeh et al., 2014; Sharafzadeh, 2014). Also massaging vertebral column with plant creams containing extract of this plant accompanied by foot baths and vaginal steaming is very effective to reduce orgasmic dysfunction in women (Lewis and Elvin-Lemis, 1977). *Satureja* is useful to treat diarrhea and its pomade with olive oil is useful for different types of abdominal cramps (Yousefzadi et al., 2013; Iranpour Mobarakeh et al., 2014). This plant also has antioxidant, antidiabetic, antibacterial and anticoagulant property and reduces weight and triglyceride (Nazari et al., 2006). *Satureja* is used in treatment of runny nose, colic, otitis, sclerosis and spasms (Duke and Wain, 1981). In this research, we investigated the effects of the *Satureja bachtiarica* alcoholic extract on some components of complement system such as (C<sub>3</sub>, C<sub>4</sub>, CH50 and C<sub>1</sub> inhibitor) and IgM in rat serum.

## 2. Material and Methods

This study was conducted in 2013 in the Islamic Azad University of Shahrekord Branch laboratory. Some components of the complement system (C<sub>3</sub>-C<sub>4</sub>-CH50-C<sub>1</sub> inhibitor) and IgM were measured in Al-Mahdi Medical

\* Corresponding author:

dr\_faham@yahoo.com (Faham Khamesipour)

Published online at <http://journal.sapub.org/ijaf>

Copyright © 2014 Scientific & Academic Publishing. All Rights Reserved

Diagnostic Laboratories of Shahrekord, Iran. Sample of *Satureja bachtiarica* plant leaves were collected and alcoholic extract of the mentioned plant was prepared in the laboratory. Drying and extracting *Satureja bachtiarica* was prepared from medicinal plant research center. After cutting extraction of the plant extract was performed by using Alcoholic distillation method by the rotary device (British pharmacopeia, 1988).

## 2.1. Animals

In this research 30 female white wistar rats prepared from laboratory animals breeding center of university laboratory with weight range of  $215 \pm 15$  g, were maintained in standard cages and had access to food and water. According to ethical code available in university and by considering ethical issues relating to animals, the study tried to avoid any case including annoyance, unnecessary use of animals or even losses during the testing. Rats were divided into three classes of ten and then extract dose was determined through preliminary experiments in groups 1 and 2 and the control group received no compound (Hartwell, 1971). The group 1 and the group 2 received respectively 200mg/kg and 400mg/kg of *Satureja bachtiarica* alcoholic extract and the control group received no compound. Prescription of extract in groups continued for 30 days and after completion of this period animals were anesthetized intraperitoneally with 100mg/kg of Ketamin hydrochloride and 16 mg/kg of 2% xylazine. Blood sample was collected by the Cardiac Puncture Technique (Sumiko et al., 2001). Then its serum was separated by centrifugation at 200 rpm for (time) and components of complement system and IgM were measured. All experiments were carried out under ethical guidelines of the Islamic Azad University of Shahrekord Branch, for the care and use of laboratory animals.

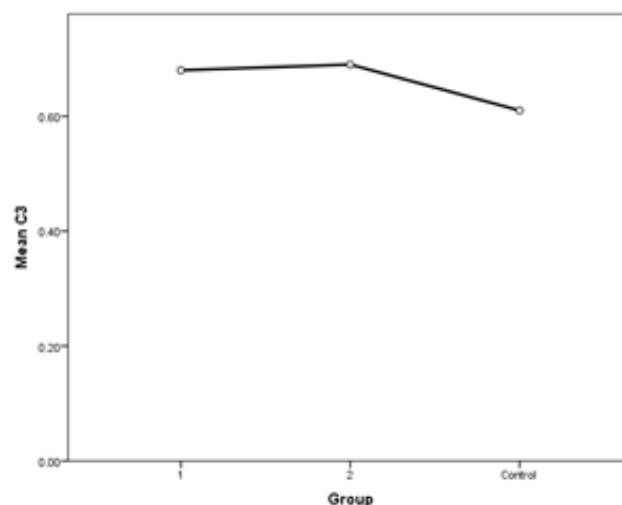
## 2.2. Statistical Analysis

Findings were statistically analyzed by means of SPSS software 18.0 (SPSS Inc., Chicago, IL, USA) and significance levels ( $p \leq 0.05$ ) were compared by means of Duncan tests.

## 3. Results

The amount of serum  $C_3$ ,  $C_4$ , CH50,  $C_1$ inhibitor and IgM in the groups are shown in Table 1. The amount of serum  $C_3$  in group one was  $0.68 \pm 0.15$  mg/dl and did not show

significant differences to the control group ( $0.61 \pm 0.59$  mg/dl) but showed a relative increase ( $p > 0.05$ ). The amount of serum  $C_3$  in group two was  $0.69 \pm 0.25$  mg/dl and showed a relative increase to the control group ( $p > 0.05$ ) (Figure 1).



**Figure 1.** The effect of *Satureja bachtiarica* alcoholic extract level of  $C_3$  of rat blood serum

The amount of serum  $C_4$  in group one was  $0.11 \pm 0.03$  mg/dl and did not show significant decrease to the control group ( $0.09 \pm 0.05$  mg/dl) but showed a relative increase ( $p > 0.05$ ). The amount of serum  $C_4$  in group two was  $0.10 \pm 0.01$  mg/dl and showed a relative increase to the control group ( $p > 0.05$ ) (Figure 2).

The amount of serum CH50 in group one was  $9.1 \pm 1$  gr/L and did not show significant decrease to the control group  $5.1 \pm 1.3$  gr/L. The amount of serum CH50 in group two was  $9.2 \pm 2.5$  gr/L and didn't show significant decrease to the control group but showed a relative increase ( $p > 0.05$ ) (Figure 3).

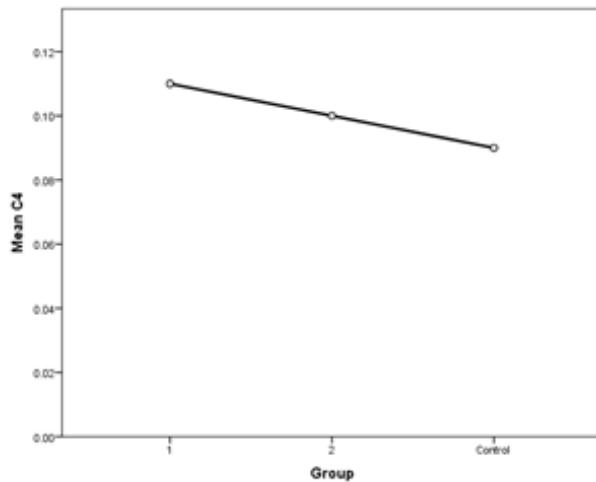
$C_1$ inhibitor: The amount of serum  $C_1$ inhibitor in group one was  $0.068 \pm 0.26$  gr/L and did not show significant decrease to the control group ( $0.05 \pm 0.01$  gr/L). The amount of serum  $C_1$ inhibitor in group two was  $0.065 \pm 0.11$  and didn't show significant difference to the control group but showed a relative increase ( $p > 0.05$ ) (Figure 4).

IgM: The amount of serum IgM in group one was  $0.34 \pm 0.08$  gr/L and did not show significant decrease to the control group ( $0.23 \pm 0.16$  gr/L). The amount of serum IgM in group two was  $0.39 \pm 0.10$  and didn't show significant difference to the control group but showed a relative increase ( $p > 0.05$ ) (Figure 5).

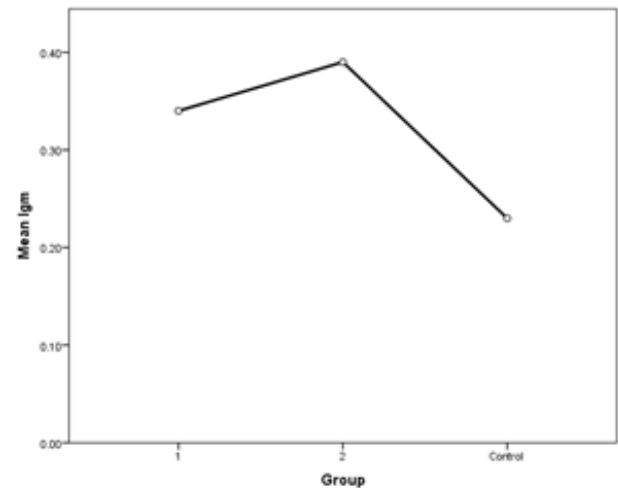
**Table 1.** The *Satureja bachtiarica* alcoholic extract effect on some components of complement system and IgM in rat serum

Groups	<i>Satureja bachtiarica</i> alcoholic extract was used	$C_3$ (SD±Mean)	$C_4$ (SD±Mean)	CH50 (SD±Mean)	$C_1$ inhibitor (SD±Mean)	IgM (SD±Mean)
Control	-	$0.61 \pm 0.59^a$	$0.09 \pm 0.05^a$	$5.1 \pm 1.3^a$	$0.05 \pm 0.01^a$	$0.23 \pm 0.16^a$
Group 1	200 mg/kg	$0.68 \pm 0.15^a$	$0.11 \pm 0.03^a$	$9.1 \pm 1^a$	$0.068 \pm 0.26^a$	$0.34 \pm 0.08^a$
Group 2	400 mg/kg	$0.69 \pm 0.25^a$	$0.10 \pm 0.01^a$	$9.2 \pm 2.5^a$	$0.065 \pm 0.11^a$	$0.39 \pm 0.10^a$

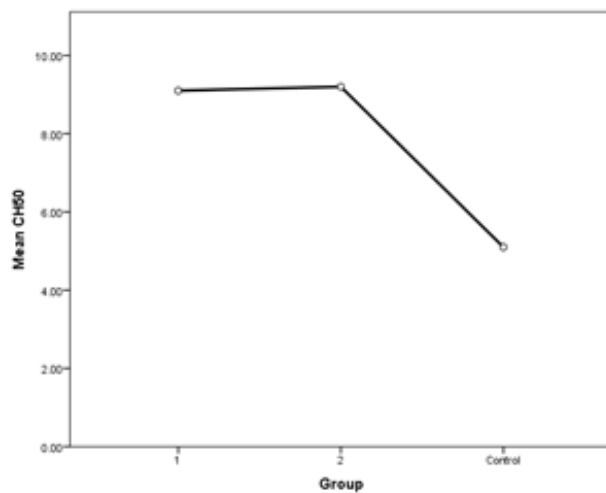
In each column numbers that have similar letters the difference is not significant ( $p < 0.05$ ).



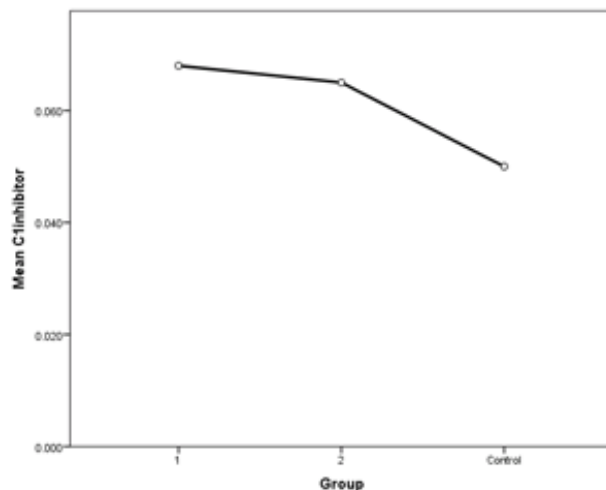
**Figure 2.** The effect of *Satureja bachtiarica* alcoholic extract level of C<sub>4</sub> of rat blood serum



**Figure 5.** The effect of *Satureja bachtiarica* alcoholic extract level of IgM of rat blood serum



**Figure 3.** The effect of *Satureja bachtiarica* alcoholic extract level of CH50 of rat blood serum



**Figure 4.** The effect of *Satureja bachtiarica* alcoholic extract level of C<sub>1</sub> inhibitor of rat blood serum

## 4. Discussion

This study was conducted to investigate the effect of *Satureja bachtiarica* alcoholic extract on some components of complement system and IgM in rat serum. In the study by Sturkie (1995), conducted to test the effect of medicinal plants on immunity surface of hen's body, using *urtica dioica* as combined with *Satureja* increases lymphocytes percent and decreases heterophils percent that indicates positive effect of this plant on enhancement of body immune system. Lymphocytes have role in cell-mediated immunity. Cell-mediated immunity involves increase in delayed hypersensitivity, Graft-versus-host, defense against intracellular organisms (like *Bacillus* and *Brucella*) and possibly defense against neoplasms and also plays role in humoral immunity with production of antibodies (Sturkie, 1995). With regard to spherocytosis property of complement system and delivering antigen to lymphocytes and phagocytosing cells (Male et al., 2006) increase in amount of complement factors can increase antigens spherocytes and increase in phagocytosis and also better identification of antigens delivered by adaptive immunity and antibody production against them.

In present study *Satureja* extract created a relative increase in all measured components of complement system and IgM compared to control group. Regarding to the stated contents this relative increase can create increase in lymphatic immune system performance and phagocytes and overall cascade of immune system performance.

To the obtained results by Iranpour Mobarakeh et al. (2014), Savory (*Satureja Hortensis* L.) plant can lead to increase blood serum proteins (Iranpour Mobarakeh et al., 2014). In another study, it was demonstrated that *Satureja khuzistanica* essential oil has significant anti-microbial activity. In addition, a cytotoxic effect of the oil against cancerous cell lines was noted (Yousefzadi et al., 2013).

Thomke and Elwinger (1998) also presented a report about different properties of *Satureja* plant including liver performance increase (Thomke and Elwinger, 1998). With regard to this report and the fact that complement system factors are created in liver it can be concluded that increase in liver performance can create increase in amount of complement system protein that in present study we observed a relative increase in complement system components.

Antioxidant property has been reported as another property of this plant (Radonic and Milos, 2003). Radical oxygen and nitrogen are continually produced in human body and are controlled by inner enzymes (superoxide decmutaz, catalysis, Glutathione peroxidase). When these radicals are produced excessively substrates are exposed to oxidation, body defense mechanisms become disable and biomolecules (DNA, lipid, protein) are destroyed (Oke et al., 2009).

## 5. Conclusions

With regard to antioxidant property of *Satureja*, it can be said that this plant increases immunity activity in the body.

## REFERENCES

- [1] Abdollahi M, Salehnia A, Mortazavi S, Ebrahimi M, Shafiee A, Fouladian F, Keshavarz K, Sorouri S, Khorasani R and Kazemi A. 2003. Antioxidant, anti diabetic, anti hyperlipidemic, reproduction stimulatory properties and safety of essential oil of *saturejakhuzistanica* in rat invivo: A oxicopharmacological study. *Med scimonit.*, 9 (9): 331-335.
- [2] British Pharmacopoeia. 1988. British pharmacopoeia, vol. 2. HMSO, London, pp. 137-138.
- [3] Duke JA. 1981. Handbook of Legumes of Economic Importance. Plenum Press. New York. 345 pp.
- [4] Duke JA and Wain KK. 1981. Medicinal plants of the World. Computer index with more than 85,000 entries. 3: 1654.
- [5] Hartwell JL. 1971. Plants used against cancer. A survey. *Lloydia.*, 34(4): 386-425.
- [6] Iranpour Mobarakeh H, Soleyman Dehkordi H, Jafarian Dehkordi M and Khamesipour F. 2014. Assessing the Effect of the Savory (*Satureja Hortensis L.*) Essence on Some Biochemical Factors in Rat's Blood Serum. *Advances in Life Sciences.*, 4(2): 73-78. DOI: 10.5923/j.als.20140402.06.
- [7] Lawrence BM. 1978. Essential oils Allured. Publishing, wheaton, I 11. 1979. 192.
- [8] Lawrence BM. 1981. Progress in essential oils savory oil. *Perfumer & Flavourist.*, 6 (4): 73-78.
- [9] Lewis WH and Elvin-Lewis MPF. 1977. Medical Botany: Plants Affecting Man's Health. A Wiley-Interscience Publication. John Wiley & Sons Inc., 605 Third Ave., New York, NY 10158. 515 pages.
- [10] Nazari A, Delfan B, Shirkhani Y and Kiyani AA. 2006. Effect of *Satureja Khuzestanica* on blood coagulation activity in rats he *Journal of Qazvin University of Medical Sciences.*, 9 (4): 15- 18 (Persian).
- [11] Male D, Brostoff J, Roth D, et al. Immunology, 7 th Ed. Boston,USA: C V Mosby; 2006. Janeway C, Travers P. Immunology – the Immune System in Health and Disease, 5 th Ed. New York, USA: Garland Publishing 2006.
- [12] Oke F, Aslim B, Ozturk S and Itundag S. 2009. Essential oil composition, antimicrobial, and antioxidant activities of *Satureja cuneifolia* Ten. *Food Chem.*, 112: 874-879.
- [13] Radonic A and Milos M. 2003. Chemical composition and in vitro evaluation of antioxidant effect of free volatile compounds from *Satureja montana* L. *Free Radic Res.*, 37(6): 673-679.
- [14] Rechinger KH. 1982. *Teucrium L.* In: Rechinger KH ed. *Flora Iranica*. Graz: Akademische Druck und Verlagsanstalt.
- [15] Sharafzadeh S. 2014. Comparison of the Main Essential Oil Components of Different Species of *Satureja* from Iran: A Review. *App. Sci. Report.*, 2 (1): 1-3.
- [16] Soodi M, Moradi S, Sharifzadeh M and Saeidnia S. 2012. *Satureja bachtiarica* methanolic extract ameliorate beta amyloid induced memory impairment. *Research in Pharmaceutical Sciences.*, 7(5): S802- S85.
- [17] Sturkie P.D. 1995. Avian physiology. 4th ed .Springer Verlag , New York , 486 P .
- [18] Sumiko M, Cisinota FM, Cuilerm ORN, Nestor MP and Vicente DS. 2001. Testosterone effect on insulin content, messenger ribonucleic acid lvel. Promoter activityand secretion in rat. *Endocrinology.*, 47:144- 170.
- [19] Thomke S and Elwinger K. 1998. Growth promotants in feeding pigs and poultry. III. Alternatives to antibiotic growth promotants. *Annales de Zootechnie.*, 47, 245-271.
- [20] Yousefzadi M, Riahi-Madvar A, Hadian J, Rezaee F, Rafiee R and Biniiaz M. Toxicity of essential oil of *Satureja khuzistanica*: In vitro cytotoxicity and anti-microbial activity. *J Immunotoxicol.*, Early Online: 1-6.
- [21] Zareii B, Seyfi T, Movahedi R, Cheraghi J and Ebrahimi S. 2014. Antibacterial effects of plant extracts of *Alcea Digitata L.*, *Satureja Bachtiarica L.* and *Ferulago Angulata L.* *J Babol Univ Med Sci.*, 16(1):31-37.