

Evaluation of the Anticestode and Antinematode Effects of the Methanol Extract of *Ferula Asafoetida* on Experimentally Infected Rats

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ABSTRACT

BACKGROUND AND OBJECTIVE: Synthetic antiparasitic medications are often associated with drug resistance and adverse side effects. In traditional medicine, *Ferula asafoetida* has been widely used in the treatment of parasitic infections, and various studies have confirmed the anti-leishmanial, anti-Giardia, and antifungal properties of this medicinal herb. This study aimed to evaluate the antiparasitic effects of the methanol extract of asafoetida.

METHODS: This experimental study was conducted on 100 male rats equally divided into 10 groups. To induce infection, animals were fed the eggs of nematode *Syphacia obvelata* and cestode *Hymenolepis nana* via gavage. Animals in groups one and two were considered as control subjects for cestode and nematode, respectively and received no medications. The third group was administered with a standard anticestodal dose of praziquantel (25 mg/kg), and the fourth group was administered with a standard antinematodal dose of piperazine (20 mg/kg). Infected animals in experimental groups five, six and seven received treatment with 2.5%, 5% and 10% concentrations of asafoetida methanol extract, respectively. Moreover, induction of nematode infection was performed on the animals of experimental groups 8, 9 and 10, which were administered with 2.5%, 5% and 10% concentrations of asafoetida methanol extract, respectively. Treatment of the animals continued for two weeks. Number of parasite eggs in the fecal samples was determined at the end of the first and second week of treatment. After the intervention, total number of the intestinal parasites was calculated and assessed in the experimental groups.

FINDINGS: In the first week of treatment, no statistically significant difference was observed between the number of *Syphacia obvelata* eggs in rats administered with the methanol extract of asafoetida even at the highest concentration compared to control subjects (166.4±3.11 vs. 235±9.5). Similarly, treatment of nematode infection with the methanol extract of asafoetida could not decrease the number of eggs and parasites by the end of the second week of treatment (P>0.05). However, rats with cestode infection receiving different concentrations of asafoetida extract showed a significant reduction in the number of eggs and parasites compared to control subjects (p<0.05).

CONCLUSION: According to the results of this study, the methanol extract of *Ferula asafoetida* had no antinematode properties in vitro, while it exhibited anticestode effects on laboratory animals with induced parasitic infection.

KEY WORDS: *Ferula asafoetida*, *Syphacia obvelata*, Piperazine, *Hymenolepis nana*, Praziquantel, Rats.

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Introduction

Parasitic infections are associated with numerous health problems and may develop epidemically in a community. Although various pharmacological agents are effective in the treatment of these infections, some of these medicines often cause drug resistance, drug residues in tissues and other side effects. Throughout history, different medicinal herbs have been used in the treatment of parasitic infections across the world (1). Frequent references in relation to the antiparasitic effects of Iranian herbs in the sources of traditional medicine have urged researchers toward inclusive investigation in this regard in order to validate the data. Recently, studies have focused on the application of medicinal herbs owing to the approval of patients, remarkable efficiency of herbal remedies and low treatment costs (2). *Ferula asafoetida* is one of the therapeutic herbs commonly recommended in the Iranian traditional medicine for the treatment of parasitic diseases (3). This plant is grown in Iran and Afghanistan and belongs to the Apiaceae family (3).

Asafoetida has a strong odor due to the presence of sulfur compounds, such as disulfides, trisulfides and tetrasulfides. Moreover, this plant contains coumarin derivatives (4). In the traditional medicine, *asafoetida* is used as a mucokinetic, diuretic, sedative and antiparasitic agent. In traditional remedies, the most commonly administered dose of *asafoetida* was 0.2-0.5 grams daily (5, 6). In veterinary medicine, *asafoetida* was an essential ingredient in preparing antiparasitic baths (3). According to the literature, the aqueous extract of *asafoetida* is administered orally in Nepal as a remedy for intestinal worms (7). Previous studies have suggested that the resin extracted from the stems and roots of *asafoetida* contributes to the growth inhibition of *Trichomonas vaginalis* protozoan in vitro (8). Furthermore, researchers have claimed that oral administration of *asafoetida* essential oil in rats with *Schistosoma mansoni* infection could decrease the number of parasites in the treated samples (9).

With this background in mind, this study aimed to evaluate the anticestode and antinematode effects of the methanol extract of *asafoetida* on rats experimentally infected with nematode *Syphacia obvelata* and cestode *Hymenolepis nana* in vitro.

Methods

Preparation of *asafoetida* herbal extract: In this study, *asafoetida* was collected from the plains of

Bojnurd city located in Khorasan province, Iran during the summer of 2014. Type and species of the plant were confirmed by Sari University of Agricultural Sciences & Natural Resources (Mazandaran province, Iran). Aerial parts of *asafoetida* were dried at room temperature and powdered afterwards. To provide the herbal extract, we used the percolation method and methanol. Using a distillation apparatus, the obtained methanol extract was condensed in vacuum and kept at the temperature of 4°C until the beginning of experiments.

Experimental animals: In total, 100 male rats (age: 60 days, weight: 25-30 grams) were purchased from Pasteur Institute of Iran (Amol Branch). The animals had been preserved in sterile conditions. Samples were transferred to the laboratory of the Faculty of Veterinary Medicine at Islamic Azad University of Babol Branch, Iran.

Induction of infection in rats: To induce infection in the experimental animals, eggs of nematode *Syphacia obvelata* and cestode *Hymenolepis nana* were separated from the fecal samples of infected rats in Pasteur Institute of Iran (Amol Branch). Samples were studied on wet mount slides using an optical microscope, and viability of the eggs was confirmed. To infect the healthy rats (as confirmed by fecal examination), they were each fed 200 cestode and nematode eggs (10).

Treatment groups and applied doses: Two weeks after the induction and confirmation of parasitic infection, the animals were randomly divided into 10 groups of 10. The first group was considered as cestode control, and the second group was determined as nematode control. None of the animals in these groups received treatment. Animals in the third study group received praziquantel at the standard anticestodal dose of 25 mg/kg, and rats in group four were administered with piperazine at the standard antinematodal dose of 20 mg/kg. Infected animals in experimental groups five, six and seven received 2.5%, 5% and 10% concentrations of the methanol extract of *asafoetida*, respectively. Similarly, nematode-infected rats in experimental groups 8, 9 and 10 received treatment with 2.5%, 5% and 10% concentrations of the methanol extract of *asafoetida*. All the mentioned dosages were administered via oral gavage (fig 1).

After the first week of treatment, animals in the experimental groups were evaluated in terms of the number of parasite eggs using the Willis method. At the end of the second week of treatment, the number of

intestinal parasites in all experimental groups was determined upon necropsy (10).

Statistical analysis: Data analysis was performed in SPSS V.13 using one-way analysis of variance (ANOVA) and the Bonferroni method, and $p < 0.05$ was considered statistically significant.



Figure 1. Drug administration in rats via gavage

Results

After one week of treatment, a significant reduction was observed in the number of *Syphacia obvelata* eggs in animals administered with piperazine compared to the control subjects and other experimental groups ($p < 0.01$). Moreover, animals receiving treatment with piperazine showed a significant difference in this regard compared to the other study groups ($p < 0.05$).

On the other hand, treatment with the methanol extract of *asafoetida* was associated with no significant reduction in the number of parasite eggs in the fecal samples of the rats, even at the highest concentration ($p < 0.05$). Nevertheless, after the first week of treatment, a statistically significant difference was observed in the number of *Hymenolepis nana* parasite eggs in the animals administered with praziquantel and 2.5%, 5% and 10% concentrations of the methanol extract of *asafoetida* compared to the control group

($p < 0.05$) (table 1). After two weeks of treatment, a significant difference was observed in the number of *Hymenolepis nana* eggs in the fecal samples of all the experimental groups compared to control subjects ($p < 0.05$). According to the results of this study, treatment of parasitic infections with the methanol extract of *asafoetida* at different concentrations significantly decreased the number of cestode eggs in the fecal samples of animals after two weeks. However, the anticestode effects of *asafoetida* methanol extract was found to be dose-dependent, so that treatment at higher doses caused a greater response in the animals. As such, there was a significant difference in the improvement of parasitic infection between the animals receiving 2.5% and 10% concentrations of the *asafoetida* extract ($p < 0.05$).

As expected, piperazine treatment eliminated all the nematodes, while praziquantel destroyed all the cestodes. Consequently, the results of necropsy revealed no intestinal worms in the infected rats treated with the mentioned drugs. No statistically significant difference was observed between the animals in the *Syphacia obvelata* group treated with *asafoetida* extract and control subjects ($p > 0.05$). In addition, treatment with the methanol extract of *asafoetida* had no significant effect on the number of intestinal nematodes even at high concentrations. On the other hand, *asafoetida* extract had a significant effect on the number of *Hymenolepis* cestodes ($p < 0.05$).

It is noteworthy that all the concentrations of the herbal extract significantly reduced the number of intestinal cestodes in the samples obtained from the animals of the experimental groups compared to control subjects ($p < 0.05$) (table 2).

Table 1. Mean number of cestode and nematode parasitic eggs in fecal samples of study groups during treatment

Groups	<i>Syphacia obvelata</i>		<i>Hymenolepis nana</i>	
	Mean±SD		Mean±SD	
	Week One	Week Two	Week One	Week Two
Control	235±9.5 ^a	214.2±12.4 ^a	111.8±10.3 ^a	124.6±8.4 ^a
Piperazine	14.6±11.4 ^b	0 ^b		
Praziquantel			19±1.5 ^b	-
10% <i>Asafoetida</i> Methanol Extract	166.4±3.11 ^a	121.2±11.5 ^a	21±1.5 ^b	5.1±0.6 ^b
5% <i>Asafoetida</i> Methanol Extract	176.3±3.13 ^a	147.9±20.7 ^a	33.6±5.7 ^b	7±2.14 ^b
2.5% <i>Asafoetida</i> Methanol Extract	210.9±9.5 ^a	195.7±15.3 ^a	59.2±5.3 ^b	43.5±4.33 ^b

Table 2. Mean of intestinal parasites after necropsy of studied rats

Groups	<i>Syphacia obvelata</i>	<i>Hymenolepis nana</i>
	Mean±SD	Mean±SD
Control	221.3±40.29 ^a	98.5±10.6 ^a
Piperazine	0 ^b	-
Praziquantel	--- ^b	0 ^b
10% <i>Asafoetida</i> Methanol Extract	177.4±13.81 ^a	1.4±0.88 ^b
5% <i>Asafoetida</i> Methanol Extract	209.7±22.38 ^a	8.6±1.02 ^c
2.5% <i>Asafoetida</i> Methanol Extract	214.6±34.57 ^a	15.7±0.8 ^c

Dissimilar letters in each column indicate a significant difference based on Bonferroni correction method ($p < 0.05$)

Discussion

According to the results of the present study, treatment with the methanol extract of asafetida could effectively reduce *Hymenolepis nana* parasites in experimentally infected rats. In another study, Maraghi et al. performed the laboratory and clinical evaluation of the antiparasitic effects of asafetida against *Hymenolepis nana* in comparison with niclosamide. According to the findings, administration of 17.7 mg/ml of asafetida extract for two weeks improved the health of 90% of the infected rats (11).

This was in congruence with the results of the current study, which marked a significant reduction in the number of mature parasites in the experimental animals receiving treatment with the methanol extract of asafetida compared to control subjects. This finding confirms the anticestode properties of asafetida plant, which has been reported by Nazemi et al. as well (12). In a study in this regard, Barati assessed the anti-*Leishmania* effects of asafetida herbal extract against *Leishmania major* promastigotes, and the obtained results were indicative of the anti-leishmanial effects of this medicinal herb (13). In this regard, findings of Ramdan et al. revealed that oral administration of asafetida essential oil could significantly decrease the number of eggs and worms in rats infected with *Schistosoma mansoni* (9). Another study by the same researchers was performed to evaluate the antiparasitic effects of asafetida against *Trichomonas vaginalis* in vitro. According to the findings, the resin extracted from the stems and roots of asafetida could significantly inhibit the growth of *Trichomonas vaginalis* in vitro (8).

According to the results obtained by Sarkari et al., treatment with the herbal extract of asafetida at the concentration of 2 mg/ml for one hour after exposure to *Trichomonas vaginalis* reduced the number of parasites by 90% (14). Furthermore, Rezaeimanesh et al. reported that the aqueous and alcoholic extracts of asafetida showed lethal effects against *Giardia* cysts; however, this effect was found to be more significant with the alcoholic extract (15). *Artemisia absinthium* is another medicinal herb grown in Iran widely used in

the traditional medicine for the treatment of parasitic infections. In a study by Youssefi et al. conducted to evaluate the antiparasitic effects of *Artemisia absinthium*, it was reported that the alcoholic extract of this herb demonstrated therapeutic effects against experimental infection with *Syphacia obvelata*. Nevertheless, the herbal extract of asafetida was observed to have no significant effect against nematode *Syphacia obvelata* in the present study, which rules out the antinematode potential of this medicinal plant (16). Medicinal herbs are healthy, abundant and sustainable sources of nature, which have been prominently considered in the traditional medicine. Due to the increasing resistance to available antiparasitic medications and subsequent side effects of these agents, discovering new and low-risk therapeutic sources in this regard is of paramount importance (8).

One of the most beneficial medicinal herbs in the treatment of parasitic diseases is *Ferula asafetida*, which is grown in Iran and has long been used in the Iranian traditional medicine. In conclusion, results of the present research indicated that the methanol extract of asafetida had no antinematode effect on the studied samples, while the anticestode effects of this medicinal plant were confirmed.

Therefore, it is recommended that different compounds of asafetida with possible anticestode properties be identified and extracted in future studies. Furthermore, complementary research in this regard could deliver this medicinal herb or its compounds as useful anticestode medications. Adequate knowledge of medicinal herbs and their associated therapeutic effects could promote public health in society, prevent the complications caused by chemical medications and reduce healthcare costs.

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References

1. Behnke JM, Buttle DJ, Stepek G, Lowe A, Duce IR. Developing novel anthelmintics from plant cysteine proteinases. *Paras vec.* 2008;1(1):1-18.
2. Khajeh M, Yamini Y, Bahramifar N, Sefidkon F, Pirmoradei MR. Comparison of essential oils compositions of *Ferula assa-foetida* obtained by supercritical carbon dioxide extraction and hydrodistillation methods. *Food Chem.* 2005;91(4):639-44.
3. Zargari A. Medicinal plants. Tehran: Tehran University Press; 1996.
4. Kojima K, Isaka K, Ondognii P, Zevgeegiin O, Gombosurengyin P, Davgiin K, et al. Sesquiterpenoid Derivatives from *Ferula ferulioides*. IV. *Chem Pharm Bull.* 2000;48(3):353-6.
5. Mohammadi R, Sepahvand A, Mohammadi SR, Mirsafaei H, Shargh RN. Antifungal activity of *Ferula assa-foetida* against clinical agents of Mucormycosis. *Journal of Isfahan Medical School.* 2009;27(100):582-8.
6. Sadraei H, Ghannadi A, Malekshahi K. Composition of the essential oil of *assa-foetida* and its spasmolytic action. *Saudi Pharma J.* 2003;11(3):136-40.
7. Bhattarai N. Folk anthelmintic drugs of central Nepal. *Int J Pharmacol.* 1992;30(2):145-50.
8. Ramadan NI, Al Khadrawy F. The in vitro effect of *assafoetida* on *trichomonas vaginalis*. *J Egypt Soc Parasitol.* 2003;33(2):615-30.
9. Ramadan N, Abdel-Aaty H, Abdel-Hameed D, El Deeb H, Samir N, Mansy S, et al. Effect of *ferula assafoetida* on experimental murine *Schistosoma mansoni* infection. *J Egypt Soc Parasitol.* 2004;34(3):1077-94.
10. Shady OMA, Basyoni MM, Mahdy OA, Bocktor NZ. The effect of praziquantel and *Carica papaya* seeds on *Hymenolepis nana* infection in mice using scanning electron microscope. *Parasitol Res.* 2014;113(8):2827-36.
11. Maraghi S, Soghra T. In vitro and in vivo assay of *Ferula assa* extract effects on *Hymenolepis nana* and comparison of it with Niclosamide. *Jundisapour J Med Sci.* 1991;23:48-56. [In Persian]
12. Nazemi RJ, Moharamipour S. Repellency of *Nerium oleander* L. *Lavandula officinalis* L. and *Ferula assafoetida* L. extracts on *Tribolium castaneum* (Herbst). *IRAN J Med Arom Plant.* 2008;23(4):443-52. [In Persian]
13. Barati M, Sharifi A, Sharifi Far S. Antileishmanial activity of *Artemisia aucheri*, *Ferula asafoetid* and *Gossypium hirsutum* extracts on *Leishmania major* promastigotes in vitro. *J Artesh Med Univ.* 2010;8(3):166-72. [In Persian]
14. Sarkari B, Hadisa T, Shahrbanoo A, Elahm F, Mehrangiz A. In Vitro anti-*Trichomonas* activity of *Ferula assafoetida* and garlic extracts. *J Gorgan Uni Med Sci.* 2009;11(3):13-7. [In Persian]
15. Rezaeiemanesh M, Shirbazou S. In-vitro giardicidal effect of aqueous and alcoholic extracts of *Asafoetida* on *Giardia lamblia* cyst. *J Birjand Univ Med Sci.* 2012;19(1):22-3. [In Persian]
16. Youssefi MR, Abuhosseini Tabari M, Sadeghi Hashjin G, Kouhi MK. Antiparasitic efficacy of worm wood (*Artemisia absinthium*) alcoholic extract on *Syphacia obvolata*. *Iran J of Veter Med.* 2012;6(1):47-50. [In Persian]