Evaluation of Toxicity Effects of Asafetida on Biochemical, Hematological, and Histological Parameters in Male Wistar Rats

Seyyed Majid Bagheri, Maryam Yadegari, Aghdas Mirjalily, Mohammad Ebrahim Rezvani

Department of Physiology, Herbal Medicine Research Center, Shahid Sadoghi University of Medical Sciences, 1Department of Anatomy, Shahid Sadoghi University of Medical Sciences, 2Neurobiomedical Research Center, Shahid Sadoghi University of Medical Sciences, Yazd, Iran

ABSTRACT

Objectives: Asafetida is traditionally used in folklore medicine for the treatment of various ailments. To validate its use in traditional medicine, it is important to evaluate its toxicity in the animal system. Therefore, this study aimed to evaluate the toxicological effects of asafetida in Wistar albino rats. Materials and Methods: Acute toxicity tests were conducted by the oral administration of 250, 500, and 1,000 mg/kg body weight of the animal. In chronic study, animals were administered with various doses of asafetida (25, 50, 100, and 200 mg/kg body weight) for a period of 6 weeks. At end of experiment, the effects of asafetida on hematological, renal, and hepatic markers and histological parameters were analyzed. Results: In acute toxicity study, no mortality was seen up to 72 h of the administration of asafetida. No signs of neurological and behavioral changes were noticed within 24 h. In the chronic study, the asafetida intake has changed the hematological parameters such as red blood cell (RBC), white blood cell (WBC), hematocrit (HCT), and platelets. Aspartate aminotransferase (AST) and lactate dehydrogenase (LDH) were significantly increased in treated animals. The plasma level of urea and creatinine were not altered by the administration of asafetida throughout the study. Histopathology study indicates hepatotoxicity, but no signs of prominent pathological changes in kidney. Conclusions: Asafetida did not show any acute toxicity, but chronic administration could have undesirable effects on hepatocytes and hematological factors.

Key words: Asafetida, hematology, kidney, liver, toxicity

INTRODUCTION

Asafetida, an oleo-gum-resin, is extracted by incisions in the stem and roots of Ferula assafoetida L. and some others Ferula species.[1] The resin is an important pharmacological and industrial agent and is being used in traditional medicines for treatment of various disorders like stomach ache, indigestion, bronchitis, asthma, and whooping cough;[2] and for treatment of intestinal parasites.[3] Nepalian people use asafetida 50–200 mg twice a week mainly as sedative, carminative, antispasmodic, diuretic, antihelmintic, emmenagogue, and aphrodisiac agent.[4] In Ayurveda, asafetida is introduced as a valuable remedy for flatulence, hysteria, gastrointestinal disorders, inflammation, stomach ache, spasmodic, and helminthic.[1] New studies
showed that this oleo-gum-resin has antiviral, antifungal, cancer chemopreventive, anti-diabetic, and cytotoxicity effect.\cite{1} It also has antispasmodic, anticonvulsant, and antinociceptive effects.\cite{5,6,7} Although there are different studies about pharmacological properties of asafetida, to our knowledge, there is no comprehensive toxicological study of asafetida on animal models. There is only one case report that methemoglobinemia has been observed after administration of asafetida in a 5-week-old black male infant.\cite{8} It is also recommended that asafetida should not be used during pregnancy as it may increase the risk of abortion.\cite{9}

The toxicity effects of asafetida have been studied mainly on parasites and some protozoan animals suggesting it antiparasitic,\cite{10,11,12} antifungal,\cite{13} and antibacterial.\cite{14} Kumar and Singh reported that different root extracts of Ferula assafoetida have anti molluscicidal activity against the snail Lymnaea acuminata.\cite{10} Bagheri et al., showed that asafetida has cytotoxicity effect on the brine shrimp.\cite{15} Some of old studies showed that asafetida has a weak exchange-inducing on sister chromatid in spermatogonia and clastogenicity in mouse spermatocytes.\cite{16,17} The study is aimed to study the toxic effects of asafetida on blood parameters, kidney and liver factors, and histopathology of kidney and liver in male Wistar rats.

**MATERIALS AND METHODS**

**Animals**

Male Wistar rats weighing between 150 and 180g and 6–8 weeks old were bred and maintained in the animal house unit of the faculty of medicine under controlled temperature at 21 ± 1°C in 12 h light: 12 h darkness schedule. Animals were housed in plastic cages and food and water was made available *ad libitum*. The study was approved by Institutional Animal Ethical Committee of Shahid Sadoughi University of Medical Science.

**Acute toxicity studies**

For acute toxicity studies, rats were divided into four groups (n = 5; each) and were administered orally with three doses of asafetida corresponding to 250, 500, and 1,000 mg/kg body weight. The control group received 1 ml of physiological saline. They were all placed under observation immediately for 24 h for any behavioral, neurological changes and then mortality for the next 72 h.\cite{18}

**Chronic toxicity studies**

The rats were divided into five groups (n = 4; each) and were administrated four different doses of asafetida corresponding to 25, 50, 100, and 200 mg/kg orally every day for 6 weeks’ duration. Another group of five rats that received 1 ml of physiological saline served as control.

**Plant oleo-gum resin**

*Ferula assafoetida* oleo-gum-resin was collected from Tabas region (Yazd province, Iran) during the summer and the plant species was botanically identified by Dr. Abbas Zarezadeh in Yazd Agricultural Research Center. The dried powder of asafetida was soaked in distilled water overnight at room temperature and the yielded suspension was used orally. Concentrations and dosages of the extract were expressed as crude amount of the dried oleo-gum-resin used in preparing the stock solution.

**Biochemical analysis**

Blood was collected from orbital sinus of rats. Serum was prepared by centrifugation (3,000 rpm, 20 min) and stored frozen until biochemical assay. The urea, creatinine, lactate dehydrogenase (LDH), and aspartate aminotransferase (AST) were determined using the suitable kits and according to the manufacturer’s instructions.

**Hematological tests**

The blood with ethylenediaminetetraacetic acid (EDTA) was used for the counting of RBC and total count of WBC and platelet by standard procedures. Hematocrit percentage (HCT %) was also determined.\cite{19}

**Histopathology evaluation**

For histopathological studies, liver and kidney tissue were collected immediately in 10% neutral buffered formalin. The tissues were processed by routine histological methods and embedded in paraffin blocks. The sections were cut by a rotary microtome and subsequently stained with Ehrlich’s hematoxylin and eosin (H and E). The histological sections were studied under Olympus light microscopy (Olympus, Japan, magnification ×40) to evaluate histopathological changes.

**Statistical analysis**

Statistical data were assessed with one-way analysis of variance (ANOVA), followed by *post hoc* Tukey’s test using Graph pad prism version 5. Results were expressed as mean ± standard error of the mean (SEM). A value of *P* < 0.05 was considered significant.

**RESULTS**

**Acute toxicity studies**

After the administration of different doses of asafetida, the rats were monitored for the signs of toxicity. The animals
showed normal activity and no mortality were observed till the end of the study.

**Biochemical factors**

The result of serum kidney and liver parameters analysis in chronic study was shown in Table 1. There is no significant difference between the urea and creatinine concentration in control and treated animals with asafoetida. Our results also showed that AST and LDH significantly increased in all treated animals with asafoetida compared to control group ($P < 0.05$).

**Hematological changes**

Table 2 reveals that the asafoetida significantly decreased the concentrations of WBC, RBC, platelets, and HCT % after 6 weeks of treatment with asafoetida. The HCT % in asafoetida 25 group did not show any significant difference.

**Histopathological findings of liver**

The liver sections of rats in control group showed normal hepatic architecture and normal hepatocytes with prominent nucleus, central vein, and portal areas with no sign of inflammation or necrosis [Figure 1a]. Histopathological examination of extract treated rats showed little degenerative changes in some hepatic cells [Figure 1b]. Findings demonstrated that with increase in the extract doses (50, 100, and 200 mg/kg respectively), hepatocytes became larger in size with prominent nucleus compared to control group. Hypotrophied Kupffer cells were also prominent with increase in extract doses and blood vessels expansion and dilated hepatic sinusoids were noticed [Figure 1c-e]. However, the general structures of lobules of liver are normal and clear.

**Histopathological findings of kidney**

Histological examination of the control group showed normal appearance and there were no changes in the renal tubular and glomerulus of kidneys [Figure 2a]. Sections of kidneys from extract groups (25, 50, 100, and 200 mg/kg) showed no signs of prominent pathological changes except occasional renal tubular necrosis [Figure 2b-d]. In some sections of extract groups, mild infiltration of inflammatory cells around the blood vessels and in the interstitial spaces was found. In extract group (200 mg/kg), some of the glomeruli appeared to be slightly expanded and there were little signs of tubular degenerative changes [Figure 2e].

**DISCUSSION**

The herbal drugs are playing an important role in healthcare programs throughout the world with a general belief that herbal drugs are always safe as these are natural. However,
adverse effects do occur which are usually mild and affect a small number of people. Asafetida is used in different countries not only as a culinary spice but also traditionally used to treat various diseases. Although there are no experimental data about toxicity effects of asafetida, in Iranian traditional, it was emphasized that intake of larger dosages of asafetida can lead to swelling of the lips, digestive complaints such as flatulence and diarrhea, discomfort, and headache. {3} The nontoxic nature of the asafetida and its oral safety at selected dose level was confirmed as the administration of the drug does not cause mortality in rats. According to Clarke and Clarke, {20} any compound or drug with the oral LD50 estimate greater than 1,000 mg/kg could be considered of low toxicity. This supports that asafetida was found to be safe up to 1,000 mg/kg body weight in terms of mortality. In the present study, acute toxicity of asafetida caused a significant increase in AST and LDH level in treated animals as compared to control group [Table 1]. There are many enzymes found in the serum that did not originally originate from the serum. During tissue damage, some of these enzymes find their way into the serum, probably by leakage. {21} Serum enzyme measurements are therefore a valuable tool in clinical diagnosis, providing information on the effect and nature of pathological damage to any tissue. The increase in serum LDH and AST activity may indicate hepatic damage probably by the altered cell membrane permeability leading to the leakage of the enzymes from the tissues to the serum. {22,23} LDH and AST are considered to be sensitive indicators of hepatocellular damage and within limit can provide a quantitative evaluation of the degree of damage to the liver. {24} In the present study, there was not any significant difference in urea and creatinine concentrations following treatment with asafetida. Destruction of glomeruli causes significant decrease in the glomerular filtration rate (GFR) and increases the blood urea and creatinine resulting chronic renal failure. {25} Since the urea and creatinine are markers of kidney function, {26} the data suggest that asafetida is not nephrotoxic. Hematological data obtained in our work indicated a significant decline in RBC and WBC and platelet count [Table 2]. Altered RBC, WBC, and platelet counts reflect that asafetida would have some mild effect on the hematological parameter. Since blood cells are originated from bone marrow, the inertness of the extract on this organ is evident and its use in pharmacological evaluation is again certified. Histopathological observations revealed that treatment with asafetida resulted little degenerative changes of some hepatic cells. Sections of kidneys from extract groups (25, 50, 100, and 200 mg/kg) showed no signs of prominent pathological changes in the cortex and the medulla, but little renal tubular necrosis was seen. Our finding also demonstrated that these changes were dose-dependent, respectively, and these histopathological evidences were in agreement with biochemical results.

CONCLUSION

The data showed that asafetida did not have any acute toxicity, but the chronic consumption of this oleo-gum-resin caused reverse effects on liver and blood parameters. This study suggested that asafetida may be used in minimum possible doses.

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Conflicts of interest

There are no conflicts of interest.

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