Radioprotective Effect of Arbutin in Megavoltage Therapeutic X-irradiated Mice using Liver Enzymes Assessment

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ABSTRACT
Introduction: Medical use of ionizing radiation has direct/indirect undesirable effects on normal tissues. In this study, the radioprotective effect of arbutin in megavoltage therapeutic x-irradiated mice was investigated using serum alkaline phosphatase (ALP), alanine aminotransferase (ALT), and aspartate aminotransferase (AST) activity measurements.

Material and Methods: In this analytical and experimental lab study, sixty mice (12 identical groups) were irradiated with 6 MV x-ray beam (2 and 4 Gy in one fraction). Arbutin concentrations were chosen 50, 100, and 200 mg/kg and injected intraperitoneal 2 hours before irradiation. Samples of peripheral blood cells were collected and serum was separated on the 1, 3, and 7 days post-x-radiation; in addition, the level of ALP, ALT, and AST were measured. Data were analyzed using one-way ANOVA, and Tukey HSD test.

Results: X-radiation (2 and 4 Gy) increased the ALT and AST activity levels on the 1, 3, and 7 days post-irradiation, but the ALP level significantly increased on the 1 and 7 days and decreased on the third day compared to the control group (P<0.001). ALP, ALT and AST activity levels in “2 and 4 Gy x irradiation + distilled water” groups were significantly higher than “2 and 4 Gy irradiation + 50, 100, and 200 mg/kg arbutin” groups on the first and seventh day post-irradiation (P<0.001).

Conclusion: Arbutin is a strong radioprotector for reducing the radiation effect on the whole-body tissues by measuring ALP, ALT and AST enzyme activity levels. Furthermore, the concentration of 50 mg/kg arbutin showed higher radioprotective effect.

Keywords
Arbutin, Liver, Enzymes, Radiation Protection
methods included modification of the radio-therapy technique and method of irradiation, like the volumetric irradiation, using beam shapers, and changing the number of radiation fields [1, 5]. Although the non-pharmacological strategy can reduce the RT damage, the probability of secondary cancers incidence has been still remarkably high [6].

Pharmaceutical strategies, including radioprotectors and radiosensitizers, as a moderator of free radical damage in the targeted radiotherapy are utilized for reducing toxicity in normal tissues [7]. Radioprotectors are divided into two groups, including natural and chemical agents.

Arbutin or hydroquinone glycoside, is a natural agent [8], that can be found in lots of plants belonging to the Ericaceae and Compositae groups [8, 9]. Peduncles, leaves, and bark of Pyrus plants contain certain values of arbutin [8]. Arbutin is a yellow powder soluble in water and alcohols, and the molecular weight is 272.25 g/mol. The maximum and minimum absorbance spectra of arbutin are 267 and 240 nm, respectively [10].

ALP is an endogenous protein and a factor that can predict the level of cancer metastasis; in addition, it is an easy test yielding quick results [11]. ALP has four types, including placental, intestinal, germ cell, and liver/bone/kidney ALP [12]. The prognostic role of ALP has been prospected in different types of cancers such as bony metastatic prostate cancer, bony metastatic breast cancer, and clear cell chondrosarcoma of the bone [11, 13, 14]. Mild to moderate elevation of the enzyme caused by free radicals from cell damage can be seen in all types of disorders [15-17]. The deviation and level of ALP have been utilized as an indicator for determining the radiation damage [3, 12, 13].

Other serum activities (aminotransferases), including ALT and AST can show liver disorders and are reliable sensitive markers of liver diseases [18]. Their levels will increase into the bloodstream when an organ (liver, heart, etc.) is diseased or damaged.

The effectiveness of a radioprotective agent in patients undergoing RT is an important challenge. In a study by Benkovic et al. [3], the effect of propolis and flavonoids as radioprotectors was investigated in gamma-irradiated mice using ALP level assessment method. In another study, Ramadan et al. [18] evaluated the radioprotective effect of silymarin in gamma-irradiated mice using ALP level assessment method. In another study, Ramadan et al. [18] evaluated the radioprotective effect of silymarin in gamma-irradiated mice using ALP level assessment method. In this study, the radioprotective effect of arbutin in megavoltage therapeutic x-irradiated mice by measuring liver enzymes, ALP, ALT, and AST activities after 1, 3, and 7 days. In addition, the biocompatibility of the arbutin with the liver enzyme activity was evaluated at three different concentrations.

Material and Methods

Animal

In this analytical and experimental lab study, sixty male NMRI mice (6 to 7 weeks old, 25±5 gr) were obtained from Iran University of Medical Science (Tehran, Iran). Mice were maintained under controlled conditions, including appropriate temperature (22 ± 1ºC), humidity, light regime (12 hr. light/12 hr. dark), and had free access to a commercial diet.

Chemical

Pure (98%) arbutin powder (Sigma, Germany) was dissolved in distilled water with concentrations of 50, 100, and 200 mg/kg and injected into mice intraperitoneal 2 hours before irradiation.

Irradiation procedure

Mice were placed in a collective cages made of Plexiglas for whole-body exposure after at least two weeks of acclimatization and conditioning. Mice were exposed by a 6-MV photon beam from an Elekta Compact accelerator
(Elekta AB, Stockholm, Sweden). The source-to-skin distance was 100 cm with a field size of 20 x 20 cm². All animals were irradiated with 2 and 4 Gy in one fraction delivered at a dose rate of 200 cGy/min.

Experimental design

Different concentrations of arbutin were injected to the mice. After two weeks, the animals were divided to twelve groups in separate Plexiglas cage, five mice for each group. The groups were selected as following:

Group 1: “Sham irradiation + distilled water (control)”, Group 2: “2 Gy x irradiation + distilled water”, Group 3: “4 Gy x irradiation + distilled water”, Group 4: “Sham irradiation + 50 mg/kg arbutin”, Group 5: “Sham irradiation + 100 mg/kg arbutin”, Group 6: “Sham irradiation + 200 mg/kg arbutin”, Group 7: “2 Gy x irradiation + 50 mg/kg arbutin”, Group 8: “2 Gy x irradiation + 100 mg/kg arbutin”, Group 9: “2 Gy x irradiation + 200 mg/kg arbutin”, Group 10: “4 Gy x irradiation + 50 mg/kg arbutin”, Group 11: “4 Gy x irradiation + 100 mg/kg arbutin”, and Group 12: “4 Gy x irradiation + 200 mg/kg arbutin”.

Blood sampling

Blood samples were collected from the mice exposed to irradiation and/or arbutin injection after 1, 3, and 7 days and serum samples were separated. The serum level of ALT (IU/L) was estimated according to Kind and King method [19], and serum ALT and AST (U/mL-1) levels were calculated based on the method proposed by Reitman and Frankel [20].

Statistical Analysis

All measurements were analyzed using SPSS 16 software (SPSS Inc., Chicago, Illinois, US). Multiple comparisons between groups were performed using one-way analysis of variance (ANOVA), and Tukey’s test as the post hoc analysis. The level of statistical significance was set at P < 0.05.

Results

According to Figures 1, 2, and 3, ALP, ALT, and AST levels in the “Sham irradiation + 50, 100, and 200 mg/kg arbutin” groups were not significantly disturbed in comparison with the control group during 1, 3, and 7 days post-irradiation (P>0.05). The ALP activity level in the

Figure 1: The effective of x-irradiation along with various concentrations of arbutin in ALP level on the 1, 3, and 7 days.
“2 and 4 Gy x irradiation + distilled water” was significantly higher on the 1 and 7 days, but it was significantly lower on the third day post-irradiation compared to the control group and all arbutin concentrations groups (P<0.001).

In addition, significantly higher ALT and AST activity levels were observed in “2 and 4 Gy x irradiation + distilled water” on the 1, 3, and 7 days post-irradiation respect to the control group and all arbutin concentrations groups (P<0.001).

ALP level in the “2 and 4 Gy x irradiation + distilled water” groups was significantly higher on the 1 and 7 days after post irradiation in comparison with the “2 and 4 Gy irradiation + 50 mg/kg arbutin”, “2 and 4 Gy irradiation + 100 mg/kg arbutin”, and “2 and 4 Gy irradiation + 200 mg/kg arbutin” groups, but it was

Figure 2: The effective of x-irradiation along with various concentrations of arbutin in ALT level on the 1, 3, and 7 days.

Figure 3: The effective of x-irradiation along with various concentrations of arbutin in AST level on the 1, 3, and 7 days.
significantly lower on the third day post-irradiation ($P < 0.001$). After 1, 3, and 7 days post-irradiation, ALT and AST levels in “2 and 4 Gy x irradiation + distilled water” groups were considerably higher than the “2 and 4 Gy irradiation + 50 mg/kg arbutin”, “2 and 4 Gy irradiation + 100 mg/kg arbutin”, and “2 and 4 Gy irradiation + 200 mg/kg arbutin” ($P < 0.001$).

Statistical analysis showed that ALP enzyme activity in the “2 Gy irradiation + 50 mg/kg arbutin” group was notably lower than the “2 Gy irradiation + 100 mg/kg arbutin” and “2 Gy irradiation + 200 mg/kg arbutin” groups on the first and seventh days post-irradiation ($P < 0.05$), and this variation was considerable on the 1, 3, and 7 days for ALT and AST levels. There were not significant differences in the “2 Gy irradiation + 100 mg/kg arbutin” compared to the “2Gy irradiation + 200 mg/kg arbutin” group for ALP, ALT, and AST activity levels ($P > 0.05$). Furthermore, no remarkable differences were found in ALP, ALT, and AST levels between the “4 Gy irradiation + 50 mg/kg arbutin” and “4 Gy irradiation + 100 mg/kg arbutin” groups ($P > 0.05$), but there was a significant difference in the “4Gy irradiation + 200 mg/kg Arbutin” compared with the “4 Gy irradiation + 50 mg/kg Arbutin” and “4 Gy irradiation + 100 mg/kg Arbutin” groups.

Figure 4 shows the results concerning ALP, ALT, and AST levels with respect to the control group in different days.

Figure 4: Effects of x-irradiation and different concentrations of arbutin at ALP (A), ALT (B), and AST (C) levels with respect to the control group in different days.
ALT, and AST enzyme levels determined in blood serum of mice compared with the control group. This figure shows clearly the relationship of the enzymes on the first, third, and seventh days compared to the control group.

Discussion

RT is an important technique for treatment in various malignancies [2]. However, exposure to ionizing radiation during RT leads to undesirable physiological and biochemical changes in humans and animals [3, 4]. The search for non-toxic selective and effective radioprotector compounds, that preferentially protect normal tissues, is one of the major challenges in radiation protection researches. In a previous study, Nadi et al. [8] showed the protective effect of arbutin in mice bones as a protector of the clastogenic and cytotoxic effects of ionizing radiation using micronuclei method.

The current study aims to detect what extent of arbutin can improve the levels of ALP, ALT, and AST on the 1, 3, and 7 days after x-irradiated mice blood serum.

In recent years, a number of serum markers, ALP, AST, and ALT, have been reported to have prognostic value in patients liver, brain, pancreas, heart, kidneys, lungs, etc. [18, 21]. The level of these enzymes will increase into the bloodstream after radiation or diseases [3, 12, 13]. Serum AST level usually rises immediately and reaches a higher level than ALT initially, after hepatocellular injury [21].

In the current study, x-irradiation (2 and 4 Gy) induced an increase in serum ALP activity after 1 and 7 days and a decrease in the third day with respect to the control value (Figure 4). Ramadan et al. [18] showed the same results with gamma-irradiation (3 Gy). Increasing of ALP activity by radiation in blood serum can be attributed to the possible release of this enzyme from various organs associated with the obstruction of the blood stream to the liver and the decreasing of it may be due to the decrease in liver Mg$^{2+}$ and Zn$^{2+}$ [18].

In this study, the level of AST and ALT increased after 1, 3, and 7 days post-exposure of 2 and 4 Gy x-rays, which is in agreement with Ramadan et al. [18]. The increase in AST and ALT activities, which may be due to the increase in the permeability of cell membranes, facilitates the passage of cytoplasmic enzymes outside the cells regarding the damage of cellular membranes during radiation [22].

After different injections of arbutin, there were not observed any remarkable changes in the level of ALP, ALT, and AST with respect to the control group. Furthermore, we found that arbutin can store ALP, ALT, and AST activity about normal range during megavoltage therapeutic x-irradiated mice. For evaluating the biocompatibility of arbutin with blood serum, measurements were made after 1, 3, and 7 days post-irradiation.

The results showed, whole-body x-irradiation (2 or 4 Gy) along with arbutin (all three concentrations) induced a significant decrease in ALT and AST levels compared to the other irradiated groups through the time intervals and it could store ALP level in the normal range on the first, third, and seventh day post-irradiation. Benkovic et al. [3] evaluated the propolis and flavonoids as a radioprotectors before gamma whole-body irradiated mice by measuring ALP activity. They indicated that these radioprotectors had protected mice from lethal effects and reduced primary DNA damage in their white blood cells. Ramadan et al. [18] investigated that the silymarin effect as a radioprotective in various doses of gamma-irradiation (3 and 6 Gy) induced hepatotoxicity. They reported that silymarin can diminish ALP, ALT, and AST due to its antioxidant effect and the ability to scavenge free radicals after irradiation.

According to our previous study [8], arbutin has a considerable protective and ameliorative effect against radiation and this protective action is related to its antioxidant properties. Therefore, it can sweep free radical, inhibit lipid peroxidation, stabilize plasma membrane, and able to restore the changes of en-
zyme activities such as ALP, ALT and AST.

Although the effects of three concentrations of arbutin were approximately similar, a minor difference was observed between them. The concentration of 50 mg/kg arbutin plus 2 Gy x radiation was the best concentration for dropping ALP, ALT and AST level. It is noticeable that in the dose of 2 Gy, the differences between 100 and 200 mg/kg of arbutin were not significant, and also 50 and 100 mg/kg for 4 Gy. Moreover, 200 mg/kg arbutin plus 4 Gy x radiation had no significant lower effect than 50 and 100 mg/kg plus 4 Gy x radiation.

As future research, it is suggested that other arbutin concentrations as a radioprotector along with x-radiation in different days should be carried out to find the precise optimized level of ALP, ALT, and AST.

**Conclusion**

In the current study, the radioprotective effect of arbutin was investigated in megavoltage therapeutic x-irradiated mice. Using ALP, ALT and AST measurements as an indicator of radiation effects, our findings demonstrated that arbutin is a strong radioprotector for reducing the radiation damage. This radioprotective effect might be correlated with the antioxidant and free radical scavenger. There were not any significant differences between the various concentrations of arbutin (50, 100, and 200 mg/kg) for storing ALP, ALT and AST levels, however, the concentration of 50 mg/kg showed higher radioprotective effects.

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**Conflict of Interest**

None

**References**


