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OPEN The effect of barley savigh consumption on blood sugar among pre-diabetic postmenopausal women

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The hormonal changes in menopause predispose women to health risks, including type 2 diabetes. Therefore, diabetes prevention in this age group is important. Considering the fewer side effects of herbal medicines compared to chemical drugs, the present study investigated the impact of barley savigh consumption on blood sugar levels in pre-diabetic postmenopausal women. This randomized clinical trial study was conducted on 64 pre-diabetic postmenopausal women referred to Community Health Centers in Gonabad city, Iran. The intervention group (n = 32) received 60 g of barley savigh daily (20 g before each meal), and the control group (n = 32) received no intervention. Fasting blood sugar (FBS) and blood sugar two hours post-prandial were checked in both groups at baseline and after two and four weeks. Data analysis was done using SPSS software (version 16) and chi-square, independent t-test, and Mann-Whitney tests at the significance level of p < 0.05. There was no significant difference in the mean FBS at baseline (109.31 ± 5.06 in the intervention and 109.03 ± 8.49 mg/dL in the control group, p = 0.904) and after two weeks after intervention (105.25 ± 5.92 in the intervention and 108.06 ± 8.75 mg/dL in the control group, p = 0.13), but the mean FBS after four weeks was significantly lower in the intervention (102.93 ± 7.75 mg/dL) compared to the control (107.90 ± 8.29 mg/dL) group (p = 0.019). The mean blood sugar two hours post-prandial was significantly lower in the intervention group compared to the control group after two (127.53 ± 8.03 and 143.68 ± 26.46 mg/dL, respectively, p = 0.006) and four weeks (127.09 ± 17.96 and 142.46 ± 27.06 mg/dL, respectively, p = 0.009). The results of the present study showed that barley savigh consumption could improve blood sugar regulation in pre-diabetic postmenopausal women.

Trial registration: The present study was conducted as a randomized clinical trial. The study protocol was approved by the Iranian Registry of Clinical Trials (IRCT code 20200929048880N1). Registration date (25/10/2020).

Keywords Diabetes, Pre-diabetes, Blood sugar, Menopause, Barley

Menopause is a physiological event in the life of women¹. Menopause is diagnosed when a woman does not have menstruation for 12 months due to the loss of ovarian follicular activity². During the transition from childbearing phase to menopause, the level of 17-beta-estradiol, which is a reproductive estrogen, drops from 100 to 250 pg/ ml to less than 10 pg/ml³. It has been shown that estrogen reduces insulin resistance, decreases insulin levels, and decreases insulin sensitivity to glucose. Therefore, the decrease in the ratio of estrogen to androgen around

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menopause can result in central accumulation of fat, abdominal obesity, insulin resistance, hyperinsulinemia, and type 2 diabetes, as well as predisposition to some cancers^{4–6}.

Diabetes is a group of metabolic diseases characterized by high blood sugar (hyperglycemia) due to either a defect in insulin secretion or insulin action⁷. Chronic hyperglycemia may damage various organs, especially the eyes, kidneys, nervous system, and cardiac blood vessels, causing loss of vision, kidney failure, leg ulcers that may result in amputation, sexual dysfunction, high blood pressure, and dyslipidemia⁷. Some of these complications could be delayed or prevented by proper interventions⁵. Diabetes is divided into several types. In type 1 diabetes, beta-pancreatic cells, which secrete insulin, are destroyed by an autoimmune reaction. This type of diabetes has an acute onset, usually before the age of 30, and requires insulin injection to control blood glucose levels. Type 2 diabetes is characterized by resistance to insulin and reduced insulin. Type 2 diabetes can be managed through exercise, diet control, and oral anti-diabetic medications8. The American Diabetes Association proposed a prediabetes state as elevated blood sugar from the normal range but below the diagnostic cut-off for diabetes. The risk of type 2 diabetes and vascular diseases is higher among pre-diabetic individuals compared to the healthy population⁹. Adhering to a healthy diet and increasing physical activity play a significant role in controlling this disease and preventing more advanced stages of diabetes in individuals at pre-diabetic stage¹⁰. Pre-diabetes is impaired glucose tolerance, fasting glucose, or mildly increased hemoglobin A1C level below the cut-off for diabetes¹¹. The prevalence of pre-diabetes among menopausal women was reported to range between 7 and 26% worldwide and between 19 and 26% in Iran based on age group and province 12-16. Annually, 10% of individuals with pre-diabetes develop diabetes¹¹.

Herbal medicines are widely prescribed worldwide due to their low side effects, low cost, availability, and effectiveness. Various plants have been traditionally used to reduce blood sugar and treat diabetes¹⁷. Barley is among the medicinal plants that have blood sugar-reducing effects¹⁸. The role of barley seeds in treating diabetes is also mentioned in the ancient texts of traditional Iranian medicine¹⁹. Barley and oat seeds contain alkaloids, beta-carotene, steroid compounds, saturated fatty acids, starch, beta-glucan fiber, B1, B2, A, C, D, and E vitamins. They are the richest plant sources of zinc. Due to the effects of these compounds, barley has been used as a laxative, diuretic, pain reliever, and heart and blood tonic, as well as reducing blood cholesterol and blood pressure and maintaining the balance between blood sugar and blood insulin²⁰. Therefore, regular consumption of barley is said to reduce the risk of chronic diseases, including diabetes, cancer, obesity, and cardiovascular diseases²¹.

The blood sugar-lowering effects of barley are due to its low glycemic index, high chromium and magnesium content, and water-soluble fibers, including beta-glucan²². Beta-glucan binds to the ingested sugars and lipids and indirectly changes the intestinal environment, thus affecting blood sugar levels (19). So far, many studies have reported barley's blood sugar reduction effects^{18,23–25}. The beneficial metabolic effects of oats in terms of appetite suppression and insulin sensitivity improvement have been attributed to its beta-glucan²⁶. Although there is evidence of the acute effects of barley products on post-prandial glucose, to the best of our knowledge, the short and long term effects of different compartments of barley on blood sugar has not been studied^{27,28}. Therefore, the present study investigated the impact of consuming barley savigh on the blood sugar levels of pre-diabetic postmenopausal women.

Methods Trial design

The present study was conducted as a randomized clinical trial. The study protocol was approved by the Iranian Registry of Clinical Trials (IRCT code: 20200929048880N1). Registration date: (25/10/2020). The study was approved by the Ethics Committee of the Gonabad University of Medical Sciences (Code: IR.GMU. REC.1399.070). The study protocol was published earlier²¹.

This clinical trial was reported based on the CONSORT Statement 2010 checklist²⁹. All methods were performed following the relevant guidelines and regulations.

Participants and setting

The study was conducted on 68 postmenopausal women referring to Community Health Centers in Gonabad city, Iran. Inclusion criteria were menopause (complete cessation of menstruation for at least 12 months), age between 45 and 65 years, pre-diabetic (FBS = 125 mg/dl or two-hour post-prandial plasma glucose level between 140 and 199 mg/dl), not consuming herbal supplements to control blood glucose, literacy, having cardiovascular or other systemic diseases, hypertension, history of smoking, alcohol and drug use, history of grain allergy or celiac disease, and being normal or overweight (Body Mass Index [BMI] between 19 and 28 kg/m²). The exclusion criteria were documented diagnosis of diabetes during the study period, taking anti-diabetic medications or consuming herbal supplements to control blood glucose, need for insulin therapy or anti-diabetic medicines during the study period, occurrence of stressful events, irregular consumption of barley or experiencing complications, including dermatitis, bloating, and diarrhea, hospitalization during the intervention, failure to complete the study questionnaires, occurrence of hypoglycemia based on self-monitoring blood glucose or signs and symptoms of hypoglycemia, and refusing to participate in the study. The complete list of inclusion and exclusion criteria has been published elsewhere²¹.

Sample size and randomization

The sample size for this clinical trial was calculated based on the findings of a previous study, where barley beta-glucan reduced mean FBS from 175.7 ± 29.1 to 140.5 ± 60.9 mg/dl²³.

$$n = \frac{(Z_{1-}\alpha/2 + Z_{1-\beta})^2 \times (S_1^2 + S_2^2)}{(\mu_1 - \mu_2)^2}$$

Based on the following equation and considering type I and II errors of 5% and 20%, respectively, the sample size was determined as 31 participants in each group, which was increased to 34 participants in each group (total sample size of 68 participants) considering 10% dropout.

Study instruments

- a. *Demographic characteristics questionnaire*: This questionnaire included age, BMI, systolic and diastolic blood pressure, duration of pre-diabetes, duration of menopause, parity, gravida, number of abortions, still-birth, and live children. The questionnaire was filled out based on interviews or from their medical records.
- b. *Three-Day Food Record*: Since the daily diet is an essential factor in blood sugar levels, it was controlled as a confounding factor in this study. For this purpose, all participants were asked to record their food intake during three-week days. A three-day food record was obtained at baseline, during the first two weeks of the intervention, and during the second two weeks. The validity of the three-day food record has been confirmed by the content validity method, and the reliability was approved using an r coefficient of > 0.999 in a previous study²⁴.
- c. *Beck Physical Activity Questionnaire*: Since physical activity is an essential factor in blood sugar levels, it was controlled as a confounding factor in this study. For this purpose, the Beck Physical Activity Questionnaire was used. This questionnaire is an international and standard tool for assessing the level of physical activity³⁰. This questionnaire includes 16 questions that are scored using a four-point Likert scale. Beck's physical activity questionnaire consists of three sections: work-related, sports, and leisure-time physical activity. Based on the intensity of physical activity, each question is scored from one, indicating the lowest level of physical activity, to five, indicating the highest level. The level of physical activity for each participant is obtained by summing up the score of each question. The validity and reliability of this questionnaire have been checked and confirmed in different countries, different groups, and Iran. In Iran, the validity and reliability of the Beck's physical activity questionnaire was approved^{27,31}.
- d. *Barley Savigh consumption and complication checklist*: Participants in the intervention group were asked to complete the checklist daily. The checklist included recording daily consumption of barely savigh and questions on the experience of bloating or increased frequency of defecation.

Intervention

The researcher referred to the community health centers of Gonabad city to identify the women based on the inclusion criteria. Eligible women were approached, and the study's aim and procedure were described. In case of agreement, informed and written consent was obtained from the women included in the study. To diagnose pre-diabetes, 5 ml of venous blood was obtained after overnight fasting (8–10 h) to measure FBS, and 5 ml was obtained two hours after eating a standard breakfast to measure 2-hour post-prandial blood glucose. All tests were performed in one laboratory to avoid laboratory errors.

The eligible participants were randomly assigned to the intervention and control groups using four permutation blocks. At the beginning of the study, all participants completed the demographic questionnaire, and FBS and 2-hour post-prandial blood sugar were recorded. Participants also completed a three-day food record and Beck's physical activity questionnaire.

The participants in the intervention group were asked to consume 60 g of barley savigh (20 g of savigh five minutes before each main meal/three times daily), preferably in dry form, for four weeks. This dose was determined based on previous studies that indicated that at least 4 g of beta-glucan and 30–80 g of available carbohydrates from barley could reduce blood sugar²⁰.

For the preparation of barley savigh, barley was bought from a local farmer in the region, and a sample from the barley was sent to the Gonabad Agricultural and Natural Resources Research and Training Center for identification. The identified sample was approved by the Gonabad Agricultural and Natural Resources Research and Training Center (Herbarium Code: Hordeum vulgare L. Family: poaceae. voucher sp.no.E-1401 FUMH). Then, the mill and bran were separated and ground. The mill flour was roasted. Bran was roasted and then ground and mixed with roasted barley flour.

Participants were asked to complete the barley savigh consumption and complications checklist daily to ensure regular consumption of barely savigh. As mentioned in the exclusion criteria, participants with irregular consumption of barley or experiencing complications were excluded from the study. The control group received no intervention.

The four-week study was chosen due to the findings of previous studies that indicated short-term changes in FBS and post-prandial glucose, which can be seen within four weeks in nutrition intervention studies that may reduce dropout from the study $^{32-34}$.

All participants in the intervention and control groups were given the same training about healthy diet, foods that increase blood sugar that should be avoided, and information on the number of appropriate meals and snacks to control the effect of confounding factors. Participants were asked not to change their physical activities during the study.

In both groups, FBS and 2-hour post-prandial blood glaucus were evaluated at the end of the second and the fourth weeks after the start of the intervention. Evaluating the FBs and 2-hour post-prandial blood glaucus in the second week was made better to determine the trend in changes in the variables. They also simultaneously filled out the Beck Physical Activity Questionnaire and Three-Day Food Record.

Statistical analysis

After collecting the data, statistical analysis was done using the Statistical Package for Social Sciences (SPSS) version 16 software. The normality of the continuous variables was evaluated using the Kolmogorov-Smirnov test. The independent t-test (for normally distributed variables) or Mann-Whitney (for non-normally distributed variables) was used to compare continuous variables between groups. The chi-square test was used to compare the distribution pattern of the categorical variables between groups. The statistical significance level was considered as p < 0.05.

Results Study dropouts

Out of 68 participants, two participants from the intervention group were excluded from the study due to lack of regular consumption of the barley savigh (n=1) and hospitalization (n=1), and two participants from the control group were excluded due to not completing the questionnaires (n=1) and refusal to continue the study (n=1). Therefore, 64 participants remained in the study (32 participants in each group) (Fig. 1).

Demographic characteristics

In both groups, 15 (46.9%) participants were rural, and 17 (53.1%) were urban residents, which was not statistically different based on the chi-square test (p = 0.617). The mean age of the participants in the intervention and control groups was 56.86 ± 5.86 and 56 ± 6.02 years, respectively. There was no significant difference in age between the two groups (p = 0.615). Demographic characteristics of the participants are shown in Table 1. There was no significant difference between groups regarding demographic characteristics (p > 0.05).

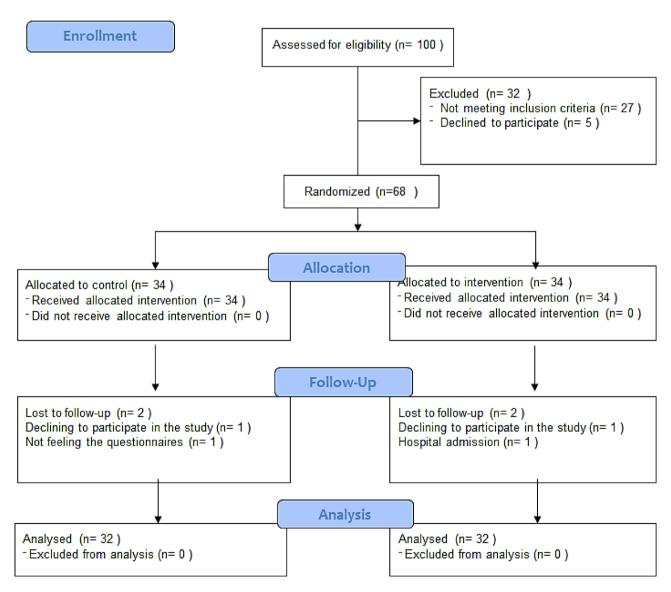


Fig. 1. The flow diagram of the study.

Variable	Intervention Mean ± SD	Control Mean±SD	p
Age (years)	56.81 ± 5.86	57.56 ± 6.02	0.615
Systolic blood pressure (mmHg)	119.90 ± 21.09	122.43 ± 21.46	0.636
Diastolic blood pressure (mmHg)	72.18 ± 6.94	72.62 ± 6.92	0.052
BMI (kg/m²)	26.03 ± 1.58	25.73 ± 1.97	0.501
Duration of pre-diabetes (years)	2.28 ± 0.92	2.32 ± 0.94	0.842
Duration of menopause (years)	7.81 ± 4.90	7.79 ± 4.96	0.990
HbA1C	5.39 ± 0.62	5.49 ± 0.59	0.486
Gravida	4.21 ± 1.51	4.12 ± 1.62	0.822
Parity	3.59 ± 1.16	3.62 ± 1.51	0.927
Number of abortions	0.62 ± 0.75	0.43 ± 0.61	0.280
Number of stillbirths	0.03 ± 0.17	0.03 ± 0.17	> 0.999
Number of live children	3.59 ± 1.16	3.62 ± 1.51	0.927

Table 1. Comparison of the demographic characteristics between the intervention and control groups. SD: Standard Deviation, BMI: Body Mass Index. The independent t-test was used for the comparison.

Variable		Intervention Mean ± SD	Control Mean ± SD	p
Physical activity (score)	Baseline	41.37 ± 3.59	40.37 ± 4.17	0.300
	Week 2	41.40 ± 3.66	40.28 ± 4.06	0.450
	Week 4	41.25 ± 3.62	40.25 ± 4.27	0.550
	Baseline	25.28 ± 1.72	25.40 ± 1.64	0.500
Bread (serving)	Week 2	26.03 ± 1.59	26.15 ± 1.60	0.650
	Week 4	25.37 ± 1.53	25.62 ± 1.82	0.760
	Baseline	7.06 ± 1.34	7.68 ± 1.30	0.081
Meat (serving)	Week 2	7.37 ± 1.07	7.59 ± 0.94	0.400
	Week 4	6.90 ± 0.92	7.18 ± 0.82	0.150
Milk (servings)	Baseline	8.65 ± 1.38	8.34 ± 1.28	0.230
	Week 2	8.00 ± 1.64	7.78 ± 1.56	0.390
	Week 4	8.00 ± 1.07	7.71 ± 1.08	0.260
Fruit (servings)	Baseline	6.25 ± 0.98	6.00 ± 1.01	0.360
	Week 2	6.96 ± 0.93	6.87 ± 1.03	0.570
	Week 4	6.21 ± 0.94	6.06 ± 1.04	0.480
	Baseline	11.84±1.64	12.15 ± 1.41	0.330
Vegetable (servings)	Week 2	12.93 ± 1.07	12.65 ± 1.23	0.330
	Week 4	11.50 ± 1.01	11.84 ± 1.48	0.450
Fat (servings)	Baseline	5.78 ± 0.90	5.40 ± 0.87	0.110
	Week 2	5.03 ± 0.73	4.90 ± 0.64	0.540
	Week 4	5.15 ± 0.72	4.68 ± 0.69	0.013*

Table 2. Comparison of nutrition intake and physical activity between the intervention and control groups at study time points. SD: Standard Deviation. Physical activity scores were presented based on Beck physical activity questionnaire. The Mann-Whitney test was used for the comparison. * Significant difference.

A comparison of physical activity and nutrition intake between the intervention and control groups at study time points is shown in Table 2. There was only a significant difference in fat intake between groups at week four (p=0.013). This finding indicated that fat intake was significantly higher in the intervention group compared to the control group at week four.

Effects of barley savigh on FBS

Comparison of the values of FBS between the intervention and control groups at study time points are shown in Table 3; Fig. 2a. There was a significant difference between groups regarding FBS at week four (p=0.019). This finding indicated that FBS was significantly lower in the intervention group compared to the control group. Within the group, the comparison revealed a significant difference in FBS between study time points (p<0.001) only among the participants in the intervention group. This finding indicated that FBS significantly decreased over time in the intervention group. Furthermore, the trend in the change in FBS was different between the groups, indicating a time-dependent reduction in FBS in the intervention group (Fig. 2a).

Variable		Intervention Mean ± SD	Control Mean ± SD	P (Between group) †
	Baseline	109.31 ± 5.06	109.03 ± 8.49	0.904
FBS (mg/dl)	Week 2	105.25 ± 5.92	108.06 ± 8.75	0.130
	Week 4	102.93 ± 7.75	107.90 ± 8.29	0.019*
P (within grou	up) ‡	P<0.001	0.350	-

Table 3. Comparison of FBS between the intervention and control groups at study time points. SD: Standard Deviation, FBS: Fasting Blood Sugar. The p values presented in the right column refer to the comparison of FBS values between the intervention and control groups at each time point (rows) and the p values in the buttom row refer to the comparison of the FBS values between baseline, weeks 2 and 4 in each intervention and control group. † The Mann-Whitney test was used for the comparison.. ‡ The Friedman test was used for the comparison.. * Significant difference.

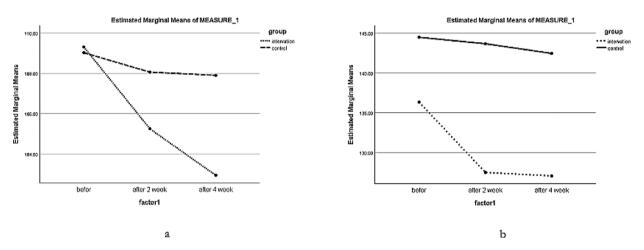


Fig. 2. Changes in FBS (**a**) and 2-hour post-prandial blood sugar (**b**) levels in the intervention and control groups after adjusting for fat intake.

Variable		Intervention Mean ± SD	Control Mean ± SD	P (Between group) †
2-hour post prandial blood sugar (mg/dl)	Baseline	136.34 ± 14.07	144.50 ± 26.09	0.120
	Week 2	127.53 ± 8.03	143.68 ± 26.46	0.006*
	Week 4	127.09 ± 17.96	142.46 ± 27.06	0.009*
P (within group) ‡		P<0.001	0.340	-

Table 4. Comparison of 2-hour post-prandial blood sugar between the intervention and control groups at study time points. SD: Standard Deviation, FBS: Fasting Blood Sugar. † The Mann-Whitney test was used for the comparison.. ‡ The Friedman test was used for the comparison.. * Significant difference.

Effects of barley savigh on 2-hour post-prandial blood sugar

Comparison of 2-hour post-prandial blood sugar between the intervention and control groups at study time points are shown in Table 4; Fig. 2b. There was a significant difference between groups regarding 2-hour post-prandial blood sugar at week two (p=0.006) and four (p=0.009). These findings indicated that 2-hour post-prandial blood sugar levels were significantly lower in the intervention group compared to the control group. The within-group comparison revealed a significant difference in 2-hour post-prandial blood sugar levels between study time points (p<0.001) only among the participants in the intervention group. This finding indicated that 2-hour post-prandial blood sugar levels significantly decreased over time in the intervention group (Table 4). Furthermore, the trend in the change in 2-hour post-prandial blood sugar was different between the groups, indicating a time-dependent reduction in 2-hour post-prandial blood sugar in the intervention group (Fig. 2b).

Discussion

Barley is a good source of fiber. The beta-glucan soluble fiber in barley can reduce the absorption of sugar in the digestive system²⁴. Therefore, this study used barley savigh as a potential blood sugar-reducing agent that can help reduce post-prandial blood sugar.

Effects of barley savigh on FBS

In the present study, FBS was significantly lower in the intervention group compared to the control group four weeks after the intervention. Furthermore, there was a significant decrease in FBS in the intervention group over time. The FBS levels slightly decreased in the control group. Still, this decrease was neither statistical nor clinically important and could be due to the effects of their education on a healthy diet. This observation might be due to the soluble fiber (beta-glucan) content of barley savigh. Although beta-glucan can have acute effects on gastric emptying and glucose absorption, its effect on gut microbiota can be the reason for its persistent effects of 5,36. This hypothesis can be justified by the observed significant reduction in FBS at the final assessment, where it can be hypothesized that the microbiota had reached the adequate quantity to affect FBS. To the best of our knowledge, no human studies evaluated the effects of barley savigh on FBS in postmenopausal women. Previous studies' findings regarding the effect of barley products on FBS were controversial. An earlier study reported that daily consumption of barley bread containing 4 g beta-glucan with or without aerobic exercise for 12 weeks reduced FBS and blood pressure and improved body composition in type 2 diabetic women²³. An animal study observed a significant decrease in fasting serum glucose four weeks after barley consumption in diabetic rats²².

In contrast, an animal study showed that none of the tested barley seed hydroalcoholic extract doses (0.1, 0.25, 0.5 g/kg) effectively reduced FBS among healthy and diabetic rats in the acute phase of treatment (first day). Nevertheless, barley hydroalcoholic extract at doses of 0.25 and 0.5 g/kg reduced FBS in diabetic rats only after 11 days of continued daily consumption²⁷. The reason for the difference in the findings of previous studies might be related to the study subjects (animals and humans) and different dosages and forms of barley product administration (barley bread vs. barley seed hydroalcoholic extract). However, considering the effects of barley priducts on FBS in the mentioned studies, it can be implied that the process of formulation, duration, and dosage of the barley extracts are the key factors that determin its effects on blood sugar and that the dose used in the present study could reduce FBS efficiently without important complications.

Effects of barley savigh on post-prandial blood sugar

In the present study, 2-hour post-prandial blood sugar two and four weeks after the intervention was significantly lower in the intervention group compared to the control group. Furthermore, there was a significant decrease in post-prandial blood sugar in the intervention group over time. Similar to FBS, the observed non-significant decrease in post-prandial blood sugar in the control group could be due to the effects of the education they received in a healthy diet. Similar to the findings of the present study, a systematic review that examined the impact of barley products consumption on post-prandial blood sugar reported that whole grains and a variety of barley foods that contain at least 4 g of beta-glucan and 30 to 80 g of carbohydrates, can reduce post-prandial blood 18 which was in line with the present study that used 60 g of barley savigh containing 40 g of carbohydrates. In a cross-over randomized controlled trial study on 67 healthy normoglycemic adults, the acute effects of administration of barley dietart fiber was evaluated. The study revealed that barley dietary fiber reduced postprandial glycemia and improved satiety in the intervention group²⁸. In another survey of 20 men, beta-glucan derived from barley better controlled post-prandial rise in blood glucose level compared to resistant starch³⁵. Beta-glucan is a soluble fiber that delays gastric emptying by increasing the smack intestine viscosity³⁷. It is also believed that it can reduce glucose absorption in the intestine³⁸. A study on 17 obese women at risk of developing insulin resistance showed that consuming breakfast cereals containing 10 g of beta-glucan in oats significantly reduced post-prandial blood sugar levels in these people³⁰. The results of a study on the effects of barley consumption on post-prandial blood sugar in Duke's type diabetic patients who did not receive antidiabetic medications showed that rice mixed with barley reduced post-prandial glucose concentration³¹. These researchers stated that using diets containing barley as a therapeutic diet may be useful in improving blood sugar control in diabetic patients³¹.

Strengths and weaknesses

One of the strengths of this study was the form of barley product that was used, which included all parts of barley seeds. One of the limitations of this study was the individual differences in terms of unknown and unpredictable neurological factors that affect blood sugar, which could not be comprehensively controlled. Also, it is suggested that the positive effects of barley savigh on glycemic control and lipid profile be evaluated over a longer period. Although insulin measurement could provide more accurate data on the impact of barley savigh on blood glucose, we could not measure insulin levels due to limited financial resources and laboratory facilities. It is suggested that insulin levels be examined in future studies.

Conclusion

Barley savigh can be important in regulating blood sugar in pre-diabetic postmenopausal women. Considering the importance of fiber consumption and its beneficial effects in preventing and controlling diabetes and other metabolic diseases, conducting more research can be helpful in this field.

Data availability

The datasets generated and analysed during the current study are not publicly available but are available from the corresponding author on reasonable request.

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Author contributions

Study concept and design: N. B., and K. K.; analysis and interpretation of data: F. HT., and S. TZ.; drafting of the manuscript: N. K. and S.D.; critical revision of the manuscript for important intellectual content: S. D. and N. B.; statistical analysis: N. K. All authors read and approved the final manuscript.

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Declarations

Ethics approval and consent to participate

The study was approved by the Ethics Committee of the Gonabad University of Medical Sciences (Code: IR.GMU.REC.1399.070). All methods were performed following the relevant guidelines and regulations.

Consent for publication

Not applicable.

Competing interests

The authors declare no competing interests.

Additional information

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