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Effectiveness of a Mindfulness-Based Stress Reduction (MBSR) Program on Stress, Anxiety, and Prenatal Attachment for High-Risk Pregnant Women: A Randomized Controlled Trial

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Abstract

Objective The present study was conducted to examine the effects of the Mindfulness-Based Stress Reduction (MBSR) Program on stress, anxiety, and prenatal attachment of the high-risk pregnant women.

Method This randomized controlled study was conducted with 94 high-risk pregnant women (48 participants in the experimental group, 46 participants in the control group). In the study, the experimental group underwent an eight-session MBSR program, consisting of two sessions per week for a total duration of 1 month. Data of the study were collected with the Prenatal Distress Questionnaire-Revised (NuPDQ), Pregnancy-Related Anxiety Questionnaire-Revised 2 (PRAQR2), and Prenatal Attachment Inventory (PAI).

Results After the intervention, the PRAQR2 total mean score in the experimental group was significantly lower than that in the control group (p < 0.05). In addition, the PAI total mean score in the experimental group was significantly higher than that in the control group (p < 0.05). However, there was no significant difference between the groups in the NuPDQ total mean score (p > 0.05).

Conclusions It was determined that the MBSR program applied to high-risk pregnant women decreased anxiety levels, increased prenatal attachment levels, and did not affect stress levels of the pregnant women. Health professionals may consider the MBSR program for high-risk pregnant women as a means by which to improve their pregnancy outcomes. **Preregistration** NCT05317065.

Keywords Anxiety · Prenatal attachment · MBSR · High-risk pregnancy · Stress

Although most pregnancies and births occur without problems, it is reported that all pregnant women are at risk, and approximately 15% of pregnant women may develop a life-threatening complication (World Health Organization,

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2017). High-risk pregnancies refer to any situation that situates the mother, fetus, or newborn at increased risk for morbidity or mortality during pregnancy or childbirth (Nesro et al., 2021; Zaidan et al., 2018). The presence of risky situations during pregnancy also has a negative effect on psychological health (Bahadır Yılmaz & Şahin, 2019; Çağlar & Yeşiltepe Oskay, 2019; Dagklis et al., 2016). It is known that the distress (Gözüyeşil & Ariöz Düzgün, 2021) and anxiety (Pisoni et al., 2016; Rodrigues et al., 2016; Smorti et al., 2023) levels of high-risk pregnant women, especially those who are hospitalized, are high. Gözüyeşil and Ariöz Düzgün (2021) reported in their study with high-risk pregnant women that more than half of the participants (63.5%) experienced distress (Gözüyeşil & Ariöz Düzgün, 2021). Furthermore, it is stated that 68% of high-risk pregnant women experience moderate to high levels of anxiety (Paz et al., 2022). It has been reported that the prevalence of pregnancy-related anxiety (26.5%) is higher than the prevalence of general anxiety (16.4%) in pregnant women (Naja et al., 2020). Pregnancy-related anxiety differs from general anxiety and depression symptoms, and refers to fears and worries specific to pregnancy, including the health of the fetus, childbirth, and future parenting concerns (Huizink et al., 2004). Pregnancy-related anxiety and distress are associated with adverse birth outcomes (Faramarzi et al., 2019; Madhavanprabhakaran et al., 2015). Moreover, it is stated that health problems experienced during pregnancy can increase the level of anxiety and affect mother-baby attachment (Yeşilçınar et al., 2023).

Prenatal attachment is a multidimensional emotional bond that includes the mother's thoughts, behaviors, and attitudes towards her unborn child (Bakır et al., 2014; Filippetti et al., 2022). This bond begins with the woman's desire to become pregnant and strengthens as she feels fetal movement and embraces her maternal roles (Topaç Tuncel & Kahyaoğlu Süt, 2019). A risky situation that may occur during pregnancy negatively affects the woman's adaptation to pregnancy and the process of gaining the maternal role that begins with pregnancy, and at the same time, it causes confusion in maternal roles by feeling guilty and can reveal sources of stress (Baltacı & Başer, 2020; Deliktaş et al., 2015). Eswi and Khalil (2012) found in their study that the attachment level of high-risk pregnant women was lower than low-risk pregnant women (Eswi & Khalil, 2012). Pisoni et al. (2016) similarly reported in their study that parents who are at risk of premature birth and are hospitalized and expecting a baby have higher anxiety and depression levels and lower attachment levels than the physiological pregnant group (Pisoni et al., 2016). Medical risks and maternal risks both increase the risk of pregnancy and adverse obstetric outcomes (Kuppusamy et al., 2023).

It is extremely important to implement psychological interventions to reduce the psychological distress of pregnant women and increase mother-baby well-being (Zhang et al., 2023). Although there is no consensus on the definition of mindfulness, it is generally defined as a mental state that involves focusing on the present moment in a non-judgmental manner, with an attitude of openness and acceptance of experiences (Lucena et al., 2020). Mindfulness-Based Stress Reduction (MBSR) program is designed in 8-week sessions and is one of the most well-known mindfulness intervention. The MBSR program includes body scans, gentle stretching, and yoga mindfulness exercises, as well as a wide range of holistic techniques and therapies to apply mindfulness to daily life experiences, including coping with stress (Creswell, 2017). Studies have also emphasized that mindfulness interventions reduce the stress and anxiety levels of high-risk pregnant women and are effective on prenatal attachment in pregnant women (Gheibi et al., 2020; Güney et al., 2022; Shreffler et al., 2019; Woolhouse et al., 2014). Additionally, Dhillon et al. (2017), in a systematic review and meta-analysis of mindfulness-based interventions in pregnancy, have stated that the MBSR program can be used to solve problems such as anxiety, depression, and perceived stress and that the practice increases the awareness levels of pregnant women, reduces prenatal anxiety, and can be used as an effective method to cope with maternal distress (Dhillon et al., 2017). Although the MBSR has been used in many areas recently, no study has been found in which the effects of the MBSR program on stress, anxiety, and prenatal attachment of high-risk pregnant women are examined together. It was aimed that the results of this study, which was conducted to examine the effect of the MBSR program on stress, anxiety, and prenatal attachment of the high-risk pregnant women, will provide important contributions to health professionals who provide care services in the antenatal period regarding the effectiveness of MBSR program. We hypothesized that the MBSR program provided to high-risk pregnant women affects prenatal distress levels, pregnancyrelated anxiety levels, and prenatal attachment levels.

Method

Participants

The current study was carried out between the dates of June and August 2022 at the 15th Family Health Center (FHC) of a city center located in the Southeastern Anatolia Region of Turkey. Seven healthcare professionals, including three physicians, three midwives, and one nurse, were working at the FHC. The total population of FHC was around 9000 on average. The working hours of FHC were between 08:00-12:00 and 13:00-17:00. Pregnant women who were diagnosed with high-risk pregnancies registered to the FHC were referred to the Healthy Living Center (HLC) in the city center and the high-risk pregnancies were reported. The population of the study consisted of pregnant women who were registered to the 15th FHC in a city center located in the Southeastern Anatolia Region of Turkey at the relevant dates and diagnosed with high-risk pregnancies and reported to the HLC. The minimum number of individuals to be included in the sample of the study was determined by power analysis. In the literature review, it was determined the mean pregnancy-related anxiety score in the high-risk pregnant women, was previously reported as 30.53 (SD = 6.49) (Duman et al., 2021). The information in the literature shows that an approximately 4- or 5-point difference in scale scores can be clinically and statistically significant in a different intervention study conducted with pregnant women using same measurement tools (Aksoy Derya et al., 2021). The sample size was calculated as

84 pregnant women (42 participants for the experimental group, 42 participants for the control group) assuming that the method applied with a 5% error level, 95% confidence interval, and 80% ability to represent the universe would reduce the anxiety mean score (30.53, SD = 6.49) by 4 points. Considering possible case losses (20%), a total of 100 pregnant women were planned to be included in the study. The final sample consisted of 94 high-risk pregnant women. The pregnant women were 18–44 years old, in a romantic relationship, then nearly half of the participants had similar income perception, and at the time of recruitment participants had a gestational age between 12 and 27.

Inclusion criteria were as follows: pregnant women who were between the 10th and 28th week of gestation, diagnosed with high-risk pregnancies, able to understand Turkish, used smartphones, had Internet network and access, had headset accessories that they could use in the interviews, and voluntarily accepted to participate in the study were included in the study.

Exclusion criteria were as follows: pregnant women who had participated in mind-body practices or any other types of prenatal classes/programs during the study period. Other exclusion criteria included unwillingness to continue to participate in the study, having a psychological problem. Psychiatric health problems were determined according to the declaration and medical history of the pregnant women.

The high-risk pregnant women were randomly assigned to two groups, namely experimental and control group. Columns between 1 and 100 were created using the "Random Integer Generator" method, which is located in the "Numbers" sub-heading of the "random.org" site for the women who met the sample selection criteria. The numbers 1 and 2 for the experimental and control groups were observed in the column. Which number would represent the experimental or control group was determined by drawing lots at the beginning of the research. The women who matched the number 1 as a result of the lottery were assigned to the experimental group, and the women who matched the number 2 were assigned to the control group (Random.org). The randomization process was conducted by a non-involved person in the sampling and analysis.

Procedure

The data of the study were collected from pregnant women who were registered to the 15th FHC in a city center in the Southeastern Anatolia Region of Turkey between the dates of June and August 2022, diagnosed with the high-risk pregnancies, and reported to the HLC. Pregnant women were invited to participate in the study by contacting them by phone. The pregnant women who accepted to participate in the study, met the inclusion criteria, and were assigned to the experimental and control groups were informed (Whatsapp) by the researcher and asked to fill in web-based online questionnaires prepared through Google Forms. The informed consent form was presented on the first page of the online questionnaire. As a pre-test, the Personal Information Form, Prenatal Distress Questionnaire-Revised (NuPDQ), Pregnancy-Related Anxiety Questionnaire-Revised 2 (PRAQR2), and Prenatal Attachment Inventory (PAI) were applied to the high-risk pregnant women in the experimental and control groups. After the MBSR program (assignments, exercises, and applications of breathing and body) was applied online by the researcher to the experimental group for 1 month, two sessions each week, a total of eight sessions, post-test data was obtained by re-applying the the NuPDQ, PRAQR2, and PAI to the experimental group and simultaneously to the control group.

Intervention

In order to evaluate the effect of the MBSR program on stress, anxiety, and prenatal attachment of the high-risk pregnant women, the MBSR was applied to the high-risk pregnant women in the experimental group for four weeks (two sessions per week for a total of eight sessions). The intervention protocol of this study covers the entire MBSR program, which consists of two sessions per week for 1 month and home assignments given throughout the program. In all sessions, mindfulness exercises such as 3-min breathing, body and breathing exercises, body scanning meditation, sitting meditation, mountain meditation, and compassion meditation were applied to the participants under the guidance of the researcher (S.O.C.), who was trained and has a certificate in the field of MBSR. It was done in sessions and individual applications in an environment where pregnant women were not disturbed at any time of the day. The sessions of the MBSR program were based on the meditation techniques of Mindfulness Therapy (Kabat-Zinn, 1990). The live online MBSR program was provided via mobile communication tools. Sessions were conducted as individual or group sessions according to choice of high-risk pregnant women by determining the days and time when they were available online. Each session lasted an average of 40-60 min. The pregnant women in the control group received usual hospital care and they were applied no intervention. After the collection of the posttest data, the participants in the control group were offered mindfulness training at no cost.

Measures

Personal Information Form This is a form consisting of 16 questions developed by the researchers in line with the literature, with the aim of determining the women's socio-demographic (age, spouse's age, educational level, employment

status, spouse's employment status, perceived income level) and obstetric (week of gestation, number of pregnancies, number of abortions/curettage, number of births, number of living children, the status of having a planned pregnancy, method of previously terminated pregnancy, the time period between the present pregnancy and the previous delivery, the status of getting pregnant with treatment, the risk status of pregnancy) characteristics (Bakır et al., 2014; Dereli Yılmaz & Kızılkaya Beji, 2010; Duman et al., 2021; Yüksel et al., 2020).

Prenatal Distress Questionnaire-Revised Prenatal distress was measured using the validated 17-item NuPDQ. The NuPDQ assesses pregnant women's concerns about social relationships, physical and emotional symptoms, themselves, and their baby. The Turkish validity and reliability study of the questionnaire was conducted by Yüksel et al. (2011). The questionnaire is in the form of 3-point Likert type. Each item in the questionnaire was scored as 0 = Notat all, 1 = A little, and 2 = A lot. A minimum of 0 and a maximum of 34 points can be obtained from the questionnaire. An increase in the scores obtained from the questionnaire is interpreted as an increase in prenatal distress (Ibrahim & Lobel, 2020; Lobel et al., 2008; Yali & Lobel, 1999; Yüksel et al., 2011). In the Turkish validity and reliability study of the scale, Cronbach's alpha coefficient was reported as 0.85 (Yüksel et al., 2011). In the present study, the Cronbach's alpha coefficient and McDonald's omega were calculated as 0.94 for the NupDQ.

The Pregnancy-Related Anxiety Questionnaire-Revised

2 The PRAQR2 was developed by Van der Bergh (1990), and Huizink et al. (2016) revised it. The PRAQR2 was developed to question the level of anxiety experienced by women about their pregnancies. The Turkish adaptation of the questionnaire was made by Aksoy Derva et al. (2018). The Turkish version of the questionnaire consists of 10 items for multiparas and 11 items for primiparas, and the questionnaire is in the format of a 5-point Likert type. There are 3 subscales in the questionnaire: *Fear of giving* birth, Worries about bearing a handicapped child, and Concern about one's own appearance. The 8th item in the questionnaire is applied only to primiparous women. Items are scored between 1 and 5, where 1 = Absolutelynot relevant and 5 = Very relevant. A minimum of 11 and a maximum of 55 points can be obtained from primiparas, and a minimum of 10 and a maximum of 50 points can be obtained from multiparas. As the score obtained from the questionnaire increases, the level of anxiety during pregnancy also increases (Aksoy Derva et al., 2018; Huizink et al., 2016; Van der Bergh, 1990). In the Turkish validity and reliability study of the questionnaire, Cronbach's alpha coefficient was reported as 0.93 in multiparas and 0.94 in nulliparas (Aksoy Derya et al., 2018). In the current study, the Cronbach's alpha coefficient was determined to be 0.94 and McDonald's omega was calculated as 0.95 for the PRAQR2.

Prenatal Attachment Inventory The PAI was developed by Muller (1993) to explain the thoughts, feelings, and situations experienced by women during their pregnancies and to determine the level of attachment to their babies in the prenatal period. Turkish validity and reliability study was carried out by Dereli Yılmaz & Kızılkaya Beji, (2013). Each item is evaluated between 1 and 4 points, and the inventory is in the 4-point Likert type. There are 21 items in total in the inventory, and the items are scored as 1 = Never, 2 = Sometimes, 3 = Often, and 4 = Always. A minimum of 21 and a maximum of 84 points can be obtained from the inventory. As the scores obtained from the inventory increase, the level of attachment also enhances (Dereli Yılmaz & Kızılkaya Beji, 2013; Muller, 1993). In the Turkish validity and reliability study of the inventory, Cronbach's alpha coefficient was reported as 0.84 (Dereli Yılmaz & Kızılkaya Beji, 2013). In this study, the Cronbach's alpha coefficient and McDonald's omega was calculated as 0.94 for the PAI.

Data Analyses

The obtained data were analyzed using the program Statistical Package for the Social Sciences (SPSS) for Windows 25.0. Kolmogorov–Smirnov test was used to determine whether the data presented a normal distribution. It was seen that the data were normally distributed. Descriptive statistics included numbers, percentage distribution, mean, standard deviation, and min–max values. The descriptive and obstetric characteristics were compared through chisquare test, and independent samples *t*-test. The mean scores of stress, anxiety, and prenatal attachment were compared before intervention using the independent samples *t*-test. To test for the intervention effects, ANCOVA test with controlling the baseline scores were conducted. Effect sizes were reported as partial η^2 . The level of significance was accepted as p < 0.05.

Results

Due to the fact that two pregnant women in the experimental group could not be reached during the post-test application, and four pregnant women in the control group did not complete the post-test voluntarily, the study was completed with a total of 94 high-risk pregnant women (48 participants for the experimental group, 46 participants for the control group). In addition, 16 pregnant women did not accept to participate in the study, and 4 pregnant women could not be included in the study because they did not have smartphones/or Internet access (Fig. 1; CONSORT Research Flowchart). Dropped out and excluded participants did not differ from the final sample on any descriptive and obstetric characteristics.

When the high-risk pregnant women in the experimental and control groups were compared in terms of descriptive characteristics, it was determined that there was no statistically significant difference between the two groups in terms of descriptive characteristics such as age, spouse's age, educational level, employment status, spouse's employment status, and income level (p > 0.05) (Table 1).

When the high-risk pregnant women in the experimental and control groups were compared in terms of obstetric characteristics, the status of having a planned pregnancy, how the previous pregnancy terminated, the time period between the current pregnancy and the previous delivery, the status of conception with treatment, gestational week, number of pregnancies, number of miscarriages/curettages, number of deliveries, and number of living children, there was no statistically significant difference in terms of obstetric characteristics (p > 0.05) (Table 2). Both groups had similar descriptive and obstetric characteristics.

When the pregnancy-related risk status of high-risk pregnant women regarding their pregnancies was examined, it was seen that 33% of them were having pregnancies at early/ advanced age, 21.3% of them had frequent interbirth intervals and anemia, 19.1% of them had obesity, 17% of them had blood incompatibility, 16% of them had consanguineous marriages, 5.3% of them had gestational diabetes and preexisting diabetes, and 42.6% of them had other risk factors (risk of premature birth, multiple pregnancies, history of anomalous baby, preeclampsia, smoking, presence of infection, risk of miscarriage, heart diseases, endocrine diseases) (Table 3).

When the pre-test and post-test NuPDQ total, PAI total, PRAQR2 total, and its subscales' mean scores of the high-risk pregnant women were examined, it was found that the pre-test NuPDQ total mean score of the high-risk pregnant women was 25.74 (SD = 3.45), and the post-test NuPDQ total mean score was 15.56 (SD = 8.61).

It was determined that the pre-test PRAQR2 total mean score of the high-risk pregnant women was 39.33 (SD = 5.05), and the post-test PRAQR2 total mean score was 29.88 (SD = 10.0). In addition, the PRAQR2's subscale of fear of giving birth total mean score was 14.94 (SD = 1.90) in the pre-test, 11.78 (SD = 3.87) in the post-test; the PRAQR2's subscale of worries about bearing a handicapped child total mean score was 13.71 (SD = 2.76) in the pre-test and 9.86 (SD = 3.73) in the post-test; the PRAQR2's subscale of concern about one's own appearance total mean score was 10.67 (SD = 1.89) in the pre-test and 8.24 (SD = 3.13) in the post-test.

It was determined that the pre-test PAI total mean score of the high-risk pregnant women was 42.16 (SD = 7.33), and the post-test PAI total mean score was 49.35 (SD = 9.66) (Table 4).

The inter-group comparisons of the pre-test and posttest total mean scores of the high-risk pregnant women in the experimental and control groups from the NuPDQ, PRAQR2, and PAI were analyzed. When the pre-test NuPDQ and PAI total mean scores of the high-risk pregnant women in the experimental and control groups were compared, it was determined that the differences between the groups were not statistically significant and that the high-risk pregnant women in the experimental and control groups had similar distress and attachment levels before the implementation of MBSR program (p > 0.05). When the pre-test PRAQR2 total mean scores of the high-risk pregnant women in the experimental and control groups were compared, it was determined that the difference between the groups was statistically significant and the anxiety levels of the high-risk pregnant women in the control group before the implementation of MBSR program were lower than those in the experimental group (p < 0.05) (Table 5).

After the intervention, a comparison of the mean score of the stress indicated no statistically significant difference between the groups after adjusting the effect of baseline score (adjusted mean difference, AMD: -12.571; 95% CI (confidence interval): -15.0 to -10.1; p > 0.05). However, the mean score of the anxiety significantly decreased in the experimental group, compared to the control group (*AMD*: -14.917; 95% CI: -17.8 to -11.9; p < 0.001), with large effect size ($\eta^2 = 0.528$). Furthermore, the mean score of the prenatal attachment significantly increased in the experimental group, compared to the control group (*AMD*: 11.554; 95% CI: 8.3 to 14.7; p < 0.001), with large effect size ($\eta^2 = 0.357$) (Table 5).

Discussion

High-risk pregnancy is a physiological and psychosocial condition that threatens the life and health of the mother, fetus, or newborn. High-risk pregnant women often experience stress and anxiety as a result of risk factors during pregnancy and as a result of hospitalization. Well-being in psychosocial health ensures that both the maternal role and maternal attachment are healthy. This study aimed to assess the impact of the MBSR program on stress, anxiety, and prenatal attachment among high-risk pregnant women. The results showed a decrease in anxiety levels and an increase in attachment levels among the high-risk pregnant women in the experimental group, while stress levels remained unchanged after the MBSR program.

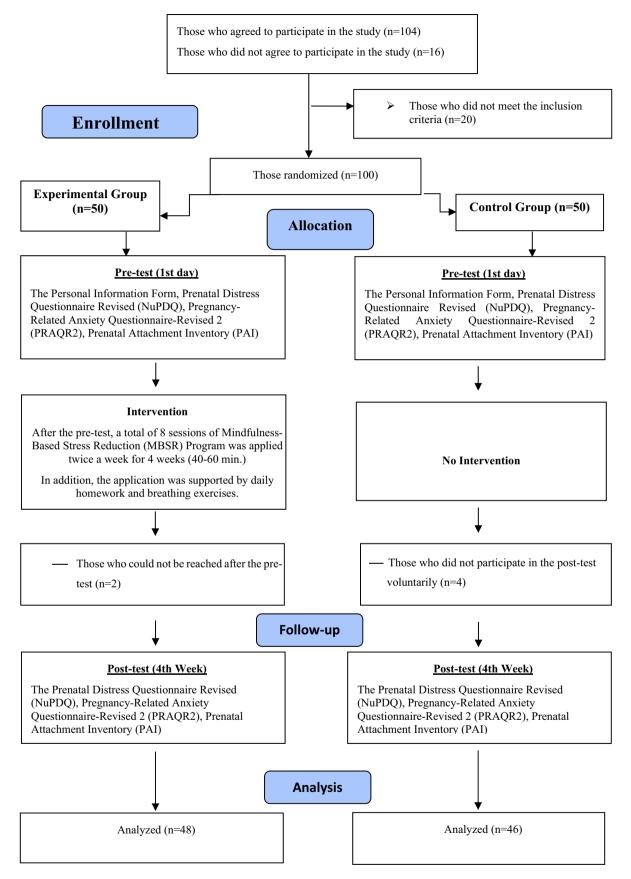


Fig. 1 CONSORT research flowchart

 Table 1
 Comparison of the high-risk pregnant women in experimental and control groups in terms of descriptive characteristics

Descriptive characteristics	Experimental (n=48) n %	Group	Control Group n %	(<i>n</i> =46)	Test and <i>p</i> values
Educational level					
Primary school	2	4.2	1	2.2	
Middle school	7	14.6	8	17.4	$\chi^2 = 0.483$
High school	22	45.8	22	47.8	p = 0.923
University or higher	17	35.4	15	32.6	
Employment status					
Unemployed	25	52.1	31	67.4	$\chi^2 = 2.286$
Employed	23	47.9	15	32.6	p = 0.131
Spouse's employment status					
Employed	42	87.5	40	87.0	$\chi^2 = 0.006$
Unemployed	6	12.5	6	13.0	p = 0.937
Perceived income level					
Income is less than expenses	18	37.5	19	41.3	$\chi^2 = 0.162$
Income is equal to expenses	24	50.0	22	47.8	p = 0.922
Income is more than expenses	6	12.5	5	10.9	
	M(SD)		M(SD)		
Age (years)	31.50 (6.46)		31.46 (6.44)		t = 0.033 p = 0.974
Spouse's age (years)	37.25 (5.87)		37.70 (5.41)		t = -0.382 p = 0.703

 χ^2 = chi-square test, t = independent samples t-test; M mean, SD standard deviation

In the study, it was determined that there was no statistically significant difference between the descriptive and obstetric characteristics of the pregnant women in the experimental and control groups. In line with these results, it can be said that the study groups were homogeneously distributed. When the NuPDQ and PAI total mean scores of the high-risk pregnant women in the experimental and control groups were compared, it was determined that the differences between the groups were not statistically significant, while the pre-test measurement of the PRAQR2 total mean score of the high-risk pregnant women in the control group was lower than those of the experimental group. Therefore, it was seen that the high-risk pregnant women in the experimental and control groups had similar distress and attachment levels before the implementation of the MBSR program. On the other hand, it was seen that the anxiety levels of the pregnant women in the control group were lower, and the pregnant women in the experimental group were more disadvantaged in terms of anxiety. These results may be due to the fact that individuals' situations and risk perceptions are different and that there are many factors from effecting anxiety.

Pregnancy, which is a physiological process, is also a complex process in which anxiety and stress can occur intensely. In fact, this situation is even more evident when pregnancy is risky (Dağlar & Nur, 2014; Gümüşdaş et al., 2014). Because pregnant women with high-risk pregnancies may experience anxiety and stress due to many factors such as health problems, uncertainty, hospitalization, and separation from the family (Baltacı & Başer, 2020). A study evaluating psychosocial health status in the literature has also stated that those with high-risk pregnancies are more likely to develop distress, anxiety, and depression compared to others, and their psycho-social health status is affected more negatively (Gümüşdaş et al., 2014). It is known that mindfulness interventions have positive effects on the distress (Pan et al., 2019; Woolhouse et al., 2014) and anxiety (İbici Akca et al., 2023; Mohamed et al., 2017; Vieten & Astin, 2008; Woolhouse et al., 2014) levels of pregnant women. Our study results show that, although the MBSR program in high-risk pregnant women does not affect the stress experienced, it is an effective method for coping with anxiety. Güney et al. (2022) reported in their study that the MBSR program was effective in reducing the distress and anxiety levels of pregnant women diagnosed with COVID-19 (Güney et al., 2022). Goetz et al. (2020) similarly found in their study that electronic mindfulness-based intervention reduced the anxiety scores of hospitalized high-risk pregnant women (Goetz et al., 2020). Zhang et al. (2023) reported in their study that the Digital Guided Self-help Mindfulness Training reduced the anxiety levels of pregnant women with psychological distress (Zhang et al., 2023). Hulsbosch et al. (2023) reported in their study that the online mindfulness-based Table 2Comparison of thehigh-risk pregnant women inthe experimental and controlgroups in terms of obstetriccharacteristics

Obstetric characteristics	Experimental $(n=48)$ n %	Group	Control Group ($n = 46$) n %		Test and <i>p</i> -values
The status of having a planned pregna	ancy				
Planned	30	62.5	23	50.0	$\chi^2 = 1.492$
Unplanned	18	37.5	23	50.0	p = 0.222
How the previous pregnancy terminated $(n = 73)$					$\chi^2 = 3.321$
Vaginal delivery	16	43.2	19	52.8	p = 0.345
Caesarean section	11	29.8	11	30.5	
Miscarriage/curettage	7	18.9	6	16.7	
Stillbirth	3	8.1	_	-	
The time period between the current	pregnancy and th	ne previo	us delivery		
First pregnancy	13	27.1	9	19.5	
Less than 24 months (2 years)	13	27.1	17	37.0	$\chi^2 = 1.314$ p = 0.518
More than 24 months (2 years)	22	45.8	20	43.5	
The status of conception with treatme	ent				
Conception with treatment	2	4.2	3	6.5	$\chi^2 = 0.259$
Spontaneous conception	46	95.8	43	93.5	p = 0.674
	M(SD)		M(SD)		
Gestational Week	18.50 (5.01)		18.76 (4.79)		t = -0.258 p = 0.797
Number of pregnancies	2.88 (1.63)		3.28 (2.13)		t = -1.042 p = 0.300
Number of Miscarriages/curettages	0.50 (0.96)		0.59 (1.22)		t = -0.383 p = 0.702
Number of deliveries	1.40 (1.10)		1.74 (1.70)		t = -1.163 p = 0.248
Number of living children	1.25 (1.08)		1.70 (1.73)		t = -1.500 p = 0.137

 χ^2 = chi-square test, t = independent samples t-test; M mean, SD standard deviation

Pregnancy-related risk situations*	n	%
Early/advanced age pregnancy	31	33.0
Frequent interbirth interval	20	21.3
Anemia	20	21.3
Obesity	18	19.1
Blood incompatibility	16	17.0
Consanguineous marriage	15	16.0
Gestational diabetes	5	5.3
Pre-existing diabetes		5.3
Other (risk of premature birth, multiple pregnancies, history of anomalous baby, preeclampsia, smoking, presence of infection, risk of miscarriage, heart diseases, endocrine diseases)		42.6

*Participants gave more than one response

intervention was not effective in reducing the distress levels of pregnant women with pregnancy distress (Hulsbosch et al., 2023). Our research results show that the MBSR program applied to high-risk pregnant women effectively reduces anxiety levels and supports the literature. However, there are differences between study findings regarding the effects of mindfulness practices on the distress levels in pregnant women. In line with these results, it can be said that there is a need for further controlled studies that reveal the relationship between mindfulness and distress.

Table 3Distribution ofthe pregnancy-related risk

situations

Table 4Distribution of the high-risk pregnant women's pre-test andpost-test mean scores from the NuPDQ total, PAI total, PRAQR2total, and its subscales

Scales	Pre-test M (SD)	Post-test M (SD)
NuPDQ total	25.74 (3.45)	15.56 (8.61)
PRAQR2 total	39.33 (5.05)	29.88 (10.01)
PRAQR2's subscales		
Fear of giving birth	14.94 (1.90)	11.78 (3.87)
Worries about bearing a handicapped child	13.71 (2.76)	9.86 (3.73)
Concern about one's own appearance	10.67 (1.89)	8.24 (3.13)
PAI total	42.16 (7.33)	49.35 (9.66)

NuPDQ Prenatal Distress Questionnaire Revised, *PRAQR2* Pregnancy-Related Anxiety Questionnaire-Revised 2, *PAI* Prenatal Attachment Inventory, *M* mean, *SD* standard deviation

Danger symptoms and fears of losing the fetus in highrisk pregnant women can negatively affect mother-infant attachment (Baltacı & Başer, 2020; Røhder et al., 2020). In the literature, it has been stated that the prenatal attachment level of high-risk pregnant women is lower than that of non-risky pregnant women (Eswi & Khalil, 2012; Pisoni et al., 2016), and that anxiety and stress that occur during hospitalization and discomfort that cause risky situations negatively affect prenatal attachment (Pisoni et al., 2014). Although it has long been accepted that mindfulness-based practices are a factor that significantly contributes to the

Table 5The comparisonof the mean score of stress,anxiety and prenatal attachmentbetween two groups beforeand after intervention withcontrolling base score effect

health of the mother and child (Golmakani et al., 2021), studies examining the effects of these practices on prenatal attachment are quite limited. Limited studies have reported that mindfulness interventions increase the prenatal attachment levels of pregnant populations (Gheibi et al., 2020; Haghighat et al., 2015; Malhotra et al., 2021; Shreffler et al., 2019). In this study, similarly, it was determined that the prenatal attachment levels of the high-risk pregnant women who received the MBSR program were significantly higher than those who did not receive the MBSR program. Muzik et al. (2012) smilarly reported that mindfulness yoga significantly increased maternal-fetal attachment of pregnant women at high risk for psychopathology (Muzik et al., 2012). According to these results, it is possible to say that the MBSR program is an effective application that increases the prenatal attachment of high-risk pregnant women and based on the available evidence, MBSR can also be used among non-medical methods for high-risk pregnant women.

Limitations and Future Research

The limitations of the study are that pregnant women who did not have a smartphone or Internet access could not be included in the study due to the online conduct of the study. We included all high-risk pregnant women in our study, regardless of risk factor rather than a specific risky condition associated with pregnancy. Anxiety and stress levels may vary depending on the severity of the risk situation during pregnancy and the risk perception of the pregnant

Scales	Experimental Group $(n=48)$ M (SD)	Control Group (n=46) M (SD)	<i>AMD</i> (CI=95%)	Test and <i>p</i> -values	η^2
Before intervention	on				
NuPDQ total	26.06 (2.74)	25.41 (4.06)	-	t = 0.903 p = 0.369	-
PRAQR2 total	40.52 (3.70)	38.09 (5.95)	-	t = 2.369 p = 0.020*	-
PAI total	40.98 (6.46)	43.39 (8.02)	-	t = -1.607 p = 0.111	-
After intervention					
NuPDQ total	9.52 (7.39)	21.87 (4.09)	- 12.5 (- 15.0 to - 10.1)	F = 3.682 p = 0.058	0.039
PRAQR2 total	22.96 (8.94)	37.11 (4.38)	-14.9 (-17.8 to -11.9)	F=101.849 p=0.000**	0.528
PAI total	54.65 (8.33)	43.83 (7.71)	11.5 (8.3 to 14.7)	<i>F</i> = 50.607 <i>p</i> = 0.000 **	0.357

AMD adjusted mean difference, CI confidence interval, NuPDQ Prenatal Distress Questionnaire Revised, PRAQR2 Pregnancy-Related Anxiety Questionnaire-Revised 2, PAI Prenatal Attachment Inventory, M mean, SD standard deviation

 η^2 = partial eta kare, *t* = independent sample *t*-test, *F* = ANCOVA test with adjusting the baseline score was used

p* < 0.05; *p* < 0.001

woman. Therefore, it may be recommended to consider a specific risky situation associated with pregnancy for future studies. This way, the relationship between MBSR program and anxiety and distress can be understood more clearly in this population. Additionally, more studies focusing on low-cost psychological interventions are needed. Future research should include face-to-face practices as an alternative to improve the efficacy and persistence of the intervention, including self-administration at home, and include sample groups of pregnant women experiencing high levels of distress in pregnancy. Other limitations include the relatively small number of participants and the fact that all were selected from only a single health center. Multicenter studies that allow high-risk pregnant women with different socio-cultural characteristics to be included in the research will help generalize the findings. In addition, verifying our results with more sample groups in terms of generalizability of the findings will be important.

According to the results of the study, the MBSR program applied to high-risk pregnant women decreased anxiety, increased prenatal attachment levels, but did not affect stress levels of the pregnant women. Pregnancy is a period in which some important physical and psychological changes are seen, and the addition of risks related to pregnancy to these changes can make the process more difficult and complex, causing it to become inextricable for the woman and her family. For this reason, it is extremely important to manage the changes that occur in women who have a high-risk pregnancy in order to protect and increase the health of the mother and baby. Further research is necessary to validate these findings and explore the full potential of such interventions. Although the literature is limited, studies, like ours, on some populations, have shown that a mindfulness program may help to increase prenatal attachment and reduce anxiety. For this reason, it is recommended that health professionals integrate this program into the care they offer in order to reveal the positive effects of the MBSR program.

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Author Contributions E.I.A.: conceptualization; data curation; formal analysis; investigation; methodology; supervision; visualization; validation; writing—original draft; writing—review and editing. S.O.C.: data curation; investigation; methodology; validation; writing—original draft; writing—review and editing. N.G.: investigation; methodology; validation; writing—original draft; writing—review and editing.

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Data Availability The datasets used and/or analyzed during the current study are available from the corresponding author upon reasonable request.

Declarations

Ethics Statement Approval was obtained from the Scientific Research and Publication Ethics Committee of Inonu University (Decision Number: 2022/3280). In addition, necessary permissions were obtained from the relevant institution in order to conduct the research. The highrisk pregnant women who participated in the study were informed that they were free to participate in the study or not and that they could leave the study at any time.

Informed Consent Informed consent and consent to publish were obtained from all individual participants included in the study.

Conflict of Interest The authors declare no competing interests.

Use of Artificial Intelligence Artificial Intelligence was not used.

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