RESEARCH

Open Access



The association between global and prime diet quality scores and the risk of bacterial vaginosis: a secondary analysis of case-control study

Sanaz Mehrabani¹, Mozhgan Hafizi Moori², Morvarid Normohammadi^{3,4}, Marzieh Shoja⁵, Sevda Eskandarzadeh⁴, Seyyedeh Neda Kazemi⁶, Bahram Rashidkhani⁷, Mehran Nouri^{8*} and Ghazaleh Eslamian^{9*}

Abstract

Introduction The present aimed to examine the relationship between Global Diet Quality (GDQ) and Prime Diet Quality (PDQ) scores and the likelihood of bacterial vaginosis (BV) among women.

Methods This case-control study was conducted among patients referred to a gynecological clinic in Tehran using the convenience sampling method. All the participants were examined by a gynecologist to rule out BV based on the presence of three or four of Amsel criteria. A valid semi-quantitative food frequency questionnaire (FFQ) containing 168 food items was used to estimate participants' dietary intake. To calculate the GDQ score, 25 food groups were considered, while 21 food groups were used for the PDQ score, based on the of daily consumption (in grams). All statistical analysis were performed using SPSS, and the association between GDQ and PDQ scores and the odds of BV was evaluated using binary logistic regression.

Results After adjusting for age, energy intake, fat intake, BMI, physical activity, familial history of BV, pregnancy history, menstrual cycle, smoking history, and the number of sexual partners in the previous month, significant associations remained between highest tertile of GDQ total (odds ratio (OR) = 0.219, confidence interval (CI) 95%: 0.101-0.475) and positive score (OR = 0.235, CI 95%: 0.103-0.533), as well as PDQ total (OR = 0.277, CI 95%: 0.131-0.583) and healthy score (OR = 0.397, CI 95%: 0.185-0.854) with the odds of BV, compared to the first tertile.

Conclusion A high diet quality, as indicated by high GDQ and PDQ scores, was associated with decreased risk of BV. These findings suggest that dietary intervention may be a viable strategy for the prevention and management of BV.

Keywords Diet, Diet quality, Global diet, Prime diet, Bacterial vaginosis

*Correspondence: Mehran Nouri mehran_nouri71@yahoo.com Ghazaleh Eslamian gh_eslamian@yahoo.com

Full list of author information is available at the end of the article



© The Author(s) 2025. **Open Access** This article is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License, which permits any non-commercial use, sharing, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if you modified the licensed material. You do not have permission under this licence to share adapted material derived from this article or parts of it. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by-nc-nd/4.0/.

Introduction

Bacterial vaginosis (BV) often presents with vaginal discharge, odor, and discomfort in women of childbearing age [1]. It is associated with various gynecological and obstetric complications, including pelvic inflammatory disease, spontaneous abortion, preterm birth, early pregnancy loss in IVF, and an increased risk of acquiring and transmitting HIV/STIs [2–6]. The global prevalence of BV is significant, affecting a considerable number of individuals worldwide, with reported occurrence rate ranging from 23 to 29% in different regions [7].

The clinical presentation of BV often includes a vaginal pH greater than 4.5, a thin and homogeneous whitish discharge, the presence of "clue" cells, and an amine odor upon the application of 10% potassium hydroxide [8, 9]. Treating BV presents challenges, as the condition recurs in a significant number of women—approximately 60% after antibiotic treatment [10]. BV is characterized by a decline in the abundance of healthy vaginal bacteria and a concomitant overgrowth of pathogenic bacteria [11, 12]. While sexual activity is associated with imbalances in the vaginal flora, other nonsexual risk factors can also contribute to BV development [13].

There are evidences linking BV to nutritional status and diet [14, 15]. For instance, prior research has revealed an inverse association between the intake of certain micronutrients, such as vitamins, folate and calcium, and the risk of sever BV [16]. Another recent study found a positive association between adherences to an unhealthy dietary pattern and BV risk, while an inverse association was observed between adherences to an ovo-vegetarian dietary pattern and the odds of BV [17].

To assess dietary quality, the use of a dietary metric is necessary. The two dietary scores commonly used for this purpose are the Prime Diet Quality Score (PDQS) and the Global Diet Quality Score (GDQS). The PDQS, developed by Fung et al. [18], is a simple, food-based diet quality score that differentiate healthy foods from unhealthy ones based on two main criteria: (1) evidence from the literature regarding the relationship between food and the risk of non-communicable diseases, and (2) the global contribution of nutrients. The association between PDQS and several non-communicable diseases, such as dyslipidemia, hypertension, diabetes, and depression, has been previously investigated [19–21].

Additionally, the GDQS was designed as an easy-touse cost-effective tool to monitor nutritional deficiencies and assess the risks of non-communicable diseases based on diet [22]. According to the GDQS, 25 food groups are categorized into three groups: (1) "unhealthy foods," which lower the overall diet quality score; (2) "healthy foods," which enhance the overall diet quality score; and (3) two food groups classified as "unhealthy in excessive amounts," which increase the diet quality score when consumed in moderation but lower it when consumed in excess [23].

Previous research has suggested that a high-quality diet, characterized by high scores in both the Prime and Global Diet Quality Scores, is associated with a decreased likelihood of developing several health-related conditions [24, 25]. However, no research has examined the correlation between diet quality, as assessed through GDQ and PDQ scores, and the risk of BV.

Therefore, the present study was designed to assess the relationship between GDQ and PDQ scores and the probability of BV among women. This study may provide important insight into the role of dietary quality in the BV risk and contribute to improving reproductive health among woman.

Methods and materials

Sample size

The sample size of the present study was calculated based on dietary pattern exposure using the case-control study formula by Fahim et al. [26]. Previous data indicated that 73% of the Iranian population adheres to unhealthy diets [27]. Assuming an odds ratio (OR) 2.5 for BV in women with unhealthy dietary patterns compared to those who do not adhere, and to achieve 80% statistical power with a 5% alpha error [26], we recruited 151 women with BV for the case group and 143 healthy women for the control group.

This study was conducted in accordance with the ethical standards of the Declaration of Helsinki and was approved by the Ethics Committee of Shahid Beheshti University of Medical Sciences (IR.SBMU.NNFTRI. REC.1399.054). Informed consent was obtained from all participants after they had read and signed the consent form. Some details of this study have been previously published [17, 28, 29].

Study population

We conducted this case-control study among patients referred to the gynecological clinic of Imam Hossein hospital in Tehran, Iran, using the convenience sampling method. All participants were examined by a gynecologist to diagnose or rule out BV based on the presence of three or four Amsel criteria, which include: (1) a homogenous and diluted vaginal discharge, (2) a vaginal pH greater than 4.5, (3) the presence of at least 20% clue cells during saline microscopy, and (4) a fishy odor upon adding 10% potassium hydroxide to the discharge slide [8].

Eligible participants met the following inclusion criteria: willingness to participate in the study and signing the consent form; an age range of 15–45 years; not being pregnant or menopausal; not using antibiotics, probiotics, hormonal contraceptives, vaginal douches, or immunosuppressive medications; and not suffering from systemic illnesses, immune system diseases, chronic infections, diet-related chronic diseases (such as diabetes and cardiovascular disease), or any uterine cavity abnormalities (such as fibroids and polyps), as well as no history of hysterectomy. The only differing inclusion criteria were a diagnosis of BV for the case group and the absence of ongoing or previous BV or BV treatment for the control group.

Participants in both groups were excluded if they failed to complete 60% or more of the food frequency questionnaire (FFQ), if their reported energy intake deviated by more than ± 3 standard deviations (SD) from the average energy intake, or if they were unwilling to continue participation.

A socio-demographic questionnaire was used to collect information on participants' age, family history of BV, education level, occupational status, smoking habits, and number of sex partners. Questions about alcohol and opium were excluded due to the specific religious and cultural beliefs of Iranians.

For anthropometric assessments, a trained examiner measured participants' weight while wearing light clothing a reliable scale with a precision of 100 g. Height in a straight standing position without shoes, with an accuracy of 1 mm. Waist circumference (WC) was assessed to determine central adiposity using a non-stretched tape, measured to the nearest 1 mm. Body Mass Index (BMI) was calculated by dividing weight (kg) by the square of height (meter).

Dietary intake assessment

A valid semi-quantitative food frequency questionnaire (FFQ) containing of 168 food items [30] was used to estimate participants' dietary intake during the year prior to the interview. Each item had a standard and commonly used serving size for Iranians, following the Willet method [31].

Global diet quality score (GDQS)

For the calculation of GDQS index, daily consumption (in grams) of food items was categorized into 25 food groups. These included 16 healthy food groups (fish and shellfish, poultry and game meats, eggs, low fat dairy, whole grains, cruciferous vegetables, dark green leafy vegetables, deep orange vegetables, other vegetables, deep orange fruits, citrus fruits, other fruits, deep orange tubers, nuts and seeds, legumes, and liquid oils), two optimal food groups (red meats and high fat dairy), and seven unhealthy food groups (refined grains and baked goods, white roots and tubers, juices, sugar-sweetened beverages, sweets and ice creams, fried foods, and processed meats). Each food group was categorized into three or four intake levels. For the healthy food groups, the scoring system assigned:

- 0 points for lower intake of all 16 food groups.
- 0.25 points for moderate intake and 0.5 points for higher intake of cruciferous vegetables, deep orange vegetables, other vegetables, and deep orange tubers.
- 1 point for moderate intake and 2 points for higher intake of citrus fruits, deep orange fruits, other fruits, whole grains, liquid oils, fish and shellfish, poultry and game meats, and low-fat dairy.
- 2 points for moderate intake and 4 points for higher intake eggs, dark green leafy vegetables, deep orange vegetables, nuts and seeds, legumes.

For the optimal food groups:

- 0 points were assigned for lower or very high intake (gram per day).
- 1 point was assigned for moderate intake.
- 2 points were assigned for higher intake of red meats and high-fat dairy.

For the unhealthy food groups:

- 2 points were given for lower intake.
- 1 point for moderate intake.
- 0 points for higher intake of refined grains and baked goods, white roots and tubers, juices, sugarsweetened beverages, sweets and ice creams, fried foods, and processed meats.

Finally, the total GDQS score was obtained by summing the scores of all 25 food groups, yielding a range from 0 to 49 [32].

Prime diet quality score (PDQS)

The PDQS contains 21 food groups, categorized into two groups: healthy and unhealthy. Initially, all 21 food groups were classified into tertiles. For healthy food groups—low fat dairy, poultry, whole grains, fish and shellfish, legumes and soy, nuts and seeds, liquid vegetable oils, deep orange fruits, citrus fruits, other fruits, deep orange vegetables, cruciferous vegetables, dark green leafy vegetables, and other vegetables—points were assigned as follow: 0 for lower intake, 1 for moderate intake, and 2 for higher intake.

Conversely, for healthy food groups—low fat dairy, poultry, whole grains, fish and shellfish, legumes and soy, nuts and seeds, liquid vegetable oils, deep orange fruits, citrus fruits, other fruits, deep orange vegetables, cruciferous vegetables, dark green leafy vegetables, and other vegetables—points were assigned in reverse: 2 for lower intake, 1 for moderate intake, and 0 for higher intake. The PDQS comprises 21 food groups, with a total ranging from 0 to 42 total [33].

Statistical analysis

All statistical analysis were preformed using SPSS (Statistical Package for the Social Sciences program; version 23; Chicago, IL, United States). The Chi-square was used to compare categorical variables, while Mann-Whitney U test and independent sample t-test were employed to compare non-parametric and parametric variables between the case and control groups.

Median (confidence interval (CI) 25th -75th) or mean \pm SD were used for continues variables, and percentage were reported for categorical variables. To evaluate the association between GDQS and PDQS with the odds of BV, binary logistic regression was conducted using two models: a crude model and an adjusted model. The OR with a CI 95% was calculated.

In the adjusted model, potential confounders—including age (years), energy intake (kcal/day), fat intake (g/ day), BMI (kg/m²), physical activity (MET/h/day), familial history of BV (yes/no), pregnancy history (yes/no), menstrual cycle (yes/no), smoking history (yes/no), and number of sexual partners in previous month—were controlled.

Results

The baseline characteristics of the study population between the healthy and bacterial vaginosis groups are presented in Table 1. The median age (p-value = 0.019), the percentage of familial history of BV (p-value < 0.001), and smoking (p-value < 0.001) were significantly different

between the two groups. Additionally, the median intake of total fat was significantly higher in the case group (p-value = 0.022), while fiber intake was higher in the control group (p-value = 0.013).

The food group intake of the study population between the healthy and bacterial vaginosis groups is shown in Table 2. The mean total GDQ and PDQ scores, as well as their components, were significantly higher in the control group compared to the case group (p-value < 0.001 for all, except for healthy and unhealthy PDQ scores).

The median intake of other fruits (p-value = 0.028), deep dark leafy vegetables (p-value < 0.001), other vegetables (p-value = 0.006), deep orange tubers (p-value = 0.018), legumes (p-value < 0.001), and poultry and game meats (p-value = 0.034) were significantly higher in the control group. However, the median intake of refined grains and baked goods (p-value = 0.004), processed meats (p-value = 0.008), sugar-sweetened beverages (p-value < 0.001), and fried foods (p-value = 0.003) was higher in the case group.

The association between tertiles of Global and Prime Diet Quality Scores and the odds of bacterial vaginosis is presented in Table 3, based on two models: crude and adjusted.

In the crude model, compared to the first tertile, lower odds of BV were observed in the highest tertile of the total GDQ score total (OR = 0.246, CI 95%: 0.136–0.444), positive GDQ score total (OR = 0.367, CI 95%: 0.205–0.658), negative GDQ score total (OR = 0.432, CI 95%: 0.249–0.750), total PDQ score total (OR = 0.237, CI 95%: 0.128–0.437), healthy PDQ score total (OR = 0.457,

Table 1 Baseline characteristics and nutrients intake in the study population: comparison between healthy and bacterial vaginosis groups

Variables	Case (n = 143)	Control (<i>n</i> = 151)	P-value
Baseline characteristics			
Age (year) ¹	30.0 (25.0-33.5)	32.0 (24.0-37.0)	0.201
BMI (kg/m ²) ¹	26.1 (23.3–28.7)	24.6 (22.2–27.9)	0.019
Physical activity (MET/h/day) ¹	25.0 (15.0–50.0)	25.0 (15.0–60.0)	1.000
Familial history of BV, yes, % ²	77 (53.8)	37 (24.5)	<0.001
Pregnancy history, yes, % ²	80 (55.9)	81 (53.6)	0.726
Menstrual cycle, irregular, % ²	49 (34.3)	49 (32.5)	0.805
Smoking history, yes, % ²	25 (17.5)	2 (1.3)	<0.001
Number of sexual partners in previous month, more than 2, % 2	6 (4.5)	4 (3.0)	0.800
Dietary intakes			
Energy (kcal/day) ¹	2481.12 (1718.84-3274.81)	2223.33 (1761.26-3042.55)	0.306
Protein (g/day) ¹	76.16 (58.96–99.12)	80.75 (59.77-110.86)	0.234
Total fat (g/day) ¹	99.91 (68.04-124.86)	83.46 (62.14-107.95)	0.022
Carbohydrate (g/day) ¹	325.73 (216.73-449.23)	307.47 (217.77-412.20)	0.463
Fiber (g/day) ¹	20.09 (14.50-27.05)	23.75 (15.83–37.27)	0.013

Abbreviation: BMI, body mass index; kg, kilogram; m, meter; MET, metabolic equivalent of task; kcal, kilocalorie; g, gram; BV, bacterial vaginosis, kcal, kilocalorie; g, gram

¹Using the Mann-Whitney U-test, values are presented as median (25th -75th)

²Using chi-square tests for categorical variables, values are presented as percentage

Table 2 Food group	intake in the study	/ population: com	nparison between	healthy and bacterial	vaginosis groups

Variables	Case (n = 143)	Control (<i>n</i> = 151)	P-value
Positive GDQ score ¹	14.37±4.51	16.72±5.41	<0.001
Negative GDQ score ¹	9.64±1.82	10.47 ± 1.86	<0.001
Total GDQ score ¹	24.02 ± 5.08	27.19±5.37	<0.001
Healthy PDQ score ¹	12.92±4.63	14.86 ± 5.94	0.002
Unhealthy PDQ score ¹	8.59 ± 3.14	12.92 ± 4.63	0.003
Total PDQ score ¹	20.38±5.11	23.45 ± 6.04	<0.001
Citrus fruits (g/day) ²	60.00 (28.81-104.78)	53.83 (27.08-116.75)	0.795
Deep orange fruits (g/day) 2	72.69 (36.93–110.90)	74.25 (44.40-158.45)	0.245
Other fruits (g/day) ²	56.10 (29.79-109.48)	83.86 (41.18-158.87)	0.028
Deep dark and leafy vegetables (g/day) ²	25.36 (12.57–45.46)	45.22 (22.20-82.73)	<0.001
Cruciferous vegetables (g/day) ²	3.10 (0.75–6.20)	3.92 (0.76–12.61)	0.103
Deep orange vegetables (g/day) ²	3.69 (1.35-11.00)	2.67 (0.90-9.04)	0.227
Other vegetables (g/day) ²	156.42 (109.92-236.62)	203.88 (118.60-308.70)	0.006
Deep orange tubers (g/day) ²	4.86 (1.21–10.52)	7.30 (2.43–14.07)	0.018
Legumes (g/day) ²	18.64 (7.41–34.24)	29.79 (12.09-70.64)	<0.001
Nuts and seeds (g/day) ²	5.97 (2.01–14.55)	5.74 (1.51–14.28)	0.369
Whole grains (g/day) ²	2.14 (0.41-4.00)	1.06 (0.16-5.21)	0.984
Refined grains and backed goods (g/day) 2	421.00 (288.15-605.94)	337.04 (240.20-474.83)	0.004
White roots and tubers (g/day) ²	14.25 (5.70-36.64)	24.42 (5.70-36.64)	0.391
Liquid oils (g/day) ²	11.50 (4.28-18.00)	12.00 (6.00–18.00)	0.508
Red meats (g/day) 2	44.58 (19.15–64.31)	42.66 (23.31-82.85)	0.113
Processed meats (g/day) ²	2.01 (0.30-5.78)	0.71 (0.00-4.34)	0.008
Fish and selfish (g/day) 2	2.95 (0.60-8.66)	5.51 (1.52–14.38)	0.004
Poultry and game meats (g/day) 2	12.85 (8.57–25.71)	17.14 (8.57-30.00)	0.034
Eggs (g/day) ²	22.92 (7.64–38.21)	22.92 (15.28-38.21)	0.108
Low fat dairy products (g/day) ²	224.25 (87.72-369.33)	246.87 (77.34-446.42)	0.563
High fat dairy products (g/day) ²	58.33 (21.42-145.52)	48.60 (19.33-134.95)	0.159
Sweets and ice creams (g/day) ²	49.31 (29.17–68.35)	38.42 (21.36-71.00)	0.067
Sugar-sweetened beverages (g/day) ²	24.50 (7.41-176.39)	8.16 (0.69-35.00)	<0.001
Juices (g/day) ²	2.71 (0.00-10.28)	3.39 (0.00-16.53)	0.410
Fried foods (g/day) ²	14.00 (6.00-30.00)	8.90 (3.00-21.14)	0.003

Abbreviation: GDQ, global diet quality; PDQ, prime diet quality; g, gram

 1 Using an independent sample t-test, values are presented as mean \pm SD

²Using the Mann-Whitney U-test, values are presented as median (25th -75th)

CI 95%: 0.261–0.801) and healthy PDQ score total (OR = 0.509, CI 95%: 0.289–0.897).

After adjusting for age, energy intake, fat intake, BMI, physical activity, familial history of BV, pregnancy history, menstrual cycle, smoking history, and number of sexual partners in previous month, the associations remained significand for the highest tertile of the total GDQ score (OR = 0.219, CI 95%: 0.101–0.475), positive GDQ score (OR = 0.235, CI 95%: 0.103–0.533), total PDQ score (OR = 0.277, CI 95%: 0.131–0.583), and healthy PDQ score (OR = 0.397, CI 95%: 0.185–0.854) with the odds of BV, compared to the first tertile.

Discussion

The result of the current study demonstrated that the last tertile of the GDQ total score, positive GDQ score, PDQ total score, and healthy PDQ score was associated lower odds of BV compared to the first tertile. Additionally, the total GDQ and PDQ score, along with their components, were significantly higher in the control group than in the BV group.

Healthy and unhealthy food groups of PDQ and GDQ scores may explain this association. The consumption of certain food groups, including fruits, vegetables, whole grains, and legumes, is included in the positive GDQ score and a healthy PDQ score, where higher intake is assigned higher points. Increased consumption of these food groups is associated with greater fiber intake. The current study showed that fiber intake was higher in the control group compared to BV subjects.

An earlier study indicated that women with a higher fiber intake had a lower risk of molecular BV [34]. Additionally, it was demonstrated that a plant-based diet rich in high-fiber foods such as legumes, fruits and vegetables reduced the odds of BV [17]. The ecology of vaginal microbiota, particularly a high proportion of

Variables	T ₁	T ₂	T ₃	P _{trend}
		OR (CI 95%)	OR (CI 95%)	
GDQ score (case/control)	63/35	49/46	31/70	
Crude	Ref.	0.592 (0.332-1.054)	0.246 (0.136–0.444)	<0.001
Adjusted	Ref.	0.497 (0.240-1.029)	0.219 (0.101–0.475)	<0.001
Positive GDQ score (case/control)	58/40	52/49	33/62	
Crude	Ref.	0.732 (0.418-1.282)	0.367 (0.205–0.658)	0.001
Adjusted	Ref.	0.493 (0.238-1.018)	0.235 (0.103–0.533)	0.001
Negative GDQ score (case/control)	53/38	43/35	47/78	
Crude	Ref.	0.881 (0.478-1.622)	0.432 (0.249–0.750)	0.002
Adjusted	Ref.	1.186 (0.560–2.513)	0.550 (0.252-1.202)	0.104
PDQ score (case/control)	61/33	54/54	28/64	
Crude	Ref.	0.541 (0.307–0.954)	0.237 (0.128–0.437)	<0.001
Adjusted	Ref.	0.601 (0.300-1.202)	0.277 (0.131–0.583)	0.001
Healthy PDQ score (case/control)	56/49	52/35	35/67	
Crude	Ref.	1.300 (0.732-2.310)	0.457 (0.261–0.801)	0.007
Adjusted	Ref.	1.213 (0.585–2.515)	0.397 (0.185–0.854)	0.014
Unhealthy PDQ score (case/control)	56/41	46/51	41/59	
Crude	Ref.	0.660 (0.375-1.164)	0.509 (0.289–0.897)	0.02
Adjusted	Ref.	0.920 (0.438-1.932)	0.673 (0.276-1.644)	0.383

Table 3 Association between tertiles of global and prime diet quality score and the odds of bacterial vaginosis

Abbreviation: OR, odds ratio; CI, confidence interval; T, tertile; BV, bacterial vaginosis; GDQ, global diet quality, PDQ, prime diet quality

Significant values are presented in bold

Values are expressed as odds ratio (95% Cls)

Obtained from logistic regression analysis

Adjusted model: adjusted for age (years), energy intake (kcal/day), fat intake (g/day), BMI (kg/m²), physical activity (MET/h/day), familial history of BV (yes/no), pregnancy history (yes/no), menstrual cycle (yes/no), smoking history (yes/no), and number of sexual partners in previous month

Lactobacillus species, decreases vulnerability to BV and inhibits pathogen invasion [35]. The *Lactobacillus* genus, the predominant bacteria found in the human vaginal microbiome, creates an acidic environment that is believed to protect women against sexually transmitted pathogens and risk of infections [36].

A Diet rich in fiber may lower the risk of bacterial infections linked to BV by impacting the vaginal microflora, increasing the abundance of *Lactobacillus* bacteria, and enhancing overall vaginal health [34]. Moreover, an in vitro study revealed that prebiotic dietary fibers promoted the proliferation of primary *Lactobacillus* species monocultures [37]. Previous studies have shown a certain level of agreement between the types of *Lactobacillus* species found in both the rectum and the vagina [38, 39]. The gut microbiota can influence the vaginal microbiome [40], and change in gut microbiota composition may contribute to BV development.

A study among BV patients demonstrated that oral intake of a mixture of three *Lactobacillus* species prolonged the time to recurrence and lowered the percentage of BV recurrences in women who had recently received treatment [41]. Additionally, a meta-analysis revealed that probiotics used as an adjuvant therapy alongside antibiotics were more effective than antibiotic treatment alone in BV patients. Furthermore, a high dose of probiotics administered orally, particularly *Lactobacillus*

rhamnosus, was found to be more effective than a low dose or vaginal administration of *Lactobacillus rhamnosus* for BV treatment [42].

Growing evidence suggests that the vaginal microbiome can be influenced by gut microbiota, as bacterial strains can transfer from the gut to the vagina [40]. Vaginal dysbiosis can be improved through probiotics via several mechanisms, including the production of lactic acid, which helps maintain normal vaginal pH. A study using a yogurt drink containing *Lactobacillus* strains found that vaginal pH decreased in the intervention group compared to the placebo group after four weeks of treatment [43]. Additionally, probiotics help reduce pathogenic organisms by competing for adhesion sites [44, 45] and promote immune defense by increasing anti-inflammatory cytokines while decreasing inflammatory cytokines [46].

Moreover, short-chain fatty acids (SCFAs) are produced through the fermentation of dietary non-digestible carbohydrates by gut microbiota. A deficiency in SCFAproducing bacteria and their metabolites—such as acetate, propionate, and butyrate—can lead to the leakage of endotoxins like lipopolysaccharides into systemic circulation (leaky gut), resulting in systemic inflammation. This inflammation or the transfer of these pathogens to the genital tract increases the risk of genital tract infections and inflammation [40].

Page 7 of 10

In addition to their fiber content, fruit and vegetables may play a protective role against BV due to their other beneficial components. The current study showed that the consumption of certain fruits and vegetables was higher in the control compared to the case group.

Fruits and vegetables—particularly green leafy vegetables, whole grains, and eggs—are important dietary sources of folate [47]. A previous study found a negative association between dietary folate intake and BV risk [48]. It is possible that folate consumption enhance the immune function, thereby reducing the likelihood of severe BV [48].

Folic acid has been shown to sustain or improve natural killer (NK) cell activity, which strengthens innate, non-specific immune function, thereby improving immunity and lowering the prevalence of BV [49]. Additionally, folic acid plays a key role in antibody synthesis and supports the Th1-mediated immune response, which stimulates the body's adaptive immune system and further reduces the risk of BV [49].

Another micronutrient that may explain the association between higher GDQ and PDQ scores and lower odds of BV is vitamin E. This vitamin is found in various dietary sources, including vegetable oils, as well as fresh fruits, vegetables, and nuts. A previous study reported a correlation between increased vitamin E consumption and a reduced likelihood of severe BV [48].

The role of vitamin E as a potent antioxidant is well established. Excessive production of reactive oxygen species (ROS) beyond the cellular antioxidant scavenging capacity can contribute to BV development [50]. As a fat-soluble antioxidant, vitamin E helps neutralize excess free radicals and regulates gene expression in the immune system, potentially reducing the risk of BV [51].

Moreover, women with insufficient levels of micronutrients such as beta-carotene and vitamins D, C, A, and E are at a higher risk of developing BV. This inadequacy in essential nutrients may be linked to a diet low in fruits and vegetables.

Additionally, flavonoid-rich foods—such as vegetables, fruits, nuts, and legumes—have been found to possess anti-inflammatory properties that can effectively inhibit the activation of nuclear factor kappa-light-chainenhancer of activated B cells (NF- κ B) [52]. NF- κ B plays a crucial role in pro-inflammatory signaling pathways, and its activation has been observed in various cell types present in the vaginal secretions of women with BV [53].

Furthermore, previous studies have shown an association between vitamin D deficiency and increased odds of BV [54]. Adequate vitamin D levels have been demonstrated to protect against BV by promoting the synthesis of cathelicidins, antimicrobial peptides found in the lysosomes of neutrophils and macrophages [55]. A recent study also found that higher serum levels of carotenoids were associated with a lower prevalence of BV [56]. As strong antioxidants, carotenoids help reduce the accumulation of reactive oxygen species. Additionally, sufficient carotenoid and vitamin A levels have been shown to be essential for maintaining the integrity of vaginal epithelial cells [57]. Moreover, carotenoids enhance the activity of neutrophils, natural killer cells, and other innate immune cells, further supporting immune function [58, 59].

On the other hand, a lower intake of certain food groups—such as red meat, processed foods, refined grains, fried foods, and sugar-sweetened beverages (SSBs)—along with higher GDQ and healthy PDQ scores may explain the observed association. The result of the current study showed that the intake of refined grains, processed meat, SSBs, and fried food was higher in BV subjects compared to control individuals.

The excessive fat content in processed and fried foods has been identified as a potential contributing factor to the increased odds of BV. Naggers et al. provided evidence suggesting that a diet high in saturated fat may increase the likelihood of severe BV [60]. Consuming excessive amounts of fat can lead to an elevated vaginal pH by altering the vaginal microflora, thereby increasing the risk of BV development [48, 60]. Furthermore, a high intake of saturated fats may negatively impact the effectiveness of the mucosal immune response, further contributing to BV susceptibility [60].

Another key characteristic of higher GDQ and PDQ scores is a lower intake of high-glycemic-index and gly-cemic-load-low-carbohydrates, including refined grains, SSBs, sweets, and ice cream. Previous research has established a link between the consumption of ultra-processed sweets, dietary glycemic index (GI), and dietary glycemic load (GL) with an increases risk of developing BV [14, 61].

Excessive consumption of high-GI/GL foods, as part of an unhealthy diet, may contribute to BV development by inducing oxidative stress and compromising the immune system's response [38, 62].

Although the primary goal of the current study was to investigate the association between GDQ and PDQ scores and the odds of BV, the findings have potential implications for public health and clinical practice. Given the high prevalence of BV among women and its substantial economic burden [7], implementing dietary interventions to modify eating patterns may be a valuable strategy for preventing and managing the disease.

If future studies confirm these findings, the role of diet quality in the onset and progression of BV could be emphasized in public health education to raise awareness among women and help reduce BV risk. Based on GDQ and PDQ scores, increasing the consumption of high-fiber foods—such as fruits, vegetables, whole grains, and legumes—along with antioxidant-rich foods, while reducing the intake of sugar and processed foods, may help lower the odds of BV among women.

his study has several strengths. One notable advantage is the use of a comprehensive and globally recognized diet quality score, which provides valuable insights into the potential impact of overall diet quality on BV risk. Additionally, accounting for potential confounding factors—such as age, energy intake, fat intake, BMI, physical activity, and family history of bacterial vaginosis—enhances the reliability of the findings. Furthermore, to minimize the risk of information bias, a qualified nutritionist administered the questionnaires without prior knowledge of the study outcomes.

Nevertheless, this study has several limitations that should be considered. One major limitation is its casecontrol design, which inherently restricts the ability to establish causality. While our findings indicate associations, we cannot conclude that one factor directly leads to another. Establishing causality would require alternative research designs, such as randomized controlled trials.

Additionally, despite efforts to minimize bias, casecontrol studies are susceptible to selection bias, measurement bias, and recall bias, which may affect the accuracy of the results. Moreover, diet quality was assessed using a FFQ, a method prone to recall bias since participants report their dietary intake retrospectively.

Another limitation is the use of convenience sampling, which may affect the generalizability of our findings. Since participants were not selected randomly, the sample may not fully represent women with BV [63].

It is also important to consider the long-term impact of diet quality on BV recurrence. Conducting studies with longer follow-up periods could provide more valuable insights into the relationship between diet and BV risk and recurrence. Furthermore, future research should explore the correlation between dietary quality indices and the composition of vaginal microbiota. Such studies could offer a deeper understanding of the potential impact of diet on the vaginal microbial community.

Conclusion

A high diet quality, as indicated by high GDQ and PDQ scores, was associated with a decreased risk of BV. These findings suggest that dietary interventions may serve as an effective strategy for the prevention and management of BV.

Acknowledgements

We sincerely thank all field investigators, staff, and participants of the present study.

Author contributions

S.M., M.H.M., M.N., M.S., S.E. and M.N.; Contributed to writing the first draft. M.N. and S.N.K.: Contributed to data collecting. B.R. and M.N.; Contributed to all data and statistical analysis, and interpretation of data. M.N. and G.E.; Contributed the research concept, supervised the work and revised the manuscript. All authors read and approved of the final manuscript.

Funding

Not applicable.

Data availability

No datasets were generated or analysed during the current study.

Declarations

Ethics approval and consent to participate

This study was conducted in accordance with the ethical standards of the declaration of Helsinki and was approved by the Ethics Committee of Shahid Beheshti University of Medical Sciences (IR.SBMU.NNFTRI.REC.1399.054). All participants read and signed the informed consent form.

Consent for publication

Not applicable.

Competing interests

The authors declare no competing interests.

Author details

¹Nutrition and Food Security Research Center, Isfahan University of Medical Sciences, Isfahan, Iran
²Department of Midwifery, Faculty of Nursing and Midwifery, Ahvaz Branch, Islamic Azad University, Ahvaz, Iran

³Preventative Gynecology Research Center, Imam Hossein Hospital, Shahid Beheshti University of Medical Sciences, Tehran, Iran

⁴Department of Nutrition, School of Public Health, Iran University of Medical Sciences, Tehran, Iran

⁵Nutrition and Metabolic Diseases Research Center and Clinical Sciences Research Institute, Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran

⁶Department of Obstetrics and Gynecology, School of Medicine, Preventative Gynecology Research Center, Imam Hossein Hospital, Shahid Beheshti University of Medical Sciences, Tehran, Iran

⁷Department of Community Nutrition, Faculty of Nutrition and Food Technology, National Nutrition and Food Technology Research Institute, Shahid Beheshti University of Medical Sciences, Tehran, Iran ⁸Infertility and Reproductive Health Research Center, Health Research

Institute, Babol University of Medical Sciences, Babol, Iran ⁹Department of Cellular and Molecular Nutrition, Faculty of Nutrition and Food Technology, National Nutrition and Food Technology Research

Institute, Shahid Beheshti University of Medical Sciences, Tehran, Iran

Received: 16 January 2025 / Accepted: 21 February 2025 Published online: 28 February 2025

References

- Bagnall P, Rizzolo D. Bacterial vaginosis: A practical review. JAAPA: Official J Am Acad Physician Assistants. 2017;30(12):15–21.
- Işik G, Demirezen Ş, Dönmez HG, Beksaç MS. Bacterial vaginosis in association with spontaneous abortion and recurrent pregnancy losses. J Cytol. 2016;33(3):135–40.
- Mohanty T, Doke PP, Khuroo SR. Effect of bacterial vaginosis on preterm birth: a meta-analysis. Arch Gynecol Obstet. 2023;308(4):1247–55.
- Eckert LO, Moore DE, Patton DL, Agnew KJ, Eschenbach DA. Relationship of vaginal bacteria and inflammation with conception and early pregnancy loss following in-vitro fertilization. Infect Dis Obstet Gynecol. 2003;11(1):11–7.
- Atashili J, Poole C, Ndumbe PM, Adimora AA, Smith JS. Bacterial vaginosis and HIV acquisition: a meta-analysis of published studies. AIDS. 2008;22(12):1493–501.

- Taylor BD, Darville T, Haggerty CL. Does bacterial vaginosis cause pelvic inflammatory disease? Sex Transm Dis. 2013;40(2):117–22.
- Peebles K, Velloza J, Balkus JE, McClelland RS, Barnabas RV. High global burden and costs of bacterial vaginosis: A systematic review and Meta-Analysis. Sex Transm Dis. 2019;46(5):304–11.
- Amsel R, Totten PA, Spiegel CA, Chen KC, Eschenbach D, Holmes KK. Nonspecific vaginitis: diagnostic criteria and microbial and epidemiologic associations. Am J Med. 1983;74(1):14–22.
- 9. Wang J. Bacterial vaginosis. Prim Care Update OB/GYNS. 2000;7(5):181–5.
- Bradshaw CS, Morton AN, Hocking J, Garland SM, Morris MB, Moss LM, Horvath LB, Kuzevska I, Fairley CK. High recurrence rates of bacterial vaginosis over the course of 12 months after oral metronidazole therapy and factors associated with recurrence. J Infect Dis. 2006;193(11):1478–86.
- 11. O'Hanlon DE, Moench TR, Cone RA. Vaginal pH and microbicidal lactic acid when lactobacilli dominate the microbiota. PLoS ONE. 2013;8(11):e80074.
- Stoyancheva G, Marzotto M, Dellaglio F, Torriani S. Bacteriocin production and gene sequencing analysis from vaginal Lactobacillus strains. Arch Microbiol. 2014;196:645–53.
- Thoma ME, Klebanoff MA, Rovner AJ, Nansel TR, Neggers Y, Andrews WW, Schwebke JR. Bacterial vaginosis is associated with variation in dietary indices. J Nutr. 2011;141(9):1698–704.
- Noormohammadi M, Eslamian G, Kazemi SN, Rashidkhani B, Omidifar F. Association between consumption of ultra-processed foods and bacterial vaginosis: a case-control study. Iran J Obstet Gynecol Infertility. 2022;24(12):67–76.
- Noormohammadi M, Eslamian G, Kazemi SN, Rashidkhani B. Dietary acid load, alternative healthy eating index score, and bacterial vaginosis: is there any association? A case-control study. BMC Infect Dis. 2022;22(1):803.
- Neggers YH, Nansel TR, Andrews WW, Schwebke JR, Yu KF, Goldenberg RL, Klebanoff MA. Dietary intake of selected nutrients affects bacterial vaginosis in women. J Nutr. 2007;137(9):2128–33.
- Noormohammadi M, Eslamian G, Kazemi SN, Rashidkhani B. Association between dietary patterns and bacterial vaginosis: a case-control study. Sci Rep. 2022;12(1):12199.
- Fung TT, Isanaka S, Hu FB, Willett WC. International food group–based diet quality and risk of coronary heart disease in men and women. Am J Clin Nutr. 2018;107(1):120–9.
- Cano-Ibáñez N, Serra-Majem L, Martín-Peláez S, Martínez-González MÁ, Salas-Salvadó J, Piquer MDC, Lassale C, Hernandez JAM, Alonso-Gómez ÁM, Wärnberg J. Association between the prime diet quality score and depressive symptoms in a mediterranean population with metabolic syndrome. Crosssectional and 2-year follow-up assessment from PREDIMED-PLUS study. Br J Nutr. 2022;128(6):1170–9.
- Alvarez-Alvarez I, Toledo E, Lecea O, Salas-Salvadó J, Corella D, Buil-Cosiales P, Zomeño MD, Vioque J, Martinez JA, Konieczna J. Adherence to a priori dietary indexes and baseline prevalence of cardiovascular risk factors in the PREDIMED-Plus randomised trial. Eur J Nutr. 2020;59:1219–32.
- Gicevic S, Gaskins AJ, Fung TT, Rosner B, Tobias DK, Isanaka S, Willett WC. Evaluating pre-pregnancy dietary diversity vs. dietary quality scores as predictors of gestational diabetes and hypertensive disorders of pregnancy. PLoS ONE. 2018;13(4):e0195103.
- Nguyen PH, Tran LM, Hoang NT, Deitchler M, Moursi M, Bergeron G. The global diet quality score is associated with nutrient adequacy and depression among Vietnamese youths. Ann N Y Acad Sci. 2023;1528(1):48–57.
- Norde MM, Bromage S, Marchioni DML, Vasques AC, Deitchler M, Arsenaut J, de Carvalho AM, Velloso L, Willett W, Giovannucci E, Geloneze B. The global diet quality score as an indicator of adequate nutrient intake and dietary quality– a nation-wide representative study. Nutr J. 2024;23(1):42.
- Fung TT, Li Y, Bhupathiraju SN, Bromage S, Batis C, Holmes MD, Stampfer M, Hu FB, Deitchler M, Willett WC. Higher global diet quality score is inversely associated with risk of type 2 diabetes in US women. J Nutr. 2021;151:S168–75.
- 25. Cano-Ibáñez N, Serra-Majem L, Martín-Peláez S, Martínez-González MÁ, Salas-Salvadó J, Corella Piquer MD, Lassale C, Martínez Hernandez JA, Alonso-Gómez ÁM, Wärnberg J, et al. Association between the prime diet quality score and depressive symptoms in a mediterranean population with metabolic syndrome. Cross-sectional and 2-year follow-up assessment from PREDIMED-PLUS study. Br J Nutr. 2022;128(6):1170–9.
- Fahim NK, Negida A, Fahim AK. Sample size calculation Guide Part 3: how to calculate the sample size for an independent Case-control study. Adv J Emerg Med. 2019;3(2):e20.
- 27. Azadbakht L, Mirmiran P, Hosseini F, Azizi F. Diet quality status of most Tehranian adults needs improvement. Asia Pac J Clin Nutr. 2005;14(2):163–8.

- 28. Noormohammadi M, Eslamian G, Kazemi SN, Rashidkhani B. Is there any association between adherence to the mediterranean diet and dietary total antioxidant capacity with bacterial vaginosis?? Results from a Case–Control study. BMC Womens Health. 2022;22(1):244.
- Khademian A, Noormohammadi M, Moori MH, Makhtoomi M, Esmaeilzadeh S, Nouri M, Eslamian G. The association between dietary phytochemical index and bacterial vaginosis risk: secondary analysis of case-control study. J Health Popul Nutr. 2024;43(1):135.
- Mirmiran P, Esfahani FH, Mehrabi Y, Hedayati M, Azizi F. Reliability and relative validity of an FFQ for nutrients in the Tehran lipid and glucose study. Public Health Nutr. 2010;13(5):654–62.
- 31. Willett W. Nutritional epidemiology, third Edn. New York: Oxford University Press; 2013.
- Bromage S, Batis C, Bhupathiraju SN, Fawzi WW, Fung TT, Li Y, Deitchler M, Angulo E, Birk N, Castellanos-Gutiérrez A, et al. Development and validation of a novel Food-Based global diet quality score (GDQS). J Nutr. 2021;151(12 Suppl 2):s75–92.
- Kronsteiner-Gicevic S, Mou Y, Bromage S, Fung TT, Willett W. Development of a diet quality screener for global use: evaluation in a sample of US women. J Acad Nutr Diet. 2021;121(5):854–e871856.
- Shivakoti R, Tuddenham S, Caulfield LE, Murphy C, Robinson C, Ravel J, Ghanem KG, Brotman RM. Dietary macronutrient intake and molecular-bacterial vaginosis: role of fiber. Clin Nutr. 2020;39(10):3066–71.
- Greenbaum S, Greenbaum G, Moran-Gilad J, Weintraub AY. Ecological dynamics of the vaginal Microbiome in relation to health and disease. Am J Obstet Gynecol. 2019;220(4):324–35.
- Miller EA, Beasley DE, Dunn RR, Archie EA. Lactobacilli Dominance and Vaginal pH: Why Is the Human Vaginal Microbiome Unique? Frontiers in microbiology 2016, 7:1936.
- Collins SL, McMillan A, Seney S, van der Veer C, Kort R, Sumarah MW, Reid G. Promising prebiotic candidate established by evaluation of lactitol, lactulose, Raffinose, and oligofructose for maintenance of a lactobacillus-dominated vaginal microbiota. Appl Environ Microbiol. 2018;84(5):e02200–02217.
- Antonio MA, Rabe LK, Hillier SL. Colonization of the rectum by Lactobacillus species and decreased risk of bacterial vaginosis. J Infect Dis. 2005;192(3):394–8.
- Petricevic L, Domig KJ, Nierscher FJ, Krondorfer I, Janitschek C, Kneifel W, Kiss H. Characterisation of the oral, vaginal and rectal Lactobacillus flora in healthy pregnant and postmenopausal women. Eur J Obstet Gynecol Reproductive Biology. 2012;160(1):93–9.
- 40. Amabebe E, Anumba DOC. Female gut and genital tract Microbiota-Induced crosstalk and differential effects of Short-Chain fatty acids on immune sequelae. Front Immunol. 2020;11:2184.
- Reznichenko H, Henyk N, Maliuk V, Khyzhnyak T, Tynna Y, Filipiuk I, Veresniuk N, Zubrytska L, Quintens J, Richir K, Gerasymov S. Oral intake of lactobacilli can be helpful in symptomatic bacterial vaginosis: A randomized clinical study. J Low Genit Tract Dis. 2020;24(3):284–9.
- 42. Chen R, Li R, Qing W, Zhang Y, Zhou Z, Hou Y, Shi Y, Zhou H, Chen M. Probiotics are a good choice for the treatment of bacterial vaginosis: a meta-analysis of randomized controlled trial. Reproductive Health. 2022;19(1):137.
- Laue C, Papazova E, Liesegang A, Pannenbeckers A, Arendarski P, Linnerth B, Domig K, Kneifel W, Petricevic L, Schrezenmeir J. Effect of a yoghurt drink containing Lactobacillus strains on bacterial vaginosis in women–a double-blind, randomised, controlled clinical pilot trial. Beneficial Microbes. 2018;9(1):35–50.
- Falagas M, Betsi GI, Athanasiou S. Probiotics for the treatment of women with bacterial vaginosis. Clin Microbiol Infection: Official Publication Eur Soc Clin Microbiol Infect Dis. 2007;13(7):657–64.
- 45. Mendling W. Vaginal microbiota. Adv Exp Med Biol. 2016;902:83–93.
- Lee JE, Lee S, Lee H, Song YM, Lee K, Han MJ, Sung J, Ko G. Association of the vaginal microbiota with human papillomavirus infection in a Korean twin cohort. PLoS ONE. 2013;8(5):e63514.
- Delchier N, Herbig AL, Rychlik M, Renard CM. Folates in fruits and vegetables: contents, processing, and stability. Compr Rev Food Sci Food Saf. 2016;15(3):506–28.
- Neggers YH, Nansel TR, Andrews WW, Schwebke JR, Yu K-f, Goldenberg RL, Klebanoff MA. Dietary intake of selected nutrients affects bacterial vaginosis in Women1,2,3. J Nutr. 2007;137(9):2128–33.
- 49. Maggini S, Pierre A, Calder PC. Immune Function and Micronutrient Requirements Change over the Life Course. Nutrients 2018, 10(10).

- Rimessi A, Previati M, Nigro F, Wieckowski MR, Pinton P. Mitochondrial reactive oxygen species and inflammation: molecular mechanisms, diseases and promising therapies. Int J Biochem Cell Biol. 2016;81 (Pt B):281–93.
- Liu EH, Liao WZ, Chen HK, Huang XY, Li RX, Liang HW, Guo XG. Association between serum vitamin E and bacterial vaginitis in women: a cross-sectional study. BMC Womens Health. 2024;24(1):316.
- 52. Serafini M, Peluso I, Raguzzini A. Flavonoids as anti-inflammatory agents. Proc Nutr Soc. 2010;69(3):273–8.
- Kalia N, Singh J, Kaur M. Immunopathology of recurrent vulvovaginal infections: new aspects and research directions. Front Immunol. 2019;10:458450.
- Mojtahedi SF, Mohammadzadeh A, Mohammadzadeh F, Jalili Shahri J, Bahri N. Association between bacterial vaginosis and 25-Hydroxy vitamin D: a casecontrol study. BMC Infect Dis. 2023;23(1):208.
- Bodnar LM, Krohn MA, Simhan HN. Maternal vitamin D deficiency is associated with bacterial vaginosis in the first trimester of pregnancy. J Nutr. 2009;139(6):1157–61.
- Tan M-Z, Feng Y-X, Hong D-Y, Guo X-G. Association between serum carotenoids and bacterial vaginosis infection among American women. BMC Infect Dis. 2024;24(1):20.
- Valenti P, Rosa L, Capobianco D, Lepanto MS, Schiavi E, Cutone A, Paesano R, Mastromarino P. Role of lactobacilli and lactoferrin in the mucosal cervicovaginal defense. Front Immunol. 2018;9:376.
- Terao R, Murata A, Sugamoto K, Watanabe T, Nagahama K, Nakahara K, Kondo T, Murakami N, Fukui K, Hattori H, Eto N. Immunostimulatory effect

of Kumquat (Fortunella crassifolia) and its constituents, β -cryptoxanthin and R-limonene. Food Funct. 2019;10(1):38–48.

- Milani A, Basirnejad M, Shahbazi S, Bolhassani A. Carotenoids: biochemistry, Pharmacology and treatment. Br J Pharmacol. 2017;174(11):1290–324.
- Neggers YH, Nansel TR, Andrews WW, Schwebke JR, Yu K-f, Goldenberg RL, Klebanoff MA. Dietary intake of selected nutrients affects bacterial vaginosis in women, 3. J Nutr. 2007;137(9):2128–33.
- Noormohammadi M, Eslamian G, Kazemi SN, Rashidkhani B, Malek S. Association of Dietary Glycemic Index, Glycemic Load, Insulin Index, and Insulin Load with Bacterial Vaginosis in Iranian Women: A Case-Control Study. Infectious diseases in obstetrics and gynecology 2022, 2022:1225544.
- Kawahito S, Kitahata H, Oshita S. Problems associated with glucose toxicity: role of hyperglycemia-induced oxidative stress. World J Gastroenterology: WJG. 2009;15(33):4137.
- 63. Etikan I, Musa SA, Alkassim RS. Comparison of convenience sampling and purposive sampling. Am J Theoretical Appl Stat. 2016;5(1):1–4.

Publisher's note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.