

Preplanned Studies

Menstrual Characteristics and the Risk of Spontaneous Abortion — 9 PLADs, China, 2013–2024

Tsomo Tenzin^{1,2,3,&}; Yumei Wei^{4,&}; Yiyao Jin^{1,2,3}; Jiaxin Li^{1,2,3}; Xiaowei Wang^{1,2}; Shenfengxiang Feng^{1,2}; Xinyi Lyu^{1,2,3}; Yuanyuan Xiong^{1,2}; Hanbin Wu^{1,2}; Jueming Lei^{1,2}; Jihong Xu^{1,2}; Yuan He^{1,2}; Yuanyuan Wang^{1,2}; Ya Zhang^{1,2}; Hongguang Zhang^{1,2}; Ying Yang^{1,2,3,#}; Xu Ma^{1,2,#}

Summary

What is already known about this topic?

Menstrual irregularity is a hallmark clinical feature of polycystic ovary syndrome (PCOS), which is an established risk factor for spontaneous abortion. However, robust population-level evidence directly linking specific patterns of menstrual irregularity to spontaneous abortion risk remains lacking, as prior studies have been limited by small sample sizes.

What is added by this report?

This nationwide study of 3.9 million women demonstrates that abnormal menstrual characteristics — including irregular menstrual cycles and abnormal menstrual period duration — are independent risk factors for spontaneous abortion, each exhibiting a dose-response relationship. The co-occurrence of long cycles and prolonged periods confers the highest overall risk.

What are the implications for public health practice?

These findings offer a scientific basis for formulating public health policies aimed at reducing spontaneous abortion (SA) risk, particularly in resource-limited settings. Menstrual characteristics constitute a simple, low-cost, and readily accessible tool for stratifying SA risk across the perinatal continuum — from preconception counseling through early pregnancy management.

across nine Chinese provinces who participated in the National Free Preconception Care Project (NFPCP) between 2013 and 2024. Logistic regression was used to examine associations between menstrual characteristics and SA risk. Multilevel models, restricted cubic splines, and subgroup analyses were employed as extended analytical approaches to confirm the robustness of findings.

Results Irregular menstrual cycles [adjusted odds ratio (aOR)=1.11, 95% confidence interval (CI): 1.05, 1.17] and irregular periods (aOR=1.23, 95% CI: 1.15, 1.32) were independently associated with SA. The combination of long cycles (38–53 days) and prolonged periods (>7 days) conferred the highest risk (aOR=1.82, 95% CI: 1.33, 2.49). Dose-response relationships and subgroup analyses yielded patterns consistent with the primary analysis.

Conclusions These findings support the utility of menstrual characteristic assessment as an effective tool for SA risk stratification before and during pregnancy. Greater clinical attention should be directed toward women with abnormal menstrual cycles and/or periods, who warrant targeted monitoring and intervention — measures that offer meaningful benefit even for those who have already achieved conception.

ABSTRACT

Introduction Menstrual irregularity is a hallmark clinical feature of polycystic ovary syndrome (PCOS), a well-established risk factor for spontaneous abortion (SA). However, robust population-level evidence directly linking specific menstrual patterns to SA risk remains lacking, as prior studies have been limited by small sample sizes and single-center designs.

Methods We analyzed data from 3.9 million women

Spontaneous abortion (SA), defined as pregnancy loss before viability, affects approximately 23 million women each year. Polycystic ovary syndrome (PCOS), a common endocrinopathy affecting up to 13% of reproductive-age women, is associated with a 49–53% increased risk of SA (1). PCOS is frequently underdiagnosed (2), and its complex diagnostic criteria can delay confirmation by more than 2 years (3). Since anovulation — affecting 75%–85% of PCOS patients (4–5) — is central to the condition and is closely associated with menstrual dysfunction, the

characteristic patterns of these menstrual abnormalities may themselves serve as simple, accessible early warning signs. Menstrual characteristics (cycle length, bleeding duration) directly reflect endocrine health in women of reproductive age (6). Accordingly, assessment of menstrual characteristics represents a simple, low-cost, and clinically accessible tool for stratifying SA risk during preconception and early pregnancy. Incorporating this assessment into routine care could enable earlier identification of high-risk women, facilitating targeted monitoring and timely intervention.

Current research on menstrual characteristics has focused predominantly on fertility assessment during preconception, with limited robust evidence linking specific menstrual patterns to SA risk — a gap attributable in part to the small sample sizes and single-center designs of prior studies (7–9). To address this limitation, we analyzed data from over 3 million reproductive-age women across nine Chinese regions within the National Free Preconception Care Project (NFPCP) to investigate the independent and combined associations of menstrual characteristics with SA risk.

This retrospective cohort study drew on data from the NFPCP, a population-based initiative launched in 2010 by the National Health and Family Planning Commission and the Ministry of Finance to provide

free preconception health check-ups to rural and urban couples throughout mainland China. The project encompasses free health examinations, risk assessments, consultations, early pregnancy follow-ups, and pregnancy outcome follow-ups; its design, organization, and implementation have been described previously (8,10). The present study included women aged 20–49 years from nine provinces spanning eastern, central, and western China (Supplementary Table S1, available at <https://weekly.chinacdc.cn/>) who participated in the NFPCP between January 1, 2013, and December 31, 2024. Women were excluded if they met any of the following criteria: 1) multiple births; 2) ectopic pregnancy; 3) medically induced abortion; or 4) missing data on both menstrual cycle and menstrual period. After applying these exclusion criteria, a final cohort of 3,865,348 women was retained for analysis (Figure 1).

Menstrual characteristics (cycle length and period length) were assessed by trained health workers through structured interviews using a standardized household health questionnaire; data were entered directly into a web-based electronic collection system and transmitted to the national database center. Based on self-reported averages, menstrual cycles were categorized as regular (24–38 days) or irregular (≤ 24 days, 38–53 days, or > 53 days), and menstrual period as normal (3–7 days) or abnormal (< 3 days or > 7 days)

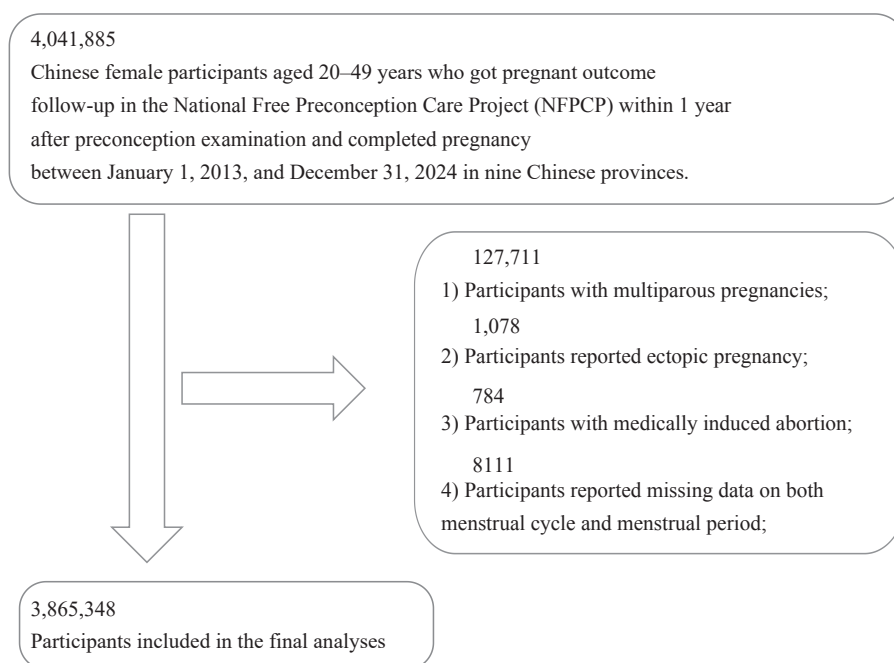


FIGURE 1. Flowchart of the study population selection. Abbreviation: NFPCP=National Free Preconception Care Project.

(11–12). The primary outcome was SA, defined as fetal death or pregnancy loss before 28 weeks of gestation. Baseline characteristics were compared across menstrual regularity groups using descriptive statistics and chi-square (χ^2) tests.

The primary analysis employed logistic regression models to estimate odds ratios (ORs) and 95% confidence intervals (95% CIs) for associations between menstrual characteristics and SA. To minimize confounding, we adjusted for a comprehensive set of covariates spanning demographic, physiological, and lifestyle factors across three sequential models: unadjusted (Model 1), adjusted for high-risk factors (Model 2), and fully adjusted (Model 3); detailed covariate definitions are provided in [Supplementary Table S2](#) (available at <https://weekly.chinacdc.cn/>). As an extended analytical approach, multilevel logistic regression models incorporating random intercepts at the economic region level were fitted to account for the clustering of nine provinces within three economic regions (Model 4). Dose–response relationships were evaluated using restricted cubic splines with four knots. Subgroup analyses stratified by age, body mass index (BMI), and adverse pregnancy history were performed to assess the robustness of the primary findings. Baseline characteristics were compared across menstrual regularity groups using descriptive statistics and chi-square tests. All analyses were performed in R version 4.1.0 (R Foundation for Statistical Computing, Vienna, Austria); two-sided $P < 0.05$ was considered statistically significant.

The final analysis included 3,865,348 reproductive-age women (mean age 27 ± 4.28 years), among whom 64,834 SA events were recorded (incidence: 1.68%). Complete menstrual cycle and menstrual period data were available for 3,860,915 and 3,791,110 women, with irregularity prevalences of 3.05% and 1.63%, respectively. Women with irregular menstrual cycles had higher rates of overweight or obesity and adverse pregnancy history. Women with abnormal menstrual periods showed higher prevalences of scanty menstrual flow and adverse pregnancy history (all $P < 0.001$; [Table 1](#) and [Supplementary Table S3](#), available at <https://weekly.chinacdc.cn/>).

Logistic regression models confirmed that both irregular menstrual cycles [adjusted OR (aOR)=1.11, 95% CI: 1.05, 1.17] and abnormal menstrual periods (aOR=1.23, 95% CI: 1.15, 1.32) were independently associated with increased SA risk after full covariate adjustment (Model 3), with consistent estimates

obtained from multilevel models (Model 4). Further categorization revealed dose-response relationships for both menstrual characteristics. Very long cycles (>53 days) conferred the highest cycle-related risk (aOR=1.25, 95% CI: 1.06, 1.48), and prolonged menstrual periods (>7 days) were also a significant risk factor (aOR=1.31, 95% CI: 1.20, 1.42) ([Table 2](#)). Restricted cubic spline (RCS) analyses revealed a J-shaped relationship for cycle length and a U-shaped relationship for menstrual period duration (both P for nonlinearity <0.001 ; [Figure 2](#)). In joint analyses, isolated irregularities in either dimension were each significantly associated with SA risk (cycles only: aOR=1.10; period only: aOR=1.24). Combined abnormalities showed an elevated point estimate (aOR=1.14, 95% CI: 0.90, 1.44) that did not reach statistical significance, a result likely attributable to the limited number of women in this category yielding wide confidence intervals. Notably, the combination of long cycles (38–53 days) and prolonged periods (>7 days) conferred the highest overall risk (Model 3: aOR=1.82, 95% CI: 1.33, 2.49), with multilevel models (Model 4) producing consistent estimates across all joint exposure categories ([Table 3](#)). Finally, stratified analyses suggested potentially enhanced associations among women aged ≥ 35 years, those with overweight or obesity, and those with a history of adverse pregnancy outcomes ([Figure 3](#)).

DISCUSSION

In this cohort of 3 million reproductive-age women spanning more than a decade, we demonstrated that deviations in menstrual characteristics — both cycle length and bleeding duration — are independently and dose-dependently associated with SA risk, following a J-shaped relationship for cycle length and a U-shaped relationship for period duration. The combination of long cycles and prolonged periods conferred the highest risk, a pattern that remained consistent across all subgroups examined.

These findings extend prior evidence linking menstrual irregularities to adverse pregnancy outcomes. We confirmed that longer menstrual cycles are associated with increased SA risk, consistent with existing literature connecting cycle irregularity to miscarriage and ovulatory dysfunction ([13–14](#)). We further identified prolonged menstrual periods (>7 days) as a significant independent risk factor for SA. Both patterns are consistent with underlying oligo-anovulation or luteal-phase deficiency — conditions

TABLE 1. Baseline characteristics of the study population by menstrual cycle and menstrual period.

Maternal characteristic	Menstruation		P	Menstruation		P
	Menstrual cycle regularity			Menstrual period regularity		
	Regularity (N=3,743,087)	Irregularity (N=117,828)		Regularity (N=3,729,300)	Irregularity (N=61,810)	
Age at last menstrual period, years (n, %)			<0.001			<0.001
20–24	1,478,896 (39.51)	42,220 (35.83)		1,476,231 (39.58)	18,311 (29.62)	
25–29	1,561,268 (41.71)	51,791 (43.95)		1,556,429 (41.73)	26,447 (42.79)	
30–34	506,842 (13.54)	18,032 (15.30)		503,009 (13.49)	11,900 (19.25)	
35–49	196,081 (5.24)	5,785 (4.91)		193,661 (5.19)	5,152 (8.34)	
BMI (kg/m ²)			<0.001			<0.001
Underweight	375,595 (10.03)	11,720 (9.95)		373,815 (10.02)	7,173 (11.60)	
Normal weight	2,672,162 (71.39)	74,583 (63.30)		2,658,515 (71.29)	41,536 (67.20)	
Overweight	551,434 (14.73)	22,657 (19.23)		551,471 (14.79)	10,120 (16.37)	
Obesity	139,047 (3.71)	8,590 (7.29)		140,861 (3.78)	2,896 (4.69)	
Unknown	4,849 (0.13)	278 (0.24)		4,668 (0.13)	85 (0.14)	
Blood sugar status (n, %)			<0.001			<0.001
Normal glucose level	3,216,942 (85.94)	100,104 (84.96)		3,204,832 (85.94)	52,510 (84.95)	
Pre-diabetes	479,579 (12.81)	15,870 (13.47)		477,901 (12.81)	8,490 (13.74)	
Diabetes	29,522 (0.79)	1,204 (1.02)		29,639 (0.79)	504 (0.82)	
Unknown	17,044 (0.46)	650 (0.55)		16,958 (0.45)	306 (0.50)	
Blood pressure status (n, %)			<0.001			<0.001
Normal	3,332,227 (89.02)	100,573 (85.36)		3,319,996 (89.02)	53,053 (85.83)	
Hypertension	39,778 (1.06)	2,118 (1.80)		39,659 (1.06)	1,035 (1.67)	
Unknown	371,082 (9.91)	15,137 (12.85)		369,675 (9.91)	7,722 (12.49)	
Education level (n, %)			<0.001			<0.001
Junior high school or below	2,426,323 (64.82)	66,952 (56.82)		2,408,328 (64.58)	39,942 (64.62)	
Senior high school or above	1,181,982 (31.58)	46,838 (39.75)		1,188,935 (31.88)	20,604 (33.33)	
Unknown	134,782 (3.60)	4,038 (3.43)		132,067 (3.54)	1,264 (2.04)	
Ethnicity (n, %)			<0.001			<0.001
Han	3,610,322 (96.45)	114,083 (96.82)		3,599,376 (96.52)	58,988 (95.43)	
Others	78,800 (2.11)	2,118 (1.80)		77,690 (2.08)	2,384 (3.86)	
Unknown	53,965 (1.44)	1,627 (1.38)		52,264 (1.40)	438 (0.71)	
Residence type (n, %)			<0.001			<0.001
Urban	233,895 (6.25)	12,246 (10.39)		234,371 (6.28)	6,240 (10.10)	
Rural	3,509,192 (93.75)	105,582 (89.61)		3,494,959 (93.72)	55,570 (89.90)	
Dysmenorrhea status (n, %)			<0.001			<0.001
None	2,902,978 (77.56)	76,971 (65.32)		2,886,189 (77.39)	44,034 (71.24)	
Mild	801,085 (21.40)	36,225 (30.74)		803,670 (21.55)	16,268 (26.32)	
Severe	29,654 (0.79)	2,903 (2.46)		30,188 (0.81)	1,347 (2.18)	
Unknown	9,370 (0.25)	1,729 (1.47)		9,283 (0.25)	161 (0.26)	
Menstrual flow (n, %)			<0.001			<0.001
Slight	72,082 (1.93)	6,726 (5.79)		69,712 (1.87)	6,110 (9.90)	
Moderate	3,594,677 (96.22)	104,608 (90.03)		3,584,151 (96.30)	52,417 (84.93)	
Large	69,000 (1.85)	4,857 (4.18)		68,144 (1.83)	3,188 (5.17)	

Continued

Maternal characteristic	Menstruation		P	Menstruation		P
	Menstrual cycle regularity			Menstrual period regularity		
	Regularity (N=3,743,087)	Irregularity (N=117,828)		Regularity (N=3,729,300)	Irregularity (N=61,810)	
Adverse pregnancy history (n, %)			<0.001			<0.001
No	3,186,350 (85.13)	90,078 (76.45)		3,173,876 (85.11)	44,634 (72.21)	
Yes	556,737 (14.87)	27,750 (23.55)		555,454 (14.89)	17,176 (27.79)	
Parity (n, %)			<0.001			<0.001
Nulliparous	2,085,449 (55.71)	61,670 (52.34)		2,081,289 (55.81)	25,327 (40.98)	
Parous	1,657,638 (44.29)	56,158 (47.66)		1,648,041 (44.19)	36,483 (59.02)	

Note: All variables are presented as N (%). All $P < 0.001$. Baseline characteristics were stratified by menstrual cycle regularity — regular (24–38 days) versus irregular (<24 days, 38–53 days, or >53 days) — and by menstrual bleeding duration (menstrual period) — normal (3–7 days) versus abnormal (<3 or >7 days). All variables were classified into appropriate categories, and chi-square (χ^2) tests were used to calculate P .

Abbreviation: BMI=body mass index.

TABLE 2. Individual associations between menstrual characteristics and SA risk.

Characteristic	Total participants	SA cases <i>n</i> (%)	OR (95% CI)			
			Model 1	Model 2	Model 3	Model 4
Menstrual cycle						
Binary classification						
Regular (24–38 days)	3,743,087	62,165 (1.66)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)
Irregular	117,828	2,559 (2.17)	1.31 (1.26, 1.37)	1.23 (1.18, 1.29)	1.11 (1.05, 1.17)	1.11 (1.05, 1.17)
Detailed classification						
24D–38D	3,743,087	62,165 (1.66)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)
<24D	69,579	1,454 (2.09)	1.26 (1.20, 1.33)	1.22 (1.15, 1.29)	1.12 (1.04, 1.20)	1.10 (1.02, 1.17)
39D–53D	38,707	865 (2.23)	1.35 (1.26, 1.45)	1.22 (1.13, 1.32)	1.06 (0.97, 1.17)	1.09 (0.99, 1.12)
>53D	9,542	240 (2.52)	1.53 (1.34, 1.74)	1.40 (1.21, 1.61)	1.25 (1.06, 1.48)	1.25 (1.06, 1.48)
Menstrual period						
Binary classification						
Regular (3D–7D)	3,729,330	61,780 (1.66)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)
Irregular	61,810	1,643 (2.66)	1.62 (1.54, 1.70)	1.44 (1.36, 1.52)	1.23 (1.15, 1.32)	1.26 (1.17, 1.35)
Detailed classification						
3D–7D	3,729,330	61,780 (1.66)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)
<3D	29,729	743 (2.50)	1.52 (1.41, 1.64)	1.35 (1.25, 1.46)	1.11 (0.99, 1.25)	1.18 (1.05, 1.32)
>7D	32,081	900 (2.81)	1.71 (1.60, 1.83)	1.53 (1.42, 1.64)	1.31 (1.20, 1.42)	1.31 (1.20, 1.42)

Note: Data are presented as odds ratio (95% CI). Estimates with 95% CIs excluding 1 are considered statistically significant at $P < 0.05$. Models examined associations between menstrual cycle regularity (regular: 24–38 days; irregular: <24, 38–53, or >53 days) and menstrual period duration (normal: 3–7 days; abnormal: <3 or >7 days) with SA risk. Four models were fitted: Model 1 (unadjusted), Model 2 (adjusted for high-risk factors), and Model 3 (fully adjusted) using conventional logistic regression; Model 4 (fully adjusted) using multilevel logistic regression as an extended analytical approach. All covariates are listed in [Supplementary Table S2](#).

Abbreviation: OR=odds ratio; CI=confidence interval; SA=spontaneous abortion.

relevant to PCOS, in which hyperandrogenism and progesterone insufficiency may compromise endometrial receptivity and impair early pregnancy maintenance (15–16). Direct mechanistic verification was beyond the scope of the present study. Nevertheless, our findings provide a scientific basis for public health policies targeting SA prevention,

particularly in resource-limited settings. Menstrual characteristics represent a simple, low-cost, and readily accessible tool for stratifying SA risk across the perinatal continuum — from preconception counseling through early pregnancy. Early identification of women with menstrual abnormalities facilitates timely intervention for underlying conditions

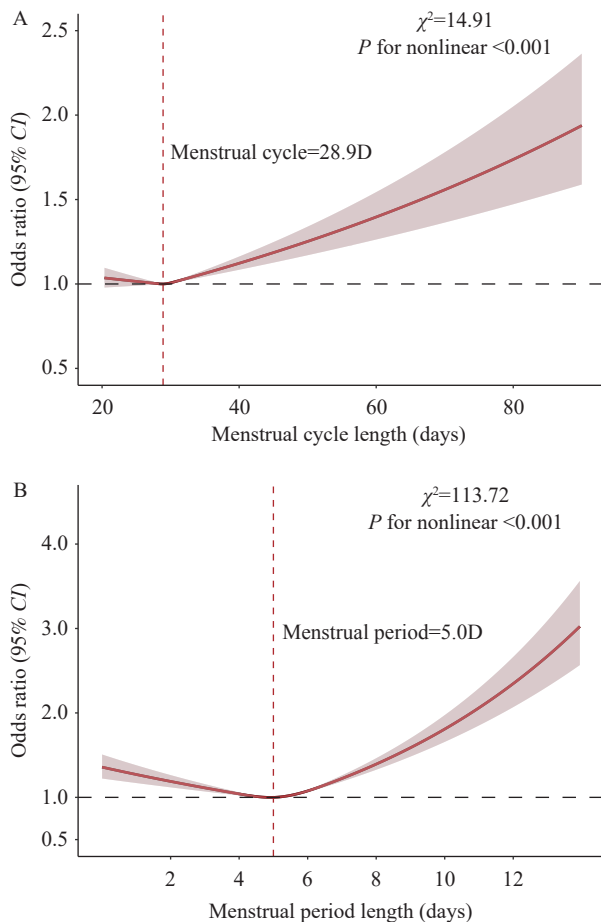


FIGURE 2. Dose-response relationship between menstrual characteristics and SA risk (restricted cubic spline analysis). (A) Menstrual cycle length (RCS-adjusted logistic regression); (B) Menstrual period length (RCS-adjusted logistic regression).

Note: This figure displays restricted cubic spline (RCS) curves from logistic regression models examining the dose-response relationships between menstrual cycle length and menstrual period duration with spontaneous abortion (SA) risk. The red dashed horizontal line represents an OR of 1.0, indicating no difference in risk relative to the reference. Solid red lines indicate the estimated ORs, and shaded ribbons denote the corresponding 95% CIs. Logistic regression models were adjusted for all covariates listed in Supplementary Table S2. Bolded values indicate statistical significance ($P<0.05$). The χ^2 and P for nonlinearity are reported in each panel to evaluate the significance of the non-linear association.

Abbreviation: RCS=Restricted Cubic Spline; SA=Spontaneous Abortion; χ^2 =Chi-square statistic; OR=odds ratio; CI=confidence interval.

such as PCOS or obesity-related anovulation, before pregnancy complications emerge. Evidence supports that lifestyle modification and low-cost pharmacological interventions can restore ovulatory cycles, improve metabolic health, and enhance

reproductive outcomes (5,11). Although menstrual data are routinely recorded in clinical practice, they remain substantially underutilized in pregnancy risk assessment. Greater clinical attention should therefore be directed toward women with abnormal cycles or periods — not only during preconception counseling, but also after conception has occurred — as both groups stand to benefit from targeted monitoring and timely intervention. Embedding this risk-aware approach throughout the perinatal continuum has the potential to meaningfully improve pregnancy outcomes.

This study has several notable strengths. First, the large-scale design, drawing on long-term data from nine Chinese regions, provided sufficient statistical power to detect moderate yet clinically meaningful associations between menstrual characteristics and SA risk — associations that prior smaller studies were underpowered to establish. Second, the application of multiple analytical approaches to evaluate distinct menstrual dimensions, combined with detailed exposure categorization, enhanced both the robustness and granularity of our findings. Importantly, all menstrual characteristics were ascertained prior to pregnancy, which minimizes recall bias and establishes a clear temporal sequence between exposure and outcome. Finally, the availability of comprehensive individual-level covariate data enabled well-powered and informative subgroup analyses across key demographic and clinical strata.

This study has several limitations that warrant consideration. First, menstrual characteristics were self-reported at the preconception health visit, which may introduce misclassification arising from recall or reporting inaccuracies. Second, the absence of biochemical or hormonal biomarkers precluded direct verification of ovulatory status or quantification of endocrine dysfunction. Third, residual confounding cannot be excluded, as certain variables remained unmeasured or unidentified — including genetic factors (such as embryonic or parental chromosomal abnormalities) and environmental exposures at the population level (e.g., air pollution, climate, temperature, humidity, and atmospheric pressure). Fourth, although participants were drawn from nine provinces, the majority were rural residents; consequently, our findings may not be fully generalizable to all reproductive-age women in China, particularly those of higher socioeconomic status or those who do not seek preconception care.

Conflicts of interest: No conflicts of interest.

TABLE 3. Combined associations between menstrual characteristics and SA risk.

Menstrual Characteristic Pattern	Total participants	SA cases n (%)	OR (95% CI)			
			Model 1	Model 2	Model 3	Model 4
Binary joint classification						
Both normal	3,657,947	60,231 (1.62)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)
Irregular cycle only	67,043	1,444 (2.11)	1.31 (1.25, 1.39)	1.20 (1.13, 1.27)	1.10 (1.01, 1.17)	1.11 (1.03, 1.20)
Irregular period only	56,991	1,498 (2.56)	1.61 (1.53, 1.70)	1.42 (1.34, 1.50)	1.24 (1.16, 1.34)	1.27 (1.18, 1.37)
Both abnormal	4,726	140 (2.88)	1.82 (1.54, 2.16)	1.45 (1.20, 1.76)	1.14 (0.90, 1.44)	1.18 (0.93, 1.49)
Detailed joint classification						
Both normal	3,657,947	60,231 (1.62)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)	1.00 (Reference)
Short cycle & short period	1,414	33 (2.28)	1.43 (1.01, 2.02)	1.01 (0.67, 1.51)	0.97 (0.65, 1.47)	0.68 (0.38, 1.20)
Short cycle & regular period	21,392	420 (1.93)	1.20 (1.09, 1.32)	1.11 (1.00, 1.23)	1.07 (0.97, 1.20)	1.12 (0.97, 1.29)
Short cycle & long period	804	30 (3.60)	2.23 (1.61, 3.33)	1.75 (1.15, 2.66)	1.52 (0.99, 2.32)	1.44 (0.90, 2.31)
Regular cycle & short period	27,655	690 (2.43)	1.53 (1.42, 1.65)	1.34 (1.24, 1.46)	1.31 (1.20, 1.42)	1.22 (1.08, 1.38)
Regular cycle & long period	29,336	808 (2.68)	1.69 (1.58, 1.82)	1.50 (1.39, 1.62)	1.39 (1.29, 1.50)	1.30 (1.19, 1.42)
Long cycle & short period	472	15 (3.08)	1.96 (1.17, 3.28)	1.36 (0.75, 2.48)	1.37 (0.75, 2.51)	1.31 (0.61, 2.79)
Long cycle & regular period	36,704	802 (2.14)	1.33 (1.24, 1.43)	1.20 (1.11, 1.29)	1.19 (1.10, 1.29)	1.07 (0.97, 1.17)
Long cycle & long period	1,463	46 (3.05)	1.94 (1.45, 2.60)	1.98 (1.46, 2.69)	1.82 (1.33, 2.49)	1.64 (1.15, 2.34)
Very long cycle & short period	146	2 (1.35)	0.83 (0.21, 3.35)	0.78 (0.19, 3.17)	0.75 (0.18, 3.04)	0.58 (0.08, 4.21)
Very long cycle & regular period	8,947	222 (2.42)	1.52 (1.33, 1.74)	1.41 (1.22, 1.63)	1.40 (1.21, 1.62)	1.29 (1.08, 1.53)
Very long cycle & long period	427	14 (3.17)	2.02 (1.19, 3.45)	1.08 (0.51, 2.28)	1.04 (0.50, 2.21)	0.59 (0.19, 1.86)

Note: Exposure categories were defined based on combinations of menstrual cycle length and menstrual period duration. The reference group comprised women with both normal cycle length and normal period duration. Both normal: normal menstrual cycle length (24–38 days) and normal period duration (3–7 days). Irregular cycle only: irregular menstrual cycle (<24 or >38 days) with normal period duration. Irregular period only: normal menstrual cycle with irregular period duration (<3 or >7 days). Both abnormal: irregular menstrual cycle and irregular period duration. Four models were constructed: Model 1 (unadjusted), Model 2 (adjusted for high-risk factors), and Model 3 (fully adjusted) using conventional logistic regression; Model 4 (fully adjusted) using multilevel logistic regression as an extended analytical approach. All covariates are listed in [Supplementary Table S2](#). Results are presented as odds ratios (95% CI). Associations are considered statistically significant at $P < 0.05$, corresponding to 95% CIs that exclude 1.

Abbreviation: CI=confidence interval; SA=spontaneous abortion.

Acknowledgements: All healthcare professionals and participants across administrative divisions for their invaluable contributions to the National Free Preconception Care Project.

Ethical statements: Approved by the Institutional Review Board of the National Research Institute for Family Planning (IRB-201001). Written informed consent was obtained from all National Free Preconception Care Project (NFPCP) participants. The study was conducted in accordance with the principles of the World Medical Association Declaration of Helsinki (2000), and its reporting adheres to the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) guidelines.

Funding: Supported by the National Key Research and Development Program of China (NKP) under grants 2021YFC2700705 (Ms. Yang) and 2016YFC100307 (Mr. Ma).

doi: [10.46234/ccdcw2026.045](https://doi.org/10.46234/ccdcw2026.045)

Corresponding authors: Ying Yang, yangying@nrfp.org.cn; Xu Ma, maxu_ky@nrfp.org.cn.

¹ National Research Institute for Family Planning, Chinese Academy of Medical Sciences, Beijing, China; ² National Human Genetic Resource Center, National Human Reproduction and Health Resource Center, Beijing, China; ³ Peking Union Medical College, Beijing, China; ⁴ Peking University First Hospital, Beijing, China.

& Joint first authors.

Copyright © 2026 by Chinese Center for Disease Control and Prevention. All content is distributed under a Creative Commons Attribution Non Commercial License 4.0 (CC BY-NC).

Submitted: January 03, 2026

Accepted: March 02, 2026

Issued: March 06, 2026

REFERENCES

1. Bahri Khomami M, Hashemi S, Shorakae S, Harrison CL, Piltonen TT, Romualdi D, et al. Systematic review and meta-analysis of birth outcomes in women with polycystic ovary syndrome. *Nat Commun*

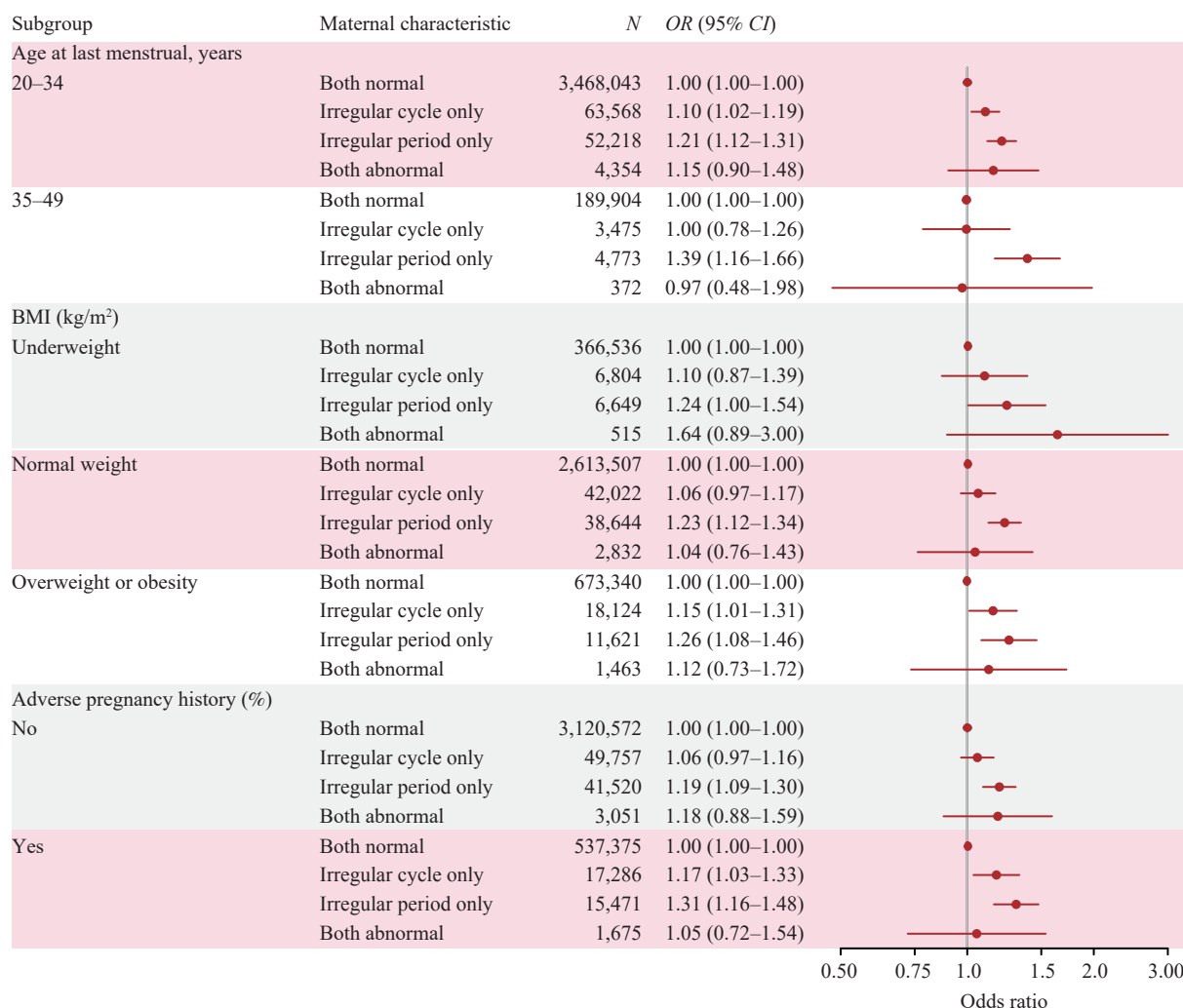


FIGURE 3. Forest plot of the association between menstrual characteristics and spontaneous abortion (SA) risk, stratified by age, BMI, and adverse pregnancy history.

Note: This forest plot presents subgroup analysis results examining the association between combined menstrual characteristic categories and SA risk. ORs and 95% CIs were derived from fully adjusted multivariable logistic regression models. Exposure categories were defined based on combinations of menstrual cycle length and menstrual period duration, with women having both normal cycle length and normal period length serving as the reference group. Both Normal: normal menstrual cycle length (24–38 days) and normal period length (3–7 days). Irregular Cycle Only: irregular menstrual cycle (<24 or >38 days) with normal period length. Irregular Period Only: normal menstrual cycle with irregular period length (<3 or >7 days). Both Abnormal: irregular menstrual cycle and irregular period length. Analyses were stratified by three key maternal characteristics: age at last menstrual period (20–34 vs. 35–49 years); body mass index (BMI) category (underweight, normal weight, or overweight/obesity); and a history of adverse pregnancy (no vs. yes). Within each stratum, four mutually exclusive exposure categories were compared, with the "Both Normal" group serving as the common reference (OR=1.00). Each estimate is represented by a solid square (point estimate, OR) with a horizontal line denoting the 95% CI. The area of each square is proportional to the precision of the estimate. The vertical dashed line at OR=1.0 indicates no association; statistical significance is indicated when the 95% CI does not cross this line (i.e., excludes 1.0). Abbreviation: OR=odds ratio; CI=confidence interval; SA=spontaneous abortion; BMI=body mass index (kg/m²).

- 2024;15(1):5592. <https://doi.org/10.1038/s41467-024-49752-6>.
2. Teede HJ, Tay CT, Laven JJE, Dokras A, Moran LJ, Piltonen TT, Costello MF, et al. Recommendations from the 2023 international evidence-based guideline for the assessment and management of polycystic ovary syndrome. *Eur J Endocrinol*. 2023 Aug 2;189(2):G43–G64. doi: 10.1093/ejendo/lvad096. PMID: 37580861.
3. Gibson-Helm M, Teede H, Dunaif A, Dokras A. Delayed diagnosis and a lack of information associated with dissatisfaction in women with polycystic ovary syndrome. *J Clin Endocrinol Metab* 2016;102(2):604–12. <https://doi.org/10.1210/jc.2016-2963>.
4. Azziz R, Carmina E, Dewailly D, Diamanti-Kandarakis E, Escobar-Morreale HF, Futterweit W, et al. The androgen excess and PCOS society criteria for the polycystic ovary syndrome: the complete task force report. *Fertil Steril* 2009;91(2):456–88. <https://doi.org/10.1016/j.fertnstert.2008.06.035>.
5. Solomon CG. THE EPIDEMIOLOGY OF POLYCYSTIC OVARY SYNDROME: prevalence and associated disease risks. *Endocrinol Metab Clin North Am* 1999;28(2):247–63. <https://doi.org/10.1016/>

- S0889-8529(05)70069-4.
6. Harlow SD, Ephross SA. Epidemiology of menstruation and its relevance to women's health. *Epidemiol Rev* 1995;17(2):265 – 86. <https://doi.org/10.1093/oxfordjournals.epirev.a036193>.
 7. Cao YQ, Zhao XL, Dou Z, Gong Z, Wang BJ, Xia T. The correlation between menstrual characteristics and fertility in women of reproductive age: a systematic review and meta-analysis. *Fertil Steril* 2024;122(5):918 – 27. <https://doi.org/10.1016/j.fertnstert.2024.06.016>.
 8. Zhao CY, Jia JJ, Wu HB, Xu Q, Lyu XY, Liu MY, et al. Maternal preconception serum alanine aminotransferase levels and risk of preterm birth among reproductive-aged women — China, 2013-2017. *China CDC Wkly* 2024;6(10):181 – 8. <https://doi.org/10.46234/ccdcw2024.037>.
 9. Wang ZW, Yan JJ, Chen HF, He LM, Xu SH. The reproductive endocrine feature and conception outcome of women with unknown etiological menstrual cycle (36-45 days) with long follicular phase. *Gynecol Endocrinol* 2022;38(9):742 – 7. <https://doi.org/10.1080/09513590.2022.2101636>.
 10. Wei YM, Xu Q, Yang HX, Yang Y, Wang L, Chen H, et al. Preconception diabetes mellitus and adverse pregnancy outcomes in over 6.4 million women: a population-based cohort study in China. *PLoS Med* 2019;16(10):e1002926. <https://doi.org/10.1371/journal.pmed.1002926>.
 11. Zhang Q, Wang YY, Zhang Y, Zhang HG, Yang Y, He Y, et al. The influence of age at menarche, menstrual cycle length and bleeding duration on time to pregnancy: a large prospective cohort study among rural Chinese women. *BJOG* 2017;124(11):1654 – 62. <https://doi.org/10.1111/1471-0528.14469>.
 12. Munro MG, Critchley HOD, Fraser IS, the FIGO Menstrual Disorders Committee. The two FIGO systems for normal and abnormal uterine bleeding symptoms and classification of causes of abnormal uterine bleeding in the reproductive years:2018 revisions. *Int J Gynecol Obstet* 2018;143(3):393 – 408. <https://doi.org/10.1002/ijgo.12666>.
 13. Rowland AS, Baird DD, Long S, Wegienka G, Harlow SD, Alavanja M, et al. Influence of medical conditions and lifestyle factors on the menstrual cycle. *Epidemiology* 2002;13(6):668 – 74. <https://doi.org/10.1097/00001648-200211000-00011>.
 14. Small CM, Manatunga AK, Klein M, Feigelson HS, Dominguez CE, McChesney R, et al. Menstrual cycle characteristics: associations with fertility and spontaneous abortion. *Epidemiology* 2006;17(1):52 – 60. <https://doi.org/10.1097/01.ede.0000190540.95748.e6>.
 15. Chen YH, Guo JY, Zhang QW, Zhang CL. Insulin resistance is a risk factor for early miscarriage and macrosomia in patients with polycystic ovary syndrome from the first embryo transfer cycle: a retrospective cohort study. *Front Endocrinol* 2022;13:853473. <https://doi.org/10.3389/fendo.2022.853473>.
 16. Huang WL, Liu C, Ding L, Li Y, Zhou HS, Wang SW, et al. The effect of β -cell dysfunction on reproductive outcomes of PCOS undergoing IVF or ICSI embryo transfer cycles: a retrospective cohort study. *Front Endocrinol* 2024;15:1327041. <https://doi.org/10.3389/fendo.2024.1327041>.

SUPPLEMENTARY MATERIAL

SUPPLEMENTARY TABLE S1. Geographic distribution of the study population by economic region and PLAD.

Economic region	PLADs	Participants <i>N</i>	Region Subtotal <i>N (%)</i>	SA cases <i>N</i>	SA rate <i>N (%)</i>
Eastern Region		876,531	22.68	1,801	0.21
	Tianjin	51,651	1.34	233	0.45
	Shandong	793,426	20.53	1,387	0.17
	Liaoning	31,454	0.81	181	0.58
Central Region		2,359,289	61.04	5,373	0.23
	Henan	1,716,248	44.40	3,142	0.18
	Jilin	52,736	1.36	234	0.44
	Anhui	590,305	15.27	1997	0.34
Western Region		629,528	16.29	5163	0.82
	Sichuan	302,767	7.83	3,940	1.30
	Gansu	285,458	7.39	682	0.24
	Qinghai	41,303	1.07	541	1.31

Note: Data were derived from the National Free Preconception Care Project (NFPCP) database.

Abbreviation: SA=spontaneous abortion; PLAD=provincial-level administrative division.

SUPPLEMENTARY TABLE S2. Definitions of covariates.

Covariates	Definition
Demographic Factors	
Education level	The highest level of educational attainment of the participant (junior high school and below, senior high school or above).
Ethnicity	The ethnic group to which the participant belongs (Han or other).
Residence type	The type of area where the participant lives (rural or urban).
Individual Physiological Factors	
Age	Maternal age at the last menstrual period of the participant (20–24, 25–29, 30–34, 35–39, and ≥ 40 years).
Reproductive system status	Whether the participant has a history of infertility or adnexal inflammation (yes or no).
Endocrine system abnormalities	
BMI level	Calculated as weight in kilograms divided by height in meters squared [underweight (less than 18.5 kg/m ²), normal (18.5–23.9 kg/m ²), overweight (24.0–27.9 kg/m ²), or obese (28.0 kg/m ² or more)].
Blood sugar status	Self-reported hyperglycemia or fasting blood glucose ≥ 6.1 mmol/L (yes or no).
Blood pressure status	Self-reported hypertension or systolic blood pressure (SBP) ≥ 140 mmHg or diastolic blood pressure (DBP) ≥ 90 mmHg (yes or no).
Thyroid function status	Self-reported thyroid disease or abnormal thyroid function (yes or no).
Immune system status	Whether the participant's white blood cell count was $\geq 4.0 \times 10^9/L$ and $\leq 10.0 \times 10^9/L$ (yes or no).
Obstetric History Covariates	
Adverse pregnancy history	Participants had any adverse pregnancy outcomes (yes or no).
Parity	Whether the participant had previous deliveries (nulliparous or parous).
Menstruation-related Covariates	
Dysmenorrhea status	Participants self-reported whether they experienced dysmenorrhea during their menstrual period (none, mild, or severe).
Menstrual flow	Participants self-reported menstrual blood volume (slight, moderate, or large).
Age of menarche status	Whether the menarche age reported by participants is normal (between 10–15 years old) (normal or abnormal).
Lifestyle and Behavior-Related Factors	
Smoking status	Whether the participant actively smokes (yes or no).
Secondhand smoking status	Whether the participant smokes passively (yes or no).
Alcohol consumption	Whether the participant has consumed alcohol (yes or no).
Pressure status	Whether the participant experiences work and life pressure, social pressure, or economic pressure (yes or no).
Medication status	Whether the participant is currently on medication (yes or no).

Note: Four logistic regression models were constructed. Model 1 was unadjusted. Model 2 was adjusted for high-risk factors, including age, BMI, fasting plasma glucose, blood pressure, adverse pregnancy history, and menstruation-related covariates. Model 3 was fully adjusted, incorporating demographic factors (education, ethnicity, and residence), physiological factors (age, BMI, blood glucose, blood pressure, thyroid function, and immune status), reproductive factors (reproductive system status, adverse pregnancy history, parity, dysmenorrhea, menstrual flow, and age at menarche), and lifestyle factors (smoking, secondhand smoke exposure, alcohol consumption, psychological stress, and medication use). Model 4 applied the same full covariate adjustment as Model 3 within a multilevel logistic regression framework to account for clustering by economic region.

Abbreviation: BMI=body mass index.

SUPPLEMENTARY TABLE S3. Complete baseline characteristics of the study population by menstrual cycle regularity and menstrual period regularity.

Maternal characteristic	Menstruation		P	Menstruation		P
	Menstrual cycle regularity			Menstrual period regularity		
	Regularity (N=3,743,087)	Irregularity (N=117,828)		Regularity (N=3,729,300)	Irregularity (N=61,810)	
Age at last menstrual period, years (n, %)			<0.001			<0.001
20–24	1,478,896 (39.51)	42,220 (35.83)		1,476,231 (39.58)	18,311 (29.62)	
25–29	1,561,268 (41.71)	51,791 (43.95)		1,556,429 (41.73)	26,447 (42.79)	
30–34	506,842 (13.54)	18,032 (15.30)		503,009 (13.49)	11,900 (19.25)	
35–49	196,081 (5.24)	5,785 (4.91)		193,661 (5.19)	5,152 (8.34)	
BMI (kg/m ²)			<0.001			<0.001
Underweight	375,595 (10.03)	11,720 (9.95)		373,815 (10.02)	7,173 (11.60)	
Normal weight	2,672,162 (71.39)	74,583 (63.30)		2,658,515 (71.29)	41,536 (67.20)	
Overweight	551,434 (14.73)	22,657 (19.23)		551,471 (14.79)	10,120 (16.37)	
Obesity	139,047 (3.71)	8,590 (7.29)		140,861 (3.78)	2,896 (4.69)	
Unknown	4,849 (0.13)	278 (0.24)		4,668 (0.13)	85 (0.14)	
Blood sugar status (n, %)			<0.001			<0.001
Normal glucose level	3,216,942 (85.94)	100,104 (84.96)		3,204,832 (85.94)	52,510 (84.95)	
Prediabetes	479,579 (12.81)	15,870 (13.47)		477,901 (12.81)	8,490 (13.74)	
Diabetes	29,522 (0.79)	1,204 (1.02)		29,639 (0.79)	504 (0.82)	
Unknown	17,044 (0.46)	650 (0.55)		16,958 (0.45)	306 (0.50)	
Blood pressure status (n, %)			<0.001			<0.001
Normal	3,332,227 (89.02)	100,573 (85.36)		3,319,996 (89.02)	53,053 (85.83)	
Hypertension	39,778 (1.06)	2,118 (1.80)		39,659 (1.06)	1,035 (1.67)	
Unknown	371,082 (9.91)	15,137 (12.85)		369,675 (9.91)	7,722 (12.49)	
Thyroid function status (n, %)			<0.001			<0.001
Normal	3,735,987 (99.81)	117,312 (99.56)		3,722,199 (99.81)	61,526 (99.54)	
Abnormal	7,100 (0.19)	516 (0.44)		7,131 (0.19)	284 (0.46)	
Reproductive system status (n, %)			<0.001			<0.001
Normal	1,890,043 (50.49)	74,883 (63.55)		1,885,006 (50.55)	37,559 (60.77)	
Abnormal	9,605 (0.26)	1,023 (0.87)		9,680 (0.26)	536 (0.87)	
Unknown	1,843,439 (49.25)	41,922 (35.58)		1,834,644 (49.20)	23,715 (38.37)	
Immune system status (n, %)			<0.001			<0.001
Normal	1,746,894 (46.67)	69,752 (59.20)		1,742,750 (46.73)	34,598 (55.97)	
Abnormal	147,217 (3.93)	5,886 (5.00)		146,451 (3.93)	3,382 (5.47)	
Unknown	1,848,976 (49.40)	42,190 (35.81)		1,840,129 (49.34)	23,830 (38.55)	
Dysmenorrhea status (n, %)			<0.001			<0.001
None	2,902,978 (77.56)	76,971 (65.32)		2,886,189 (77.39)	44,034 (71.24)	
Mild	801,085 (21.40)	36,225 (30.74)		803,670 (21.55)	16,268 (26.32)	
Severe	29,654 (0.79)	2,903 (2.46)		30,188 (0.81)	1,347 (2.18)	
Unknown	9,370 (0.25)	1,729 (1.47)		9,283 (0.25)	161 (0.26)	
Menstrual flow (n, %)			<0.001			<0.001
Slight	72,082 (1.93)	6,726 (5.79)		69,712 (1.87)	6,110 (9.90)	
Moderate	3,594,677 (96.22)	104,608 (90.03)		3,584,151 (96.30)	52,417 (84.93)	
Large	69,000 (1.85)	4,857 (4.18)		68,144 (1.83)	3,188 (5.17)	
Age of menarche status (n, %)			<0.001			<0.001
Normal	3,461,387 (92.47)	102,707 (87.17)		3,458,062 (92.73)	53,889 (87.18)	

Continued

Maternal characteristic	Menstruation		P	Menstruation		P
	Menstrual cycle regularity			Menstrual period regularity		
	Regularity (N=3,743,087)	Irregularity (N=117,828)		Regularity (N=3,729,300)	Irregularity (N=61,810)	
Abnormal	263,662 (7.04)	14,345 (12.17)		263,396 (7.06)	7,784 (12.59)	
Unknown	18,038 (0.48)	776 (0.66)		7,872 (0.21)	137 (0.22)	
Education level (n, %)			<0.001			<0.001
Junior high school or below	2,426,323 (64.82)	66,952 (56.82)		2,408,328 (64.58)	39,942 (64.62)	
Senior high school or above	1,181,982 (31.58)	46,838 (39.75)		1,188,935 (31.88)	20,604 (33.33)	
Unknown	134,782 (3.60)	4,038 (3.43)		132,067 (3.54)	1,264 (2.04)	
Ethnicity (n, %)			<0.001			<0.001
Han	3,610,322 (96.45)	114,083 (96.82)		3,599,376 (96.52)	58,988 (95.43)	
Non-Han	78,800 (2.11)	2,118 (1.80)		77,690 (2.08)	2,384 (3.86)	
Unknown	53,965 (1.44)	1,627 (1.38)		52,264 (1.40)	438 (0.71)	
Residence type (n, %)			<0.001			<0.001
Urban	233,895 (6.25)	12,246 (10.39)		234,371 (6.28)	6,240 (10.10)	
Rural	3,509,192 (93.75)	105,582 (89.61)		3,494,959 (93.72)	55,570 (89.90)	
Smoking status (n, %)			<0.001			<0.001
Never smoker	3,729,951 (99.65)	117,070 (99.36)		3,716,073 (99.64)	61,465 (99.44)	
Ever smoker	6,728 (0.18)	490 (0.42)		6,814 (0.18)	235 (0.38)	
Unknown	6,408 (0.17)	268 (0.23)		6,443 (0.17)	110 (0.18)	
Secondhand smoking status (n, %)			<0.001			<0.001
No	3,418,224 (91.32)	98,151 (83.30)		3,403,108 (91.25)	52,603 (85.10)	
Yes	318,264 (8.50)	19,415 (16.48)		319,601 (8.57)	9,100 (14.72)	
Unknown	6,599 (0.18)	262 (0.22)		6,621 (0.18)	107 (0.17)	
Alcohol consumption (n, %)			<0.001			<0.001
Never drinker	3,683,329 (98.40)	114,217 (96.94)		3,669,155 (98.39)	60,026 (97.11)	
Ever drinker	51,065 (1.36)	3,240 (2.75)		51,459 (1.38)	1,618 (2.62)	
Unknown	8,693 (0.23)	371 (0.31)		8,716 (0.23)	166 (0.27)	
Pressure status (n, %)			<0.001			<0.001
No	3,720,428 (99.39)	116,281 (98.69)		3,706,452 (99.39)	61,149 (98.93)	
Yes	9,324 (0.25)	1,024 (0.87)		9,511 (0.26)	445 (0.72)	
Unknown	13,335 (0.36)	523 (0.44)		13,367 (0.36)	216 (0.35)	
Medication status (n, %)			<0.001			<0.001
No use	3,637,488 (97.18)	111,814 (94.90)		3,622,825 (97.14)	58,513 (94.67)	
Use	95,995 (2.56)	5,674 (4.82)		96,858 (2.60)	3,161 (5.11)	
Unknown	9,604 (0.26)	340 (0.29)		9,647 (0.26)	136 (0.22)	
Adverse pregnancy history (n, %)			<0.001			<0.001
No	3,186,350 (85.13)	90,078 (76.45)		3,173,876 (85.11)	44,634 (72.21)	
Yes	556,737 (14.87)	27,750 (23.55)		555,454 (14.89)	17,176 (27.79)	
Parity (n, %)			<0.001			<0.001
Nulliparous	2,085,449 (55.71)	61,670 (52.34)		2,081,289 (55.81)	25,327 (40.98)	
Parous	1,657,638 (44.29)	56,158 (47.66)		1,648,041 (44.19)	36,483 (59.02)	

Note: All variables are presented as N (%). All $P < 0.001$. Baseline characteristics were stratified by menstrual cycle regularity — regular (24–38 days) versus irregular (<24 days, 38–53 days, or >53 days) — and by menstrual bleeding duration (menstrual period) — normal (3–7 days) versus abnormal (<3 or >7 days). All variables were classified into appropriate categories, and chi-square (χ^2) tests were used to calculate P .

Abbreviation: BMI=body mass index.