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Menstrual symptoms and risk of preterm birth: A population-based longitudinal study

Abstract

Objectives: To examine the prospective association between menstrual symptoms before pregnancy and preterm birth. **Methods:** Secondary analysis of data from 14 247 young Australian women born between 1973 and 1978 who participated in a longitudinal, population-based cohort study between 1996 and 2015. Women were first surveyed at 18-23 years, and seven waves of data were collected at roughly three-yearly intervals. At each survey, women were asked about “severe period pain,” “heavy periods,” and “irregular periods” within the last 12 months. From 2009 onward, information on their children was collected, including birth dates and preterm birth (<37>weeks). Logistic regression using generalized estimating equations was used to examine prospective associations between self-reported menstrual symptoms before pregnancy and risk of preterm birth. **Results:** Data from 6615 mothers who had 12 337 live singleton births were available for analysis. Among all births, women reporting severe period pain (adjusted odds ratio [aOR] 1.34 [95% CI 1.10-1.62]) or heavy periods (1.25 [1.02-1.53]) before pregnancy had higher odds of preterm birth. However, in analyses stratified by birth order, only severe period pain (2.05 [1.41-2.99]), heavy periods (1.77 [1.23-2.55]), or irregular periods (1.58 [1.10-2.28]) before a second or subsequent birth were associated with an increased risk of preterm birth. **Conclusions:** Severe period pain, heavy periods, and irregular periods before a second or subsequent birth may be associated with preterm birth.

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1 **Abstract**

2 **Objectives:** To examine the prospective association between menstrual symptoms before
3 pregnancy and preterm birth.

4 **Methods:** Secondary analysis of data from 14,247 young Australian women born between
5 1973 and 1978 who participated in a longitudinal, population-based cohort study between
6 1996 and 2015. Women were first surveyed at 18-23 years and seven waves of data were
7 collected at roughly three-yearly intervals. At each survey, women were asked about 'severe
8 period pain', 'heavy periods' and 'irregular periods' within the last 12 months. From 2009
9 onwards, information on their children was collected, including birth dates and preterm birth
10 (<37 weeks). Logistic regression using generalised estimating equations was used to examine
11 prospective associations between self-reported menstrual symptoms before pregnancy and
12 risk of preterm birth.

13 **Results:** Data from 6,615 mothers, who had 12,337 live singleton births were available for
14 analysis. Among all births, women reporting severe period pain [adjusted odds ratio (aOR)
15 1.34, 95% CI 1.10–1.62] or heavy periods (aOR 1.25, 95% CI 1.02–1.53) before pregnancy
16 had higher odds of preterm birth. However, in analyses stratified by birth order, only severe
17 period pain (aOR 2.05, 95% CI 1.41–2.99), heavy periods (aOR 1.77, 95% CI 1.23–2.55) or
18 irregular periods (aOR 1.58, 95% CI 1.10–2.28) prior to a second or subsequent birth were
19 associated with an increased risk of preterm birth.

20 **Conclusions:** Severe period pain, heavy periods and irregular periods before a second or
21 subsequent birth may be associated with preterm birth.

22

23 **Keywords** preterm birth, dysmenorrhea, heavy menstrual bleeding, irregular periods, cohort
24 study

25

26 **Background**

27 Preterm birth, defined as birth prior to 37 weeks, occurs in 5-10% of all pregnancies and the
28 prevalence has increased in high-resource countries¹⁻³. Preterm birth is a major cause of
29 neonatal mortality and child morbidity, including neurological, respiratory and cardiovascular
30 disorders⁴. Preventative strategies are challenging because there is uncertainty about the
31 underlying causes of preterm birth⁵. For approximately half of mothers, there will be no
32 identifiable cause for these conditions^{1,5,6}. However, commonly reported maternal risk
33 factors for preterm birth include very young and advanced maternal age, infections, medical
34 conditions before and during pregnancy (e.g., hypertension, diabetes), and smoking^{1,6,7}.

35

36 Hormonal and inflammatory influences before and during pregnancy also play important
37 roles in adverse pregnancy and neonatal outcomes⁸. Menstrual cycle characteristics, including
38 dysmenorrhea and irregular periods, can be important indicators of hormonal imbalances,
39 inflammation and of a woman's risk of future health problems^{9,10}. Approximately one
40 quarter of women report dysmenorrhea,¹¹ and a third report irregular periods⁹. Although
41 menstrual symptoms can decrease across the life course, some women will experience
42 chronic symptoms throughout their reproductive years¹¹. Risk factors for chronic
43 dysmenorrhea include smoking and obesity¹¹, which are also risk factors for adverse
44 pregnancy outcomes¹².

45

46 Studies examining how menstrual symptoms influence birth outcomes are scant. Two small,
47 case-control studies from Finland and Taiwan reported an association between dysmenorrhea
48 and preterm birth^{13,14}. However, the study from Finland was conducted more than three
49 decades ago, relied on women's retrospective reports of menstrual symptoms, and did not
50 consider other risk factors. The more recent study from Taiwan used medical records to

51 confirm women's birth outcomes, but dysmenorrhea was also collected retrospectively at the
52 first prenatal visit. Data from larger, more comprehensive studies, that can examine
53 prospective associations are important for identifying pre-pregnancy risk factors that can
54 inform timely preventative interventions for preterm birth.

55

56 In this paper, we use data from 12,337 singleton births from 6,615 women who participated
57 in a longitudinal, population-based cohort study to examine the prospective associations
58 between menstrual symptoms; we also account for the mother's physical, mental and
59 reproductive health history.

60

61 **Methods**

62 The Australian Longitudinal Study on Women's Health (ALSWH) is a national study
63 focusing on the biological, psychological, social and economic factors relevant to women's
64 health¹⁵. The women were randomly selected from the database of Medicare (Australia's
65 universal health insurance system), with oversampling from rural and remote areas of
66 Australia. Approximately every three years since 1996, participants have completed postal or
67 online surveys about their health and wellbeing^{15,16}. Detailed information on the study
68 methods has been published elsewhere^{16,17} and is available at <http://www.alswh.org.au/>.

69 *Participants*

70 A total of 14,247 young women participated in the baseline survey in 1996 when they were
71 aged 18-23 years. Of these, 8,349 women had 18,525 live births between 1996 and 2015 (see
72 Figure 1). Overall, we excluded 6,188 (33%) births and 1,734 (21%) mothers due to births
73 that occurred before the baseline survey in 1996 (1,102 births from 170 mothers); multiple
74 births (628 births from 95 mothers) and births from mothers who had missing data for

75 menstrual symptoms (2,184 births), the first birth (1,193 births from 743 mothers) or preterm
76 birth (1,081 births from 726 mothers). A final sample of 12,337 live singleton births from
77 6,615 mothers were available for analysis (see Figure 1). Of these, 4,454 (67.3%) mothers
78 had two or more singleton births by Survey 7 when they were aged 37-42 years.

79 *Measures*

80 Women were asked about their birth history at Survey 5 (2009), Survey 6 (2012), and Survey
81 7 (2015). For each birth, participants were asked to report the date of birth for each child and
82 if they had experienced: “Premature birth”. Using each infant’s date of birth and the date
83 when the mother returned each survey, births were matched to the mother’s survey before the
84 birth. This information was also used to ascertain maternal age at birth (calculated using the
85 mother’s date of birth at the time of the child’s date of birth), birth order (first birth vs.
86 subsequent birth) and history of preterm birth (dichotomised as yes vs. no).

87

88 Women were also asked about their reproductive health including menstrual symptoms. At
89 each survey, three separate questions were asked: “In the last 12 months, have you had: 1)
90 severe period pain; 2) heavy periods; 3) irregular periods?”. Separate variables were created
91 for each question; those who answered “sometimes” or “often” were coded as having that
92 symptom, and those who responded “never” or “rarely” were coded as not having the
93 symptom. Other aspects of reproductive health including a history of miscarriage, doctor
94 diagnosis of polycystic ovary syndrome (data collected from Survey 4 onwards) and
95 endometriosis (all dichotomised as yes vs no) were also collected.

96

97 Sociodemographic information was also collected at each survey including area of residence
98 based on an index of distance to the nearest urban centre (urban vs. rural/remote)¹⁸; highest

99 level of education (year 12 or less, trade/certificate/diploma, university degree) and current
100 relationship status (single, married/defacto partnership). Women were also asked about
101 smoking (classified as never, ex-smoker, current smoker) and body mass index [(BMI;kg/m²)
102 calculated using women's self-reported weight and height, and categorised as underweight
103 (<18.5kg/m²), normal weight (18.5-24.9 kg/m²), overweight or obese (≥25 kg/m²).
104 Information was collected on chronic health conditions prior to pregnancy including
105 hypertension or diabetes (dichotomised as yes vs. no), general mental health (using the 5-item
106 Mental Health subscale (MHI-5) of the SF-36) ¹⁹.

107 *Statistical analyses*

108 Chi-square tests and independent samples t-tests were used to compare the characteristics of
109 nulliparous women, up to three years prior to their first birth, who subsequently did, or did
110 not experience preterm birth. Using each mother's longitudinal birth outcome data, logistic
111 regression models using generalised estimating equations with exchangeable working
112 correlation matrices and robust error variances, were used to account for repeated births
113 nested within mothers. Separate models assessed the association between each menstrual
114 symptom and preterm birth.

115

116 All covariates were those reported by the woman at the survey before the index birth, except
117 maternal age, which was based on the participant's age at the time of the birth. All covariates
118 were initially included in the models together to assess associations with birth outcomes.

119 Area of residence, education, smoking, mental health, miscarriage, polycystic ovary
120 syndrome and endometriosis were not statistically significant ($p > 0.05$) or did not change
121 odds ratio estimates by more than 10%, and thus, were not included in the final models. Final
122 models were adjusted for maternal age at birth, BMI, hypertension or diabetes (combined into

123 one variable for the analysis), birth order and previous preterm birth. To account for the
124 possibility that women's menstrual symptoms changed between births, an interaction term
125 between each menstrual symptom and birth order (first birth vs. subsequent birth) was
126 included in the models and further analyses stratifying by birth order were completed.

127

128 There were moderate levels of missing data on the key maternal exposures (severe period
129 pain = 12.1%; heavy periods = 12.2%; irregular periods = 12.2%) and covariates
130 (hypertension = 14.8%; diabetes = 14.8%; BMI = 20.5%). The influence of missing data on
131 the exposures and covariates on the results was assessed in a multiple imputation analysis²⁰.

132 The variables used in the imputation procedure included the outcome variable, exposures
133 (menstrual symptoms) and covariates (maternal age, education, area of residence, marital
134 status, parity, age at menarche, smoking, BMI, hypertension, diabetes). The following SAS
135 procedures were performed: PROC MI using fully conditional specification (due to the binary
136 nature of the key exposures and outcomes) was used to create 20 imputed datasets; logistic
137 regression analyses using generalised estimating equations were performed on each of the
138 imputed datasets; and results were combined using MIANALYZE. Findings based on
139 original data are presented, as these did not differ from findings based on the imputed data.
140 All statistical analyses were performed using SAS software version 9.4 (TS1M5)²¹.

141 **Results**

142 A total of 12,337 live singleton neonates born between 1996 and 2015 were eligible for
143 analysis. Of these births, 6.3% (n = 779) were preterm. The percentage of preterm births was
144 8.5% (564/6,615) in a first birth and 3.8% (215/5722) in a second or subsequent birth.

145 Table 1 shows the characteristics of the 6,615 women who gave birth to the 12,337 children
146 between 1996 and 2015. Women who went on to have a preterm birth were more likely to be
147 younger, residing in a rural or remote area, married and to have completed 12 years or less

148 years of education at the survey prior to their first birth than women who did not have a
149 preterm birth (see Table 1). They were more likely to be overweight or obese, current
150 smokers, and to report a diagnosis of hypertension or diabetes. Women who went on to have
151 a preterm birth were more likely to have a history of miscarriage, a diagnosis of polycystic
152 ovary syndrome or endometriosis and poorer mental health prior to their first birth than
153 women who did not have a preterm birth (see Table 1).

154

155 Menstrual symptoms before pregnancy among women who did, or did not, have a preterm
156 birth are presented in Table 2. Before a first birth, severe period pain did not appear to differ
157 among women who did, and did not, have a preterm birth. Specifically, 30.0% of nulliparous
158 women who went on to have preterm birth reported severe period pain. By comparison,
159 26.3% of nulliparous women who did not have a preterm birth reported severe period pain.
160 However, menstrual symptoms differed among women who did, and did not, have a
161 subsequent preterm birth. For subsequent births, 24.9% of women who had a preterm birth
162 reported severe period pain compared to 13.4% who did not have a preterm birth ($p < 0.001$).
163 This pattern was similar for the other menstrual symptoms. Specifically, heavy periods and
164 irregular periods before pregnancy did not differ among women whose first birth was, or was
165 not, preterm. For subsequent births, 25.6% of women who had a preterm birth reported heavy
166 periods compared to 16.1% among women who did not have a preterm birth ($p < 0.001$).
167 Similarly, 25.8% of women who had a subsequent preterm birth reported irregular periods
168 compared to 16.6% who did not have a subsequent preterm birth ($p < 0.001$).

169 Longitudinal multivariable models were constructed to examine the associations between
170 each menstrual symptom and preterm birth. For all births, severe period pain before
171 pregnancy was associated with higher odds of preterm birth (OR 1.58, 95% CI 1.33–1.88),
172 and this association remained after adjustment for other key covariates [adjusted OR (aOR)

173 1.33, 95% CI 1.10–1.62] (Table 3). However, there was a significant interaction between
174 birth order and severe period pain (p for interaction = 0.01) suggesting that birth order
175 modified the association between severe period pain and preterm birth. In analyses stratified
176 by birth order, severe period pain before a first birth was not associated with preterm birth.
177 Only women reporting severe period pain before a second or subsequent birth had higher
178 odds of preterm birth (aOR 2.05, 95% CI 1.41–2.99). These results were similar for heavy
179 periods with an increased risk of preterm birth among all births (aOR 1.24, 1.01–1.53) and
180 for women reporting heavy periods before a second or subsequent birth (aOR 1.78, 1.24–
181 2.56). Although there was no overall association between irregular periods and preterm birth,
182 there was a significant interaction between birth order and irregular periods (p for interaction
183 = 0.01). In analyses stratified by birth order, only women reporting irregular periods prior to a
184 second or subsequent birth had higher odds of preterm birth (aOR 1.59, 1.10–2.29).

185 **Discussion**

186 Using a large, population-based cohort study we examined the associations between
187 menstrual symptoms and preterm birth. In this study, the strongest associations between
188 menstrual symptoms and preterm birth were found after a first birth. Specifically, the chances
189 of preterm birth were greater among women reporting severe period pain, heavy periods and
190 irregular periods before a second or greater birth.

191

192 Underlying maternal medical conditions may play an important role in explaining our
193 findings. Although there is evidence to suggest that dysmenorrhea is less likely after birth²²,
194 any benefits attributable to a recent birth may be short term for women who have underlying
195 conditions. Such underlying conditions may damage the uterine environment, manifesting as
196 menstrual problems and increased risk of poor subsequent birth outcomes. Endometriosis is a

197 common cause of severe period pain, heavy periods, and irregular periods^{10,22}, and has been
198 previously associated with preterm birth²³. A recent systematic review found limited evidence
199 to suggest that symptoms of endometriosis improve following pregnancy²⁴. The persistence
200 of menstrual problems beyond a first birth may therefore identify women who have more
201 severe symptoms and may be a marker for underlying conditions and future health problems.
202 During pregnancy, the hormonal and inflammatory changes associated with endometriosis
203 may negatively affect the uterine environment²⁵, or contribute to complications in pregnancy
204 (e.g., placenta praevia)²⁶, increasing the chances of preterm birth. Because considerable
205 delays in diagnosis are common for women who have endometriosis²⁷, some women with
206 severe period pain and heavy menstrual bleeding in our study may have had undiagnosed
207 disease. As the disease progresses over time, this may lead to further scarring and adhesions
208 that increase the risk of problems in subsequent pregnancies. However, adjusting for a history
209 of endometriosis did not change the association between menstrual symptoms and preterm
210 birth in this study.

211

212 Menstrual problems may also be an indicator of other underlying hormonal, endocrine, or
213 metabolic conditions that increase the risk of pregnancy complications. In this study, heavy
214 periods or irregular periods before a second or greater pregnancy increased the odds of
215 preterm birth. Heavy and irregular periods are a common feature of polycystic ovary
216 syndrome²⁸, which is also related to a range of adverse pregnancy outcomes including
217 preterm birth^{29,30}. However, in this study, adjusting for polycystic ovary syndrome did not
218 appreciably change our findings. Alternatively, heavy periods have been associated with
219 problems in the production of progesterone³¹. Because progesterone plays a key role in the
220 maintenance of pregnancy, insufficient levels of progesterone in women may increase the
221 risk of preterm birth³². Clinically, progesterone is used to prevent preterm birth in women

222 who have premature cervical shortening in pregnancy, however, the efficacy of the treatment
223 has been contentious³³⁻³⁵. Chronic conditions associated with increasing age (e.g., obesity,
224 diabetes) can also disrupt the production of progesterone and lead to heavy periods³¹;
225 however, these conditions and maternal age were taken into account in our analyses. Other
226 gynaecological conditions (e.g., uterine fibroids, adenomyosis)³¹, which were not assessed in
227 this study, might help to explain the association between menstrual problems and preterm
228 birth.

229

230 Interventions during a first birth may further help to explain our findings. Recent evidence
231 suggests that having had a caesarean birth is a risk factor for preterm birth among
232 multiparous women³⁶. In Australia, caesarean birth rates are high (33% of all births in 2015)
233 and have increased over time³⁷. In another study also based on data from the young women
234 participating in ALSWH, 29% of first, singleton births were caesarean births, which is
235 consistent with national estimates³⁸. The risks of caesarean birth are well-established and
236 increase with each subsequent caesarean birth³⁹. Adhesions, surgical injury or infections
237 following a caesarean birth⁴⁰ may change women's menstrual symptoms following a first
238 birth. Uterine changes associated with caesarean birth may help to explain the increased odds
239 of preterm birth in second and greater births in this study.

240

241 There may also be complex interactions between hormonal, metabolic, and inflammatory
242 pathways that influence a woman's uterine environment after a first pregnancy, and
243 negatively affect their future pregnancy outcomes⁸. The composition of the vaginal
244 microbiome may help to explain how hormonal and inflammatory pathways interact to
245 influence menstrual symptoms and the risk of preterm birth, particularly in subsequent

246 pregnancies. Bacterial vaginosis⁴¹, and more recently certain species in the vaginal
247 microbiome^{42,43}, have been linked to preterm birth. There is evidence to suggest that the
248 composition of the vaginal microbiome changes during pregnancy and differs from the
249 composition in the postpartum period, which may be due to the significant hormonal changes
250 during these periods⁴⁴. Menstrual symptoms are often important markers for hormonal
251 imbalances in women⁸ and may also be linked to changes in the vaginal microbiome.

252

253 Hypothetically, imbalances in vaginal microbiota following a first birth may give rise to
254 inflammatory processes⁴² and present as menstrual problems, thereby increasing adverse
255 outcomes in subsequent pregnancies. During pregnancy, the up-regulation of pro-
256 inflammatory responses among women with an abundance of certain bacterial species may
257 trigger the release of prostaglandins, or erode mucosal barriers that normally protect women
258 from infection, and increase the risk of premature labour⁴⁵. If vaginal microbiota were an
259 underlying mechanism, women presenting with a history of menstrual problems following a
260 first birth might benefit from treatments that modify the vaginal microbiome in their next
261 pregnancy. Although oral probiotic supplements to modify the vaginal microbiome are
262 potentially a low-risk intervention that can be leveraged to reduce preterm birth rates, their
263 effectiveness has yet to be demonstrated⁴⁶. The associations observed in this study, and the
264 role of the vaginal microbiome in preterm birth, warrants further investigation.

265 The main advantage of this longitudinal, population-based cohort study is that prospective
266 relationships between menstrual symptoms and subsequent preterm birth were examined. We
267 were also able to assess the contribution of an extensive range of covariates including
268 sociodemographic factors, and physical, psychological and reproductive health. Although
269 some women were lost to follow-up or withdrew over the 20-year study period, continued
270 participation has remained relatively stable at more than 80%⁴⁷. Women who remain in this

271 study are broadly representative of the general population of Australian women in this age
272 group with a few exceptions. A slightly higher proportion of women who remain in the study
273 are married or in de facto relationships and have university degrees ^{16,17}.

274

275 This study has some important limitations. All information was based on self-report
276 questionnaires rather than hospital records, which may mean that preterm birth, menstrual
277 symptoms and other covariates assessed in this study were over- or under-reported. Another
278 study comparing self-reported perinatal outcomes in ALSWH to hospital records
279 demonstrated high agreement between the two sources ⁴⁸. However, because ALSWH is a
280 broad-ranging health study not specific to birth outcomes, information was not available to
281 distinguish between early and late preterm births or to determine whether the preterm birth
282 was spontaneous or medically indicated. The timing of the ALSWH surveys also means that
283 information on women's menstrual symptoms and the other key covariates may have
284 occurred three years prior to the index birth. In addition, infants born to women who had two
285 or more pregnancies between successive surveys were excluded as menstrual symptom data
286 before that birth were missing. Therefore, we are unable to establish whether a causal
287 association exists between menstrual symptoms and preterm birth.

288

289 Overall, our findings suggest that menstrual problems that continue or emerge after a first
290 birth are risk factors for preterm birth. These findings emphasise the potential clinical
291 importance of reviewing women's menstrual symptoms in early pregnancy, especially among
292 women in a second or subsequent pregnancy. In Australia, there have been recent efforts to
293 reduce preterm births rates through increased health provider education and the
294 implementation of evidenced-based recommendations⁴⁹. These recommendations reinforce
295 the need to measure cervical length at all mid-pregnancy morphology scans and to provide

296 vaginal progesterone pessaries for women at risk based on their cervical length or maternal
297 history⁴⁹. Currently, Australian health providers should counsel women based on these
298 recommendations. In the future, women's menstrual symptom history could be used in
299 conjunction with other aspects of maternal medical history to better identify women at risk of
300 preterm birth. However, future research is required to confirm the association between
301 menstrual symptoms and women's birth outcomes.

302

303 **References**

- 304 1. Goldenberg RL, Culhane JF, Iams JD, Romero R. Epidemiology and causes of
305 preterm birth. *The lancet*. 2008;371(9606):75-84.
- 306 2. Beck S, Wojdyla D, Say L, et al. The worldwide incidence of preterm birth: a
307 systematic review of maternal mortality and morbidity. *Bulletin of the World Health*
308 *Organization*. 2010;88(1):31-38.
- 309 3. Ananth CV, Goldenberg RL, Friedman AM, Vintzileos AM. Association of Temporal
310 Changes in Gestational Age With Perinatal Mortality in the United States, 2007-2015.
311 *JAMA Pediatr*. 2018;172(7):627-634.
- 312 4. Lawn JE, Cousens S, Zupan J, Team LNSS. 4 million neonatal deaths: when? Where?
313 Why? *The lancet*. 2005;365(9462):891-900.
- 314 5. Challis JR, Lye SJ, Gibb W, Whittle W, Patel F, Alfaidy N. Understanding preterm
315 labor. *Annals of the New York Academy of Sciences*. 2001;943(1):225-234.
- 316 6. Slattery MM, Morrison JJ. Preterm delivery. *The Lancet*. 2002;360(9344):1489-1497.
- 317 7. Valero de Bernabé J, Soriano T, Albaladejo R, et al. Risk factors for low birth weight:
318 a review. *European Journal of Obstetrics & Gynecology and Reproductive Biology*.
319 2004;116(1):3-15.
- 320 8. Vannuccini S, Clifton VL, Fraser IS, et al. Infertility and reproductive disorders:
321 impact of hormonal and inflammatory mechanisms on pregnancy outcome. *Human*
322 *Reproduction Update*. 2016;22(1):104-115.
- 323 9. West S, Lashen H, Bloigu A, et al. Irregular menstruation and hyperandrogenaemia in
324 adolescence are associated with polycystic ovary syndrome and infertility in later life:
325 Northern Finland Birth Cohort 1986 study. *Human reproduction (Oxford, England)*.
326 2014;29(10):2339-2351.
- 327 10. Parazzini F, Esposito G, Tozzi L, Noli S, Bianchi S. Epidemiology of endometriosis
328 and its comorbidities. *Eur J Obstet Gynecol Reprod Biol*. 2017;209:3-7.
- 329 11. Ju H, Jones M, Mishra GD. Premenstrual syndrome and dysmenorrhea: symptom
330 trajectories over 13 years in young adults. *Maturitas*. 2014;78(2):99-105.
- 331 12. Verburg PE, Dekker GA, Venugopal K, et al. Long-term trends in singleton preterm
332 birth in South Australia from 1986 to 2014. *Obstetrics & Gynecology*.
333 2018;131(1):79-89.
- 334 13. Juang C-M, Chou P, Yen M-S, Twu N-F, Horng H-C, Hsu W-L. Primary
335 dysmenorrhea and risk of preterm delivery. *American journal of perinatology*.
336 2007;24(1):11-16.
- 337 14. Ylikorkala O, Kujansuu E. Increased rate of primary dysmenorrhea in women with
338 spontaneous premature labor. *Prostaglandins and medicine*. 1981;6(2):213-216.
- 339 15. Lee C, Dobson AJ, Brown WJ, et al. Cohort Profile: the Australian Longitudinal
340 Study on Women's Health. *International journal of epidemiology*. 2005;34(5):987-
341 991.
- 342 16. Dobson AJ, Hockey R, Brown WJ, et al. Cohort Profile Update: Australian
343 Longitudinal Study on Women's Health. *International journal of epidemiology*. 2015.
- 344 17. Brown WJ, Bryson L, Byles JE, et al. Women's Health Australia: recruitment for a
345 national longitudinal cohort study. *Women & health*. 1999;28(1):23-40.
- 346 18. Australian Institute of Health and Welfare (AIHW). *Rural, regional and remote*
347 *health: a guide to remoteness classifications*. Canberra: AIHW;2004.
- 348 19. McCallum J. The SF-36 in an Australian sample: validating a new, generic health
349 status measure. *Australian Journal of Public Health*. 1995;19(2):160-166.
- 350 20. Berglund PA. *An introduction to multiple imputation of complex sample data using*
351 *SAS v9.2*. Cary, NC: SAS Institute Inc;2010.

- 352 21. SAS Institute Inc. Cary, NC, USA2016.
- 353 22. Ju H, Jones M, Mishra G. The prevalence and risk factors of dysmenorrhea.
354 *Epidemiologic reviews*. 2013;mxt009.
- 355 23. Kim SG, Seo HG, Kim YS. Primiparous singleton women with endometriosis have an
356 increased risk of preterm birth: Meta-analyses. *Obstet Gynecol Sci*. 2017;60(3):283-
357 288.
- 358 24. Leeners B, Damaso F, Ochsenein-Kolble N, Farquhar C. The effect of pregnancy on
359 endometriosis-facts or fiction? *Hum Reprod Update*. 2018;24(3):290-299.
- 360 25. Leone Roberti Maggiore U, Ferrero S, Mangili G, et al. A systematic review on
361 endometriosis during pregnancy: diagnosis, misdiagnosis, complications and
362 outcomes. *Hum Reprod Update*. 2016;22(1):70-103.
- 363 26. Zullo F, Spagnolo E, Saccone G, et al. Endometriosis and obstetrics complications: a
364 systematic review and meta-analysis. *Fertil Steril*. 2017;108(4):667-672 e665.
- 365 27. Ballard KD, Seaman HE, de Vries CS, Wright JT. Can symptomatology help in the
366 diagnosis of endometriosis? Findings from a national case-control study--Part 1.
367 *BJOG*. 2008;115(11):1382-1391.
- 368 28. Teede H, Deeks A, Moran L. Polycystic ovary syndrome: a complex condition with
369 psychological, reproductive and metabolic manifestations that impacts on health
370 across the lifespan. *BMC Med*. 2010;8:41.
- 371 29. Boomsma CM, Fauser BC, Macklon NS. Pregnancy complications in women with
372 polycystic ovary syndrome. *Semin Reprod Med*. 2008;26(1):72-84.
- 373 30. Palomba S, de Wilde MA, Falbo A, Koster MP, La Sala GB, Fauser BC. Pregnancy
374 complications in women with polycystic ovary syndrome. *Hum Reprod Update*.
375 2015;21(5):575-592.
- 376 31. Hapangama DK, Bulmer JN. Pathophysiology of heavy menstrual bleeding. *Womens
377 Health (Lond)*. 2016;12(1):3-13.
- 378 32. Romero R, Dey SK, Fisher SJ. Preterm labor: one syndrome, many causes. *Science*.
379 2014;345(6198):760-765.
- 380 33. Eichelberger KY, Manuck TA. Progesterone has no place in the prevention of preterm
381 delivery: AGAINST: A call for a measured response to the OPPTIMUM trial. *BJOG:
382 An International Journal of Obstetrics & Gynaecology*. 2016;123(9):1511-1511.
- 383 34. Campbell S. Prevention of spontaneous preterm birth: universal cervical length
384 assessment and vaginal progesterone in women with a short cervix: time for action!
385 *American Journal of Obstetrics and Gynecology*. 2018;218(2):151-158.
- 386 35. Norman JE, Marlow N, Messow CM, et al. Vaginal progesterone prophylaxis for
387 preterm birth (the OPPTIMUM study): a multicentre, randomised, double-blind trial.
388 *Lancet (London, England)*. 2016;387(10033):2106-2116.
- 389 36. Jelliffe-Pawlowski LL, Baer RJ, Blumenfeld YJ, et al. Maternal characteristics and
390 mid-pregnancy serum biomarkers as risk factors for subtypes of preterm birth. *BJOG
391 : an international journal of obstetrics and gynaecology*. 2015;122(11):1484-1493.
- 392 37. Australian Institute of Health and Welfare 2017 (AIHW). *Australia's mothers and
393 babies 2015—in brief. Perinatal statistics series no. 33. Cat no. PER 91*. Canberra:
394 AIHW.
- 395 38. Hure A, Powers J, Chojenta C, Loxton D. Rates and Predictors of Caesarean Section
396 for First and Second Births: A Prospective Cohort of Australian Women. *Maternal
397 and Child Health Journal*. 2017;21(5):1175-1184.
- 398 39. Huang X, Lei J, Tan H, Walker M, Zhou J, Wen SW. Cesarean delivery for first
399 pregnancy and neonatal morbidity and mortality in second pregnancy. *Eur J Obstet
400 Gynecol Reprod Biol*. 2011;158(2):204-208.

- 401 40. Marshall NE, Fu R, Guise J-M. Impact of multiple cesarean deliveries on maternal
402 morbidity: a systematic review. *American Journal of Obstetrics and Gynecology*.
403 2011;205(3):262.e261-262.e268.
- 404 41. Donders GG, Van Calsteren K, Bellen G, et al. Predictive value for preterm birth of
405 abnormal vaginal flora, bacterial vaginosis and aerobic vaginitis during the first
406 trimester of pregnancy. *BJOG : an international journal of obstetrics and*
407 *gynaecology*. 2009;116(10):1315-1324.
- 408 42. Fettweis JM, Serrano MG, Brooks JP, et al. The vaginal microbiome and preterm
409 birth. *Nature Medicine*. 2019;25(6):1012-1021.
- 410 43. Tabatabaei N, Eren A, Barreiro L, et al. Vaginal microbiome in early pregnancy and
411 subsequent risk of spontaneous preterm birth: a case-control study. *BJOG : an*
412 *international journal of obstetrics and gynaecology*. 2019;126(3):349-358.
- 413 44. MacIntyre DA, Chandiramani M, Lee YS, et al. The vaginal microbiome during
414 pregnancy and the postpartum period in a European population. *Scientific Reports*.
415 2015;5(1):8988.
- 416 45. Kindinger LM, Bennett PR, Lee YS, et al. The interaction between vaginal
417 microbiota, cervical length, and vaginal progesterone treatment for preterm birth risk.
418 *Microbiome*. 2017;5(1):6.
- 419 46. Husain S, Allotey J, Drymoussi Z, et al. Effects of oral probiotic supplements on
420 vaginal microbiota during pregnancy: a randomised, double-blind, placebo-controlled
421 trial with microbiome analysis. *BJOG : an international journal of obstetrics and*
422 *gynaecology*. 2020;127(2):275-284.
- 423 47. Powers J, Loxton D. The impact of attrition in an 11-year prospective longitudinal
424 study of younger women. *Annals of epidemiology*. 2010;20(4):318-321.
- 425 48. Gresham E, Forder P, Chojenta CL, Byles JE, Loxton DJ, Hure AJ. Agreement
426 between self-reported perinatal outcomes and administrative data in New South
427 Wales, Australia. *BMC Pregnancy & Childbirth*. 2015;15(1):1.
- 428 49. Keelan JA, Newnham JP. Recent advances in the prevention of preterm birth.
429 *F1000Res*. 2017;6:F1000 Faculty Rev-1139.
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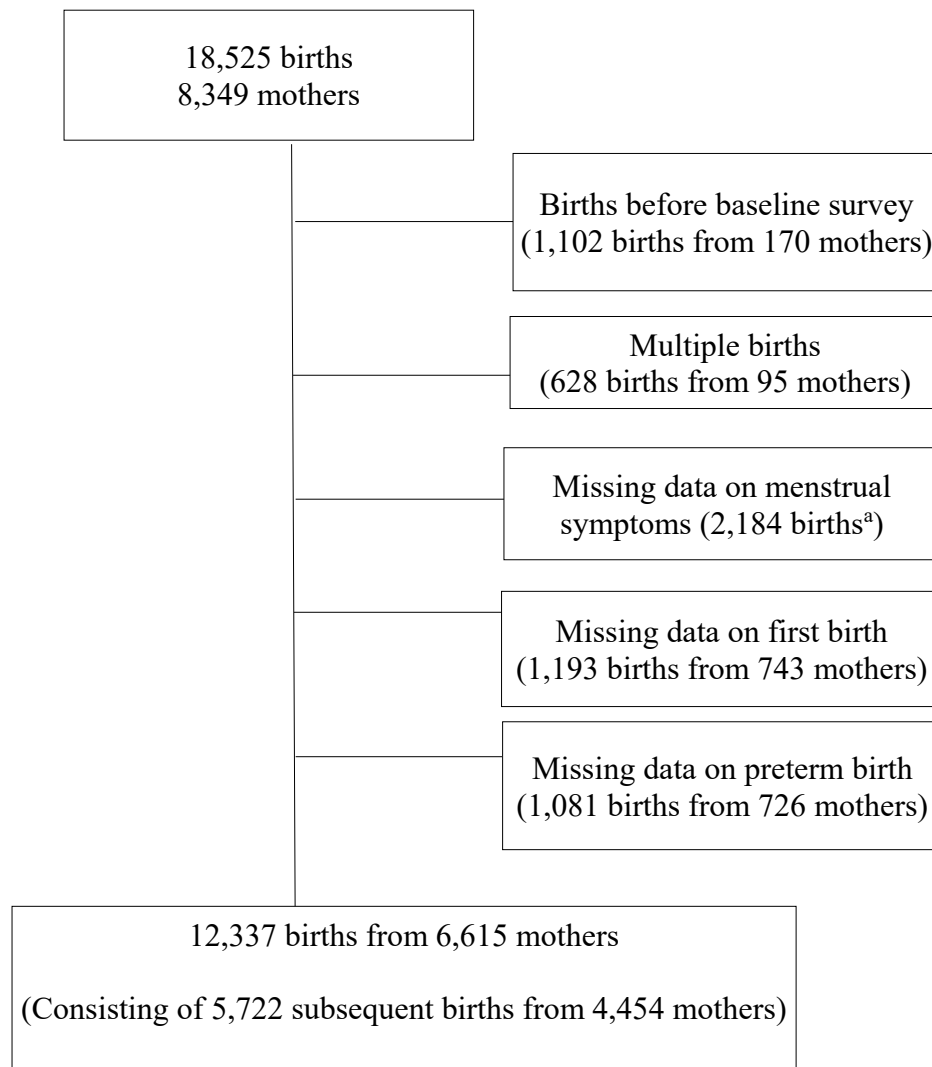
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460 ^aData from births were only missing because the mother had one record of menstrual

461 symptoms but two births between surveys.

462



463 **Figure legends**

464 Figure 1. Flow diagram of the births between 1996 and 2015 from Australian mothers who
465 met the inclusion criteria.

466 Table 1. Characteristics of Australian women prior to the first birth who subsequently did, or
 467 did not, have a preterm birth between 1996-2015. N = 6615

	No Preterm birth <i>n</i> = 5887	Preterm birth <i>n</i> = 728
	<i>n</i> (%)	<i>n</i> (%)
Age (years)***		
18-24	1179 (22.9)	168 (26.9)
25-29	2170 (42.2)	289 (46.3)
30-34	1438 (28.0)	134 (21.5)
35-40	356 (6.9)	33 (5.3)
Area of residence*		
Urban	2926 (57.8)	330 (53.6)
Rural/Remote	2133 (42.2)	286 (46.4)
Marital status**		
Single	973 (19.0)	148 (23.8)
Married/De facto	4157 (81.0)	473 (76.2)
Education**		
Year 12 or less	1343 (26.5)	193 (31.3)
Trade/certificate/diploma	1213 (23.9)	154 (25.0)
University	2516 (49.6)	269 (43.7)
Body mass index**		
Underweight	194 (4.3)	29 (5.2)

	No Preterm birth	Preterm birth
	<i>n</i> = 5887	<i>n</i> = 728
	<i>n</i> (%)	<i>n</i> (%)
Healthy weight	2900 (64.2)	316 (56.8)
Overweight	950 (21.0)	135 (24.3)
Obese	472 (10.5)	76 (13.7)
Smoking**		
Never smoked	3069 (60.3)	342 (54.9)
Ex-smoker	1109 (21.8)	139 (22.3)
Current smoker	914 (17.9)	142 (22.8)
Diabetes ^{a**}		
No	4961 (99.4)	599 (98.4)
Yes	31 (0.6)	10 (1.6)
Hypertension ^{a***}		
No	4870 (97.6)	561 (92.1)
Yes	121 (2.4)	48 (7.9)
Miscarriage**		
No	4588 (90.1)	535 (86.7)
Yes	503 (9.9)	82 (13.3)
Polycystic ovary syndrome ^{***}		
No	5352 (92.6)	621 (87.8)

	No Preterm birth	Preterm birth
	<i>n</i> = 5887	<i>n</i> = 728
	<i>n</i> (%)	<i>n</i> (%)
Yes	429 (7.4)	86 (12.2)
Endometriosis**		
No	4673 (95.8)	557 (93.6)
Yes	203 (4.2)	38 (6.4)
Mental health score (MHI-5)***	Mean ±SD	Mean ±SD
	73.1 ±16.1	70.2 ±16.7

468 *Note.* Data were reported up to three years prior to the first birth.

469 ^aPrior to pregnancy

470 SD, standard deviation.

471 **P* = <.05; ***P* = <.01; ****P* = <.001

472 Table 2. Menstrual symptoms before pregnancy among Australian women who subsequently
 473 did, or did not, have a preterm birth between 1996-2015, by birth order. N = 6615

	No Preterm birth	Preterm birth
	<i>n</i> (%)	<i>n</i> (%)
Severe period pain before first birth		
No	3867 (73.7)	340 (70.0)
Yes	1382 (26.3)	146 (30.0)
Severe period pain before subsequent birth***		
No	4268 (86.6)	133 (75.1)
Yes	659 (13.4)	44 (24.9)
Heavy periods before first birth		
No	4271 (81.5)	385 (79.2)
Yes	972 (18.5)	101 (20.8)
Heavy periods before subsequent birth***		
No	4131 (83.9)	131 (74.4)
Yes	794 (16.1)	45 (25.6)
Irregular periods before first birth		
No	4198 (80.0)	391 (80.5)
Yes	1049 (20.0)	95 (19.6)
Irregular periods before subsequent birth***		
No	4107 (83.4)	132 (74.2)

	No Preterm birth	Preterm birth
	<i>n</i> (%)	<i>n</i> (%)
Yes	819 (16.6)	46 (25.8)

474 *Note.* Data were reported up to three years prior to the birth.

475 Subsequent birth = Refers to a second or greater birth.

476 * $P = <.05$; ** $P = <.01$; *** $P = <.001$

477 Table 3. Association between menstrual symptoms reported by Australian women and
 478 preterm birth between 1996-2015. N = 12,337

	Unadjusted	Adjusted
	OR (95% CI)	OR (95% CI)
Severe period pain (all births)	1.58 (1.33–1.88)	1.34 (1.10–1.62) ^a
Stratified by birth order		
First birth	1.20 (0.98–1.47)	1.18 (0.95–1.48) ^b
Subsequent birth	2.01 (1.40–2.88)	2.05 (1.41–2.99) ^c
Heavy periods (all births)	1.34 (1.11–1.61)	1.25 (1.02–1.53) ^a
Stratified by birth order		
First birth	1.15 (0.92–1.45)	1.08 (0.84–1.39) ^b
Subsequent birth	1.65 (1.16–2.35)	1.77 (1.23–2.55) ^c
Irregular periods (all births)	1.17 (0.96–1.41)	1.05 (0.85–1.30) ^a
Stratified by birth order		
First birth	0.97 (0.77–1.23)	0.89 (0.69–1.15) ^b
Subsequent birth	1.67 (1.19–2.33)	1.58 (1.10–2.28) ^c

479 *Note.* Menstrual symptoms refer to those preceding the index pregnancy (up to 3 years before
 480 birth). Subsequent births are second or greater births.

481 ^aAdjusted for maternal age, BMI, hypertension/diabetes, parity and previous preterm birth

482 ^bAdjusted for maternal age, BMI, hypertension/diabetes

483 ^cAdjusted for maternal age, BMI, hypertension/diabetes and previous preterm birth