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Taim, BC, Ó Catháin, C, Renard, M, Elliott-Sale, Kirsty , Madigan, S and Ní Chéilleachair, N (2023) The Prevalence of Menstrual Cycle Disorders and Menstrual Cycle-Related Symptoms in Female Athletes: A Systematic Literature Review. *Sports Medicine*, 53. pp. 1963-1984. ISSN 0112-1642

DOI: <https://doi.org/10.1007/s40279-023-01871-8>

Publisher: Springer

Version: Accepted Version

Downloaded from: <https://e-space.mmu.ac.uk/632262/>

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Data Access Statement: The data analysed during the current study are available from the corresponding author upon reasonable request.

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PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

1 **Title:** The Prevalence of Menstrual Cycle Disorders and Menstrual Cycle-Related Symptoms in Female
2 Athletes: A Systematic Literature Review

3

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25

26 **Abstract**

27 **Background:** Menstrual cycle (MC) disorders and MC-related symptoms can have debilitating effects
28 on the health and performance of female athletes. As the participation of women in sports continues to
29 increase, understanding the prevalence of a range of MC disorders and MC-related symptoms may
30 guide preventive strategies to protect the health and optimise the performance of female athletes.

31
32 **Objective:** To examine the prevalence of MC disorders and MC-related symptoms among female
33 athletes who are not using hormonal contraceptives and evaluate the assessment methods used to
34 identify MC disorders and MC-related symptoms.

35
36 **Methods:** This systematic review was performed in accordance with the preferred reporting items for
37 systematic reviews and meta-analyses (PRISMA). Six databases were searched until September 2022
38 for all original research that reported the prevalence of MC disorders and/or MC-related symptoms in
39 athletes not using hormonal contraceptives, which included the definitions of the MC disorders
40 examined, and the assessment methods used. MC disorders included amenorrhea, anovulation,
41 dysmenorrhea, heavy menstrual bleeding (HMB), luteal phase deficiency (LPD), oligomenorrhea,
42 premenstrual syndrome (PMS) and premenstrual dysphoric disorder (PMDD). MC-related symptoms
43 included any affective and physical symptoms related to the MC that do not cause significant personal,
44 interpersonal, or functional impairment. The prevalence data across eligible studies were combined and
45 all studies were qualitatively synthesised to evaluate the assessment methods and tools used to identify
46 MC disorders and MC-related symptoms. The methodological quality of studies was assessed using a
47 modified Downs and Black checklist.

48
49 **Results:** Sixty studies involving 6,380 athletes were included. A wide range of prevalence was
50 observed for all types of MC disorders, with a dearth of data on anovulation and LPD. Based on pooled
51 data, dysmenorrhea (32.3%; range 7.8% – 85.6%) was the most prevalent MC disorder. Studies
52 reporting MC-related symptoms mostly examined the premenstrual and menstruation phases, where
53 affective symptoms appeared more prevalent than physical symptoms. A larger proportion of athletes
54 reported symptoms during the initial days of menstruation compared to the premenstrual phase. MC

55 disorders and MC-related symptoms were retrospectively assessed using self-report methods in 90.0%
56 of studies. Most studies (76.7%) in this review were graded as moderate quality.

57

58 **Discussion:** MC disorders and MC-related symptoms are commonplace among female athletes,
59 warranting further research examining their impact on performance and preventive/management
60 strategies to optimise athlete health. To increase the quality of future studies, researchers should adopt
61 standardised definitions of MC disorders and assessment methods such as a combination of calendar
62 counting, urinary ovulation tests and a mid-luteal phase serum progesterone measurement when
63 assessing menstrual function. Similarly, standardised diagnostic criteria should be used when
64 examining MC disorders such as HMB, PMS and PMDD. Practically, implementing prospective cycle
65 monitoring that includes ovulation testing, mid-luteal blood sampling (where feasible) and symptom
66 logging throughout the MC could support athletes and practitioners to promptly identify and manage
67 MC disorders and/or MC-related symptoms.

68

69 This review has been registered in the PROSPERO database (CRD42021268757).

70

71 **Key Points**

- 72 1. Dysmenorrhea was the most prevalent menstrual cycle disorder in female athletes, with more
73 than one in four athletes self-reporting painful menstruation. Female athletes commonly
74 experience negative menstrual cycle-related symptoms during the premenstrual and
75 menstruation phases, where affective symptoms were more prevalent than physical symptoms.
- 76 2. Definitions of menstrual cycle disorders and the assessment methods used varied widely
77 across studies. Future studies should adopt standardised definitions and best practice methods
78 to assess menstrual function within the athletic population.
- 79 3. Practically, screening for menstrual cycle-related discomforts and disorders should be
80 conducted periodically within the training environment, with access to appropriate treatment
81 and care. Athletes should monitor their menstrual cycles including ovulatory function and
82 symptomatology throughout the entirety of the cycle.

83

84

85 **Declarations**

86 **I. Funding**

87 This work was funded by the Technological University of the Shannon: Midlands Midwest and the Irish
88 Research Council under grant number GOIPG/2022/2230 awarded to Bernadette Cherianne Taim.

89

90 **II. Conflicts of Interest**

91 Bernadette Cherianne Taim, Ciarán Ó Catháin, Michèle Renard, Kirsty Jayne Elliot-Sale, Sharon
92 Madigan and Niamh Ní Chéilleachair declare that they have no conflict of interest.

93

94 **III. Availability of data and material**

95 The data analysed during the current study are available from the corresponding author upon
96 reasonable request.

97

98 **IV. Ethics approval**

99 Not applicable

100

101 **V. Consent to participate**

102 Not applicable

103

104 **VI. Consent for publication**

105 Not applicable

106

107 **VII. Code availability**

108 Not applicable

109

110 **VIII. Author contributions**

111 BCT, MR and COC conceptualized the design of this study. BCT and MR conducted the literature
112 search, screening and data extraction. BCT conducted the formal analysis. BCT, COC and NNC
113 interpreted the data analysis. BCT wrote the manuscript with critical input from COC, KES, SM, NNC
114 and MR. All authors read and approved the final manuscript.

115

116 **1. Introduction**

117 Menstrual cycle (MC) disorders are broadly defined as abnormal uterine bleeding, ovarian dysfunction,
118 pain and other symptomatology that characterise premenstrual syndrome (PMS) [1–3]. Abnormal
119 uterine bleeding includes disruptions to bleeding patterns such as cycle length disturbances (e.g.,
120 oligomenorrhea and amenorrhea) and bleeding duration and/or flow (e.g., heavy menstrual bleeding
121 (HMB)) [4]. Luteal phase deficiency (LPD) and anovulation are disorders that represent subtle ovarian
122 dysfunction and while these conditions may occur without noticeable changes in bleeding patterns,
123 continued ovarian suppression can lead to oligomenorrhea and amenorrhea [5]. Dysmenorrhea (i.e.,
124 painful menstruation), PMS and premenstrual dysphoric disorder (PMDD) are MC disorders
125 characterised by negative MC-related symptoms that cause significant impairment during or before
126 menstruation. While dysmenorrhea has been classified as a MC disorder within this review, the
127 condition can also be recognised as a MC-related symptom and a key symptom secondary to
128 gynaecological disorders such as endometriosis [6]. The symptomatology that characterises PMS and
129 PMDD, however, differs from the mild and acute premenstrual symptoms experienced by many women
130 of reproductive age that are not necessarily indicative of a disorder [3]. While studies in the general
131 population have often focused on symptoms that occur before the onset of menstruation (i.e.,
132 premenstrual), MC-related symptoms can occur at any time across the entirety of the MC. Therefore,
133 MC-related symptoms are defined as any affective and physical symptoms related to the MC that do
134 not cause significant personal, interpersonal, or functional impairment [3,7].

135

136 Female athletes may be more susceptible than non-athletic populations to MC disorders mediated by
137 an alteration in the function of the hypothalamic-pituitary-ovarian (HPO) axis, including LPD, anovulation,
138 oligomenorrhea and amenorrhea [8]. In relation to the female athlete triad [9], these disorders lie on a
139 continuum representing varying degrees of ovarian suppression, with secondary amenorrhea being the
140 most severe form [10]. While the aetiologies of these MC disorders in athletes can vary considerably
141 beyond factors related to athletic training, it is acknowledged that they are predominantly associated
142 with low energy availability (i.e., inadequate energy intake relative to exercise energy expenditure),
143 resulting in disruptions to the HPO axis and hypoestrogenism which can have deleterious long-term
144 effects on an athlete's cardiovascular and musculoskeletal health [11–14]. Athletes may also

145 experience a range of other MC disorders including HMB, dysmenorrhea, PMS and PMDD, and the
146 burden of these disorders should not be overlooked. Unlike subtle ovarian dysfunction that may be
147 unnoticeable without appropriate MC monitoring (e.g., a combination of calendar counting, urinary
148 ovulation tests and mid-luteal phase serum progesterone measurement) [15], these MC disorders can
149 significantly impair daily functioning. In the general population, women with HMB have reported a
150 significantly worse quality of life than women with normal menstrual bleeding [16]. Further, MC disorders
151 such as dysmenorrhea and PMS have been shown to pose a considerable burden to work productivity
152 (e.g., work/school absenteeism) and healthcare utilisation [17,18]. While the prevalence and burden of
153 MC-related symptoms and such MC disorders have been less commonly discussed within sport and
154 exercise medicine-based research, emerging studies have highlighted that MC-related symptoms can
155 significantly impair an athlete's perceived ability to train and perform [19,20].

156

157 In athletes, the prevalence of amenorrhea and oligomenorrhea vary widely from 6-79%, and there is
158 seemingly less consideration for the other forms of MC disorders including anovulation and LPD [21,22].
159 The large variation in the prevalence of amenorrhea and oligomenorrhea observed in athletes could be
160 explained by methodological factors such as differences in athletic calibre, sports, case definitions and
161 screening tools or diagnostic criteria used in studies [23]. Indeed, the need for higher standards of
162 methodological practices to assess menstrual function has been well acknowledged in recent times
163 [15,24,25]. In a climate where the MC is increasingly recognised as a key health and performance
164 consideration for training female athletes [26,27], an evaluation of existing identification/screening tools
165 could inform and enhance the quality of female athlete monitoring practices.

166

167 At present, the prevalence of MC disorders and MC-related symptoms in female athletes is not well
168 documented. However, as in the general population, MC disorders and MC-related symptoms can pose
169 significant burdens to athletes including a decreased quality of life, increased healthcare utilisation and
170 crucially, negative health and performance consequences that may result in athletes missing
171 training/competition or in extreme cases, bring their sporting career to a premature end [28,29]. To that
172 end, knowing the prevalence of a wide range of MC disorders and the types of MC-related symptoms
173 in athletes would contribute to a greater understanding of their burden and guide research priorities in
174 female athlete health. Furthermore, an evaluation of existing identification/screening tools used to

175 assess menstrual function and symptomatology could facilitate the development of preventive
176 strategies to optimise female athlete health and performance. Therefore, the objectives of this study
177 were to 1) identify the prevalence of MC disorders among athletes, 2) identify the prevalence of MC-
178 related symptoms reported by athletes, and 3) evaluate the assessment methods or tools used to
179 identify MC disorders and MC-related symptoms in the sporting setting.

180

181 **2. Methods**

182 The reporting of this systematic review was in accordance with the preferred reporting items for
183 systematic reviews and meta-analyses (PRISMA) guidelines [30]. Prior to the commencement of the
184 study, the review protocol was registered in the PROSPERO database (CRD42021268757).

185

186 **2.1. Study Eligibility Criteria**

187 The Population, Intervention, Comparison, Outcomes and Study design (PICOS) framework was used
188 to formulate the eligibility criteria. No specific intervention was investigated.

189

190 **2.1.1. Population**

191 The study population included female athletes who were (a) of reproductive age (i.e., ages 13-39 years);
192 (b) competing at interscholastic, collegiate, national, international, amateur and/or professional levels;
193 and (c) not using hormonal contraceptives (HCs) and/or medication that alters the endogenous
194 hormonal milieu. Pre-menarcheal, peri/post-menopausal, and pregnant athletes were excluded.

195

196 **2.1.2. Comparison**

197 Comparisons were made across age groups, sports and athletic calibre. The age groups consisted of
198 adolescents between the ages of 13 and 18, and adults between the ages of 19 and 39. This age
199 classification was based on the American College of Obstetricians and Gynaecologists' (ACOG) age-
200 based common differential diagnosis of abnormal uterine bleeding, which accounts for the immaturity
201 of the HPO axis in adolescents [1].

202

203 **2.1.3. Outcomes**

204 The primary outcome was the prevalence of MC disorders in athletes. For the scope of this review, MC
205 disorders included the following conditions: amenorrhea (primary and secondary), anovulation,
206 dysmenorrhea (primary and secondary), HMB, LPD, oligomenorrhea, PMS and PMDD. The secondary
207 outcomes were the prevalence and types of MC-related symptoms in athletes and the assessment
208 methods used to identify MC disorders or MC-related symptoms. MC-related symptoms included any
209 affective and physical symptoms related to the MC that do not cause significant personal, interpersonal,
210 or functional impairment (i.e., not classified as PMS or PMDD).

211

212 **2.1.4. Study Design**

213 All original research, including observational cohort, cross-sectional, case-control studies and
214 randomised control trials (RCT), were considered for analysis if they included the following: (a) type and
215 definition of MC disorders, (b) quantitative prevalence proportion of MC disorders, and (c) tool used to
216 identify MC disorders. Studies that only investigated MC-related symptoms were considered for analysis
217 if they included the quantitative prevalence proportion of MC-related symptoms and the tool used to
218 identify MC-related symptoms. For RCT and longitudinal studies, only baseline prevalence proportion
219 data were considered. The search was restricted to human research published in English. Unpublished
220 research and abstracts from conference proceedings were considered if the inclusion criteria were met
221 and sufficient data were provided. The search was limited to publications up to September 2022.

222

223 **2.2. Search Strategy**

224 To identify eligible studies, a comprehensive search strategy was applied to electronic databases
225 including PubMed, SPORTDiscus, CENTRAL, ProQuest, CINAHL, and OpenGrey (grey literature).
226 Databases were searched by one investigator (BCT) from inception until July 2021. The reference list
227 of included articles was also examined manually to identify eligible articles. An updated search was
228 subsequently conducted in September 2022 to identify additional articles published between July 2021
229 and September 2022. The full list of search terms used is detailed in Table 1.

230

231 ***Insert Table 1 about here***

232

233

234

235 **Table 1** Search Terms

Concept	Keywords
Menstrual Disorder	<p>“Menstrual Dysfunction” OR “Menstrual Disturbance*” OR “Menstrual Irregularity” OR “Menstrual Problem*” OR “Menstrual Cycle Pattern*” OR “Menstrual Cycle Disorder*” OR “Menstrual Disorder*” OR “Menstrual Function” OR “Menstrual Status” OR “Premenstrual Symptom*” OR “Pre-menstrual Symptom*” OR “Premenstrual Syndrome” OR “Pre-menstrual Syndrome” OR PMS OR “Premenstrual Dysphoric Disorder” OR “Menstrual Cycle Symptom*” OR “Menstrual Symptom*” OR “Heavy Menstrual Bleeding” OR Menorrhagia OR Dysmenorrhea OR Dysmenorrhoea OR “Period Pain” OR Amenorrhea OR Amenorrhoea OR “Delayed Menarche” OR Oligomenorrhea OR Oligomenorrhoea OR Anovulation OR “Anovulatory Cycle*” OR “Short Luteal Phase” OR “Luteal Phase Deficiency” OR “Luteal Phase Defect*” OR LPD</p>
Female Athlete	<p>“Athlete*” OR “Player*” OR “Female Athlete*” OR “Sport*” OR Sportswoman OR “Athletic Women”</p>

236

237 **2.3. Study Selection**

238 The study selection process was conducted in two phases by two investigators (BCT and MR). In the
 239 first phase, the eligibility criteria were applied to the titles and abstracts of studies generated from the
 240 electronic searches. Duplicates and studies that were unrelated or did not meet the eligibility criteria
 241 were removed at this phase. Abstracts with insufficient information to determine their eligibility
 242 progressed to the next phase. In the second phase, the full texts of articles that progressed from phase
 243 one were retrieved for review. Each full text was screened against the predetermined inclusion and
 244 exclusion criteria. A third reviewer (COC) was available for consensus decisions but was not required
 245 to be consulted. Figure 1 outlines the study selection process.

246

247 **2.4. Data Extraction**

248 Data were extracted from the included studies by two investigators (BCT and MR) using a standardised
 249 data extraction form developed a priori. The following data were extracted where available: study
 250 characteristics (i.e., main author, year of publication, title, country, study design, and sample size),
 251 population characteristics (i.e., sport, competitive level, age, height, mass, body mass index (BMI), lean
 252 mass, body fat percentage, and age of menarche), type(s) of MC disorders or MC-related symptoms,
 253 prevalence proportion of MC disorders or MC-related symptoms (expressed as the number of athletes
 254 reporting MC disorders or MC-related symptoms out of the total study sample), definition(s) of MC
 255 disorders, and tool used to assess menstrual function or symptoms. If the demographic data reported

256 in a study were incomplete or insufficient to determine the characteristics of the study sample, the study
257 was removed.

258

259 **2.5. Assessment of Risk of Bias**

260 The methodological quality of included studies was independently assessed by two investigators (BCT
261 and MR) using a 15-item modified Downs and Black checklist [31] which was adapted for the purpose
262 of this review (Online Resource 1). In response to each item on the checklist, investigators rated 'yes'
263 (1-point), 'no' (0-point), or 'unable to determine' (0-point). The maximum attainable score was 15 and
264 based on the points awarded across the range of items, each study was rated as high (13–15),
265 moderate (9–12), low (8–11) or very low (<8) in study quality. Discrepancies were resolved via
266 consensus discussion between the two investigators (BCT and MR). A third reviewer (NNC) was
267 available for consensus decisions but was not required to be consulted.

268

269 **2.6. Data Synthesis**

270 **2.6.1. Quantitative Synthesis**

271 As preliminary searches revealed substantial methodological heterogeneity between studies, a meta-
272 analysis was not performed. Studies that used definitions of MC disorders that were consistent with
273 standards of practice (SOP) or acceptable alternatives (AA) (Table 2) were included in the quantitative
274 synthesis, where prevalence data across studies were combined (i.e., mean prevalence and range for
275 each disorder). Prevalence data for each MC disorder were analysed separately. SOP definitions are
276 defined by experts and/or relevant organisations specialising in female health (e.g., ACOG, International
277 Federation of Gynaecology and Obstetrics, International Society for Premenstrual Disorders). AA are
278 definitions that do not deviate substantially from the SOP (i.e., similar meaning) and are defined by
279 medical organisations such as the American Academy of Family Physicians. Studies that used AA
280 definitions were deemed eligible as the exclusion of these studies could lead to an underrepresentation
281 of the true prevalence. The prevalence data of studies that examined MC-related symptoms were
282 combined across studies and further reported by individual symptoms.

283

284 Diagnostic criteria were used to identify the following conditions, HMB, PMS and PMDD. Considering
285 varying diagnostic criteria across studies, non-pooled prevalence data were narratively reported for

286 these conditions, where the range for each MC disorder was reported. The prevalence data were
287 excluded from the quantitative analysis if the definition(s) of MC disorder(s) used significantly deviated
288 from the SOP/AA and/or diagnostic criteria.

289
290 Prevalence data were further reported by age group (adult and adolescent), sport and athletic calibre.
291 When a study involved both adult and adolescent athletes, it was classified based on the mean age.
292 Sports were clustered into seven broad categories: team, endurance/long distance, middle distance,
293 speed/strength, precision/skill-dependent, racquet, and combat/weight-making sports [32]. If a study
294 failed to provide sufficient information for categorisation into one single sport classification, the study
295 was excluded from the subgroup analysis by sport. To further characterise athletic calibre and training
296 status, a 6-tiered Participant Classification Framework was used to classify study participants into
297 different tiers based on training volume and performance metrics [32].

298

299 **2.6.2. Qualitative Synthesis**

300 All studies were qualitatively synthesised to evaluate the assessment methods and tools used to identify
301 MC disorders and MC-related symptoms in the sporting setting. Studies examining MC disorders were
302 evaluated based on a tiered ranking system that grades the standard of methodological control in
303 assessing menstrual status and/or menstrual irregularities, and each study was placed into one of four
304 tiers (i.e., Gold, Silver, Bronze or Ungraded) [33]. The study design (i.e., retrospective or prospective)
305 and specific tool used (e.g., questionnaire, interview, medical/gynaecological screening, symptom log)
306 in each study were also considered.

307

308 ***Insert Table 2 about here***

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

309 **Table 2** Definitions of menstrual cycle disorders which constituted the Standards of Practice (SOP) definitions

Menstrual cycle disorder	Definition	Reference ^a
Primary Amenorrhea	The failure to reach menarche by age 15 years* when development of secondary sexual characteristics is evident, or by age 14 years when no secondary sexual characteristics are present <i>*Acceptable alternative: age 16 years</i>	[15,34,35]
Secondary Amenorrhea	The absence of at least 3 consecutive periods* in non-pregnant females with past menstruation, which can be caused by other underlying endocrinopathies <i>*Acceptable alternative: three months or 90 days</i>	[15,34]
Oligomenorrhea	Menstrual cycle length greater than 35 days	[15]
Anovulation	Menstruate but do not ovulate (unable to detect ovulation by urinary luteinising hormone surge testing or confirmed by serum hormone analysis)	[15,36,37]
Luteal Phase Deficiency	Menstrual cycles with less than 16 nmol/L of progesterone (determined by a single luteal phase serum progesterone measurement)	[15,38]
Heavy Menstrual Bleeding	A type of abnormal uterine bleeding characterised by excessive cyclic blood loss which differs from normal uterine bleeding and interferes with a woman's physical, social, emotional, or material quality of life	[39]
Primary Dysmenorrhea	Painful menstruation (in the absence of pelvic pathology; the largest contributing factor is the increased levels of prostaglandins which stimulate myometrial contractions and reduce uterine blood flow, resulting in uterine hypoxia and painful abdominal cramps)	[40]
Secondary Dysmenorrhea	Painful menstruation (due to pelvic pathology or a recognised medical condition such as endometriosis)	[40,41]
Premenstrual Syndrome	Characterised by repetitive, cyclical, psychological, physical, and behavioural symptoms occurring in the luteal phase of the normal menstrual cycle that cause significant impairment, and resolve during or shortly after the onset of menstruation; Diagnostic criteria based on International Society for Premenstrual Disorders and American Academy of Family Physicians (Prospective charting)	[3,42,43]

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

Premenstrual Dysphoric Disorder The severe form of premenstrual syndrome; A depressive disorder not otherwise specified and emphasises emotional and cognitive-behavioural symptoms; Diagnostic criteria based on DSM-5 [44]
(Prospective charting)

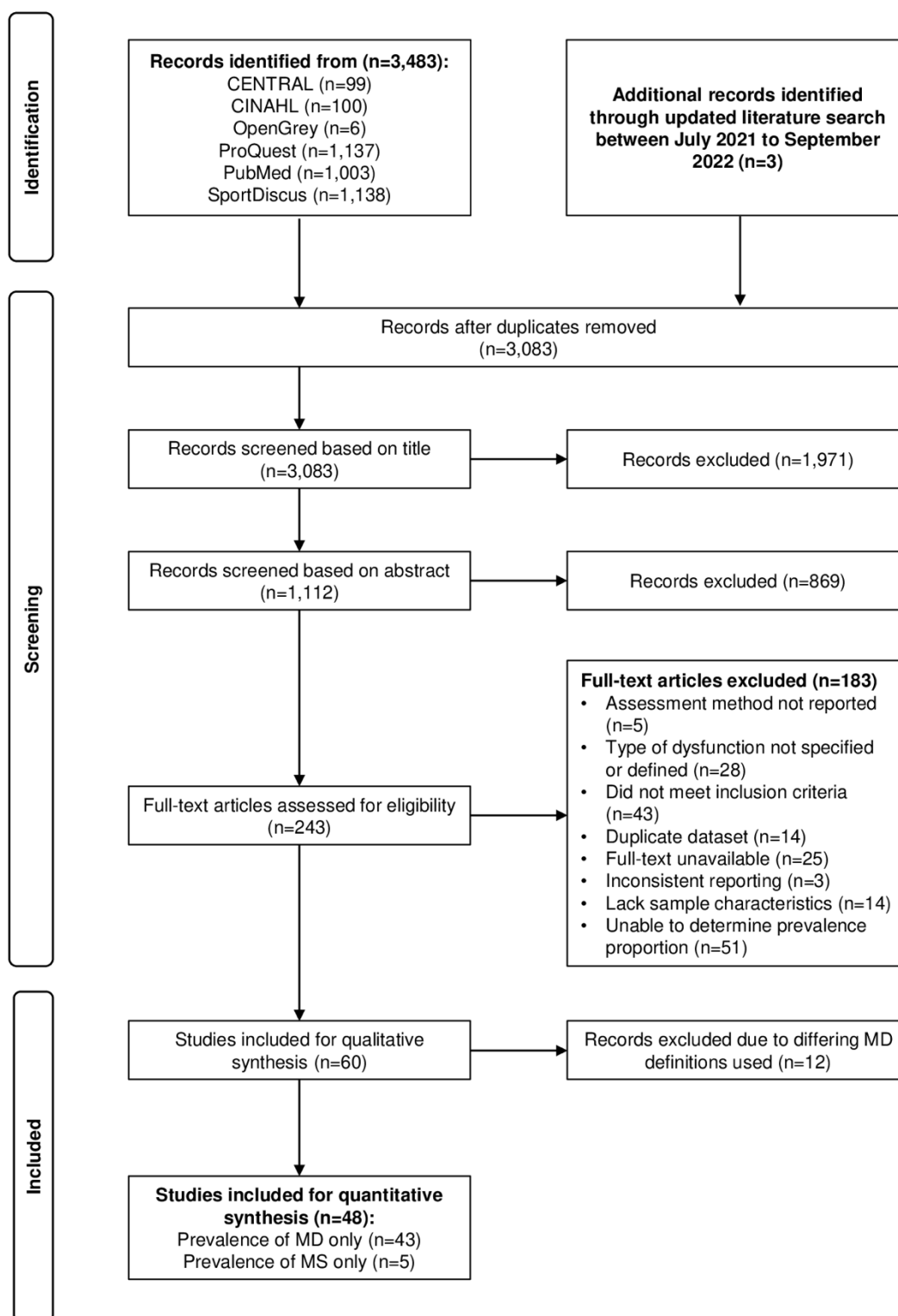
310 Abbreviations: DSM-5 = Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition. *Note:* ^aThe SOP definitions are defined by experts and/or relevant
311 organisations
312

313 **3. Results**

314 **3.1. Literature Search Results**

315 The literature search and study selection process is displayed in Figure 1. A total of 60 studies [45–104]
316 met the eligibility criteria and were included in this review with 48 included in the quantitative synthesis
317 of prevalence data. Among the 48 included studies, 43 studies reported the prevalence of MC disorders
318 only [45–48,50,52–54,56,57,59–62,64–68,70,72–78,81,82,85–90,93–95,97,98,100,102,104] and five
319 studies reported the prevalence of MC-related symptoms [69,71,79,91,103]. The remaining 12 studies
320 [49,51,55,58,63,80,83,84,92,94,96,101] were excluded due to differing definitions of MC disorders used
321 and they were only included in the qualitative synthesis.

322



323

324 **Fig. 1** Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow diagram

325 of the literature search and study selection process

326

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3.2. Study Characteristics

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3.3. Risk of Bias Assessment

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A total of 60 studies [45–104] involving 6,380 female athletes were included in the review. The included studies were published between 1981 and 2022, were all observational studies (cross-sectional, cohort, case-control) and included two unpublished masters and doctoral theses [83,100]. Based on the 6-tiered Participant Classification Framework, which classifies participants using training volume and performance metrics [32], included studies in this review consisted of the following athletic calibres; Tier 2: Trained/Developmental (n=18, 30.0%) [47,52–54,63,64,73,74,76,77,86,89,91,92,94,96,98,100], Tier 3: Highly Trained/National Level (n=20, 33.3%) [45,48,49,57,62,65,67,70–72,81,83–85,88,95,99,101,102,104], Tier 4: Elite/International Level (n=9, 15.0%) [55,59–61,66,68,69,78,80], and Tier 5: World Class (n=1, 1.7%) [79]. Twelve (20.0%) studies [46,50,51,56,58,75,82,87,90,93,97,103] did not provide sufficient information for a single classification of athletic calibre. The included studies involved various classifications of sports; Team sports (n=7) [45,57,60,71,72,79,100], endurance/long distance (n=10) [47,51,59,62,63,70,84,92,96,98], middle distance (n=5) [65,80,85,99,101], speed/strength (n=1) [88], precision/skill dependent (n=8) [46,49,50,53,56,68,75,87], and a combination thereof (n=29) [48,52,54,55,58,61,64,66,67,69,73,74,76–78,81–83,86,89–91,93–95,97,102–104]. Studies included both adults (n=43, age = 21.9 ± 3.2 years) and adolescents (n=17, age = 16.4 ± 0.9 years). A detailed description of the study and sample characteristics is presented in Online Resource 2.

The methodological quality of individual studies, as assessed by a modified Downs and Black checklist, is presented in Online Resource 3. Overall, none of the included studies was rated as 'high' quality with the majority of studies (n=46, 76.7%) rated as 'moderate' [45–48,50–54,56,57,59,61,64,65,68–70,73–85,88–91,93–95,97–100,102–104], 11 studies (18.3%) rated as 'low' [49,55,60,62,66,71,72,87,92,96,101], and three studies (5.0%) rated as 'very low' quality [58,67,86].

MC disorders were not defined in accordance with the SOP/AA definitions in twelve studies (20.0%) [49,51,55,58,63,80,83,84,92,94,96,101]. Only seven studies (11.7%) [49,51,63,82,93,94,96] confirmed that their participants were not using HCs for at least three months prior to participation.

356

357 3.4. Prevalence of Menstrual Cycle Disorders in Athletes

358 A total of 43 out of 60 studies [45–48,50,52–54,56,57,59–62,64–68,70,72–78,81,82,85–90,93–
359 95,97,98,100,102,104] were included in the quantitative synthesis of MC disorders prevalence data.
360 Studies that examined primary amenorrhea, secondary amenorrhea, oligomenorrhea and
361 dysmenorrhea used similar definitions according to the SOP/AA definitions and therefore the
362 prevalence data across studies were pooled (Table 3). In studies that examined a combination of age
363 groups (i.e., adults and adolescents), the prevalence for the respective age groups within the study was
364 extracted (where available) and pooled across studies (Table 3).

365

366 While studies that examined HMB, PMS and PMDD used definitions that were consistent with the SOP
367 definitions, the prevalence data across studies were not pooled due to variations in diagnostic criteria
368 and are instead described and reported in Table 4. There were no eligible studies that investigated the
369 prevalence of anovulation and/or LPD in athletes.

370

**371 3.4.1. Mean Prevalence of Primary Amenorrhea, Secondary Amenorrhea,
372 Oligomenorrhea and Dysmenorrhea**

373

374 ***Insert Table 3 about here***

375

376 Fourteen studies [46–48,54,56,64,68,70,76,78,81,85,86,95] were included in the combined prevalence
377 of primary amenorrhea. Eight studies [46,47,64,68,70,78,86,95] used the SOP definition (i.e., the failure
378 to reach menarche by age 15 years), while six studies [48,54,56,76,81,85] adopted the older age
379 threshold of 16 years old. For the combined prevalence of secondary amenorrhea, twenty-four studies
380 [47,50,54,56,59,61,62,64–66,70,72,75–78,81,82,85–88,98,100] were included. Nine studies
381 [47,64,66,70,72,81,85,86,98] used the SOP definition (i.e., the absence of at least 3 consecutive
382 periods in non-pregnant females with past menstruation) and 15 studies [50,54,56,59,61,62,65,75–
383 78,82,87,88,100] used the AA definition (i.e., the absence of menstruation in the previous three months
384 or 90 days).

385

386 Based on the SOP definition, 13 studies [45,47,59,60,64,70,72,75,76,85,86,95,100] were included in
387 the combined prevalence of oligomenorrhea. Seven studies [54,67,73,74,93,94,104] examined the
388 prevalence of dysmenorrhea in athletes. Only one study [67] specified the form of dysmenorrhea as
389 primary. As the remaining six studies [54,73,74,93,94,104] did not specify the type of dysmenorrhea
390 (i.e., primary or secondary), the mean overall prevalence of dysmenorrhea (i.e., painful menstruation)
391 was pooled across all seven studies.

392

393 The mean prevalence was also reported by age group (Table 3). The mean prevalence of primary
394 amenorrhea is 4.3% higher in adult than adolescent athletes, while the mean prevalence of
395 dysmenorrhea is 5.5% higher in adolescent than adult athletes (Table 3).

396

397 **3.4.2. Non-pooled Prevalence of Heavy Menstrual Bleeding, Premenstrual Syndrome and**
398 **Premenstrual Dysphoric Disorder**

399

400 The prevalence of individual PMS and PMDD symptoms is presented in Table 5.

401

402 ***Insert Table 4 about here***

403

404 ***Insert Table 5 about here***

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

405 **Table 3** Mean prevalence of Primary Amenorrhea, Secondary Amenorrhea, Oligomenorrhea and Dysmenorrhea

Menstrual cycle disorder	All athletes				Adult athletes only				Adolescent athletes only			
	n	Sample size	Mean prevalence (%)	Range (%)	n	Sample size	Mean prevalence (%)	Range (%)	n	Sample size	Mean prevalence (%)	Range (%)
Primary Amenorrhea	14	1260	7.1	0 – 32.0	8	691	9.2	0 – 32.0	7	569	4.9	0 – 11.8
Secondary Amenorrhea	24	1705	16.0	0 – 61.5	15	853	16.7	0 – 61.5	10	852	14.9	0 – 41.2
Oligomenorrhea	13	894	23.5	8.3 – 40.0	7	352	22.3	8.3 – 36.4	7	542	25.2	12.7 – 40.0
Dysmenorrhea	7	1734	32.3	7.8 – 85.6	3	1245	30.0	7.8 – 85.6	5	489	35.5	13.8 – 67.9

406 *Note:* n = number of studies

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

407 **Table 4** Prevalence of Heavy Menstrual Bleeding, Premenstrual Syndrome and Premenstrual Dysphoric Disorder (Non-pooled data)

Menstrual cycle disorder	Definition used	n	Study	Population	Sample size	Mean prevalence (%)
HMB (n=4)	Range of HMB prevalence across studies: 3.4% – 42.1%	4			821	
	A type of abnormal uterine bleeding characterised by excessive cyclic blood loss which differs from normal uterine bleeding and interferes with a woman’s physical, social, emotional, or material quality of life ^a	1	[104]	Adult	112	12.5
	Heavy amount of flow of menstrual blood	1	[74]	Adolescent	79	21.5
	Profuse menstrual flow due to no demonstrable anatomical or pathological lesion at least 5 times in the previous year	2	[94] [93]	Adult Adolescent	475 155	14.2 9.3
PMS (n=7)	Range of PMS prevalence across studies: 8.6% – 59.6%	7			1075	
	International Society for Premenstrual Disorders and American Academy of Family Physicians diagnostic criteria: Prospective daily monitoring of symptoms for at least two consecutive menstrual cycles ^a	3	[52] [57] [53]	Adult Adult Adolescent	75 52 45	49.3 59.6 48.9
	Adapted DSM-4 diagnostic criteria in a retrospective questionnaire that assessed premenstrual symptoms within the last three months	4	[89] [102] [97] [90]	Adult Adult Adult Adolescent	174 200 135 394	8.6 14.5 13.3 8.9
PMDD (n=5)	Range of PMDD prevalence across studies: 1.3% - 13.3%	5			888	
	DSM-5 diagnostic criteria: Prospective daily monitoring of symptoms for at least two consecutive menstrual cycles ^a	2	[52] [53]	Adult Adolescent	75 45	9.3 13.3

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Adapted DSM-4 diagnostic criteria in a retrospective questionnaire	3	[89]	Adult	174	2.9
that assessed premenstrual symptoms within the last three		[102]	Adult	200	2.5
months		[90]	Adolescent	394	1.3

408 Abbreviations: DSM-4/5 = Diagnostic and Statistical Manual of Mental Disorders, Fourth/Fifth Edition; HMB = Heavy Menstrual Bleeding; PMDD =

409 Premenstrual Dysphoric Disorder; PMS = Premenstrual Syndrome. *Note:* n = number of studies, ^aStandard of Practice definition

410 **Table 5** Prevalence of Premenstrual Syndrome and Premenstrual Dysphoric Disorder symptoms in
 411 athletes

PMS/PMDD symptom	n	Mean prevalence (range) (%)
Affective		
Overeating or food cravings	3	69.3 (63.9 - 72.4)
Anger or irritability	5	68.1 (44.65 - 80.6)
Anxiety or tension	5	60.7 (18.7 - 79.9)
Difficulty concentrating	3	58.6 (56.0 - 60.3)
Depressed mood or mood swings	5	54.9 (40.0 - 77.4)
Fatigue or lack of energy	4	54.7 (7.6 - 72)
Insomnia or hypersomnia	3	51.7 (43.9 - 56.5)
Tearful	3	47.2 (42.2 - 51)
Decreased interest in work, home, or social	4	33.6 (5.3 - 48)
Feeling overwhelmed	3	31.8 (29.7 - 34)
Embarrassment or confusion	1	25.3
Physical		
Abdominal congestion/swelling/bloating	1	72.0
Breast swelling/tenderness	2	65.3 (61.3 - 69.3)
Cramps or back pain	2	61.4 (55.1 - 67.7)
Physical symptoms/limb oedema	4	51.6 (6.7 - 75.5)
Headache	2	17.4 (0.0 - 34.7)
Acne	1	12.7
Diarrhoea	1	2.5

412 Abbreviations: PMDD = premenstrual dysphoric disorder; PMS = premenstrual syndrome. *Note:* n =
 413 number of studies

414

415 3.5. Prevalence of Menstrual Cycle-Related Symptoms in Athletes

416 Five studies [69,71,79,91,103] reported the prevalence and types of MC-related symptoms in athletes.

417 Among these, two studies [91,103] reported premenstrual symptoms only (i.e., symptoms occurring in

418 the luteal phase) and only provided the prevalence of individual symptoms. The remaining three studies

419 [69,71,79] reported the overall prevalence of MC-related symptoms and the prevalence of individual

420 symptoms across both the premenstrual and menstruation phases, where symptoms were reported by

421 74.0 to 82.4% of athletes. Two studies [69,79] compared the prevalence of symptoms between both

422 phases and found that a larger proportion of athletes (59.0 - 81.6%) reported symptoms on the initial

423 days of menstruation (day 1-3) compared to the premenstrual phase (i.e., lead up to menstruation)

424 (17.0 – 25%).

425

426 Among all the symptoms reported across the premenstrual and menstrual phases, affective symptoms
 427 including anger/irritability (49.2%) and anxiety/tension (45.9%), as well as physical symptoms such as
 428 abdominal cramps (58.8%) were the most prevalent. The prevalence of individual premenstrual
 429 symptoms is presented in Table 6.

430

431 ***Insert Table 6 about here***

432

433 **Table 6** Prevalence of menstrual cycle-related symptoms occurring during the premenstrual and
 434 menstruation phases in athletes

Menstrual Cycle-Related Symptom	n	Mean prevalence (range) (%)
Affective		
Insomnia or hypersomnia	2	53.3 (46.3 - 60.3)
Anger or irritability	4	49.2 (0.5 - 71.6)
Anxiety or tension	3	45.9 (0.5 - 75.0)
Overeating or food cravings	3	40.3 (0.9 - 70.4)
Decreased interest in work, home, or social	2	40.0 (29.9 - 50.0)
Fatigue or lack of energy	4	39.5 (4.1 - 72.7)
Difficulties in concentrating or coordination	3	39.4 (2.3 - 60.1)
Feeling overwhelmed	2	34.0 (25.9 - 42.0)
Depressed mood or mood swings	4	25.3 (4.0 - 61.4)
Physical		
Abdominal cramps	2	58.8 (47.5 - 70.0)
Physical symptoms/limb oedema	4	32.9 (1.8 - 78.4)
Abdominal congestion/swelling/bloating	3	17.5 (2.0 - 45.1)
Back pain	2	16.1 (15.0 - 17.1)
Breast swelling/tenderness	3	11.4 (0.9 - 31.4)
Headache	1	9.7
Vomiting/Feeling sick	2	4.3 (4.0 - 4.6)
Acne	1	0.5

435 *Note:* n = number of studies

436

437 3.6. Prevalence of Menstrual Cycle Disorders Across Different Sports

438 Twenty-five out of the 43 studies [45–47,50,53,54,56,57,59,60,62,65,67,68,70,72,75,85–
 439 88,93,94,98,100] that were included in the quantitative analysis of MC disorders were further analysed

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440 by sports classification (e.g., team sports) and respective sports (e.g., basketball) under the
441 classification (Table 6). For studies that examined a combination of sports classifications, the
442 prevalence data for each sport within the study was extracted. Therefore, the studies included in this
443 subgroup analysis by sport were conducted across various classifications of sport; team sports (n=9)
444 [45,54,57,60,67,72,93,94,100], precision/skill dependent (n=9) [46,50,53,54,56,68,75,86,87],
445 endurance/long distance (n=5) [47,59,62,70,98], speed/strength (n=3) [88,93,94], middle distance (n=2)
446 [65,85], racquet sports (n=2) [93,94], and combat/weight-making (n=1) [67].

447

448 ***Insert Table 7 about here***

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

Table 7 Prevalence of menstrual cycle disorders by sports classification and respective sports

Sport	n	Primary Amenorrhea	Secondary Amenorrhea	Oligomenorrhea	Dysmenorrhea	Heavy Menstrual Bleeding	Premenstrual Syndrome	Premenstrual Dysphoric Disorder
Team Sports	9	0	2.4 – 30.0	11.1 – 29.4	8.2 – 34.6	7.8 – 11.9	59.6	NA
Basketball [54,60,67,100]	4	0	2.4 – 5.6	12.7 – 26.1	30.6	–	–	–
Handball [60]	1	–	–	16.0	–	–	–	–
Volleyball [54,86]	2	0	0 - 30.0	11.1 – 40.0	34.6	–	–	–
Soccer [45,57,72]	3	–	3.1	13.8 – 29.4	–	–	59.6	–
Ball sports [93,94]	2	–	–	–	8.2 – 14.1	7.8 – 11.9	–	–
Endurance/long distance	5	0 – 10.0	0 – 35.0	15.0 – 34.6	NA	NA	NA	NA
Cross-country running [47]	1	0	3.8	34.6	–	–	–	–
Distance/track running [59,70,98]	3	10	0 – 35.0	15.0 – 31.8	–	–	–	–
Distance running/triathlon [62]	1	–	15.5	–	–	–	–	–
Middle distance	2	0	0 – 61.5	24.6	NA	NA	NA	NA
Distance running [65]	1	–	61.5	–	–	–	–	–
Swimming [85]	1	0	0	24.6	–	–	–	–
Speed/strength	3	NA	0	NA	15.8 – 30.0	4.2 – 42.1	NA	NA
Sprinting [88,93,94]	3	–	0	–	15.8 – 30.0	13.7 – 42.1	–	–
Jumping [94]	1	–	–	–	25.0	4.2	–	–

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Throwing [94]	1	–	–	–	18.9	17.0	–	–
Precision/skill-dependent	9	3.3 – 22.1	9.5 – 44.1	8.3 – 28.6	NA	NA	48.9	13.3
Ballet [46,50,54,56,75,87]	6	3.3 – 20.0	9.5 – 44.1	28.6	–	–	–	–
Gymnastics [53,86]	2	2.8	–	8.3	–	–	48.9	13.3
Rhythmic Gymnastics [68]	1	22.1	–	–	–	–	–	–
Racquet sports [93,94]	2	NA	NA	NA	21.3 – 31.3	9.4 – 10.6	NA	NA
Combat/weight-making	1	NA	NA	NA	36.8 – 45.8	NA	NA	NA
Taekwondo [67]	1	–	–	–	45.8	–	–	–
Judo [67]	1	–	–	–	36.8	–	–	–

450 *Note:* Data presented as range (%), n = number of studies, NA = Data not available. The rows in bold represent the total number of studies (n) examined in
 451 each sport classification, and the overall range of prevalence for each menstrual cycle disorder in each sport classification
 452

3.7. Prevalence of Amenorrhea and Oligomenorrhea Across Different Athletic Calibres

Thirty-four out of 43 studies that were included in the MC disorders quantitative analysis were further analysed by athletic calibre (Table 8) [45,47,48,52–54,57,59–62,64–68,70,72–74,76–78,81,85,86,88,89,94,95,98,100,102,104]. These studies provided sufficient information for a single classification of athletic calibre including Tier 2: Trained/Developmental (n=14) [47,52–54,64,73,74,76,77,86,89,94,98,100], Tier 3: Highly Trained/National Level (n=14) [45,48,57,62,65,67,70,72,81,85,88,95,102,104], and Tier 4: Elite/International Level (n=6) [59–61,66,68,78]. This subgroup analysis was limited to only the prevalence of primary amenorrhea, secondary amenorrhea and oligomenorrhea considering the small number of studies, or lack thereof, examining the other MC disorders.

463

464

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

465 **Table 8** Prevalence of menstrual cycle disorders by athletic calibres

Menstrual cycle disorder	Tier 2: Trained/Developmental				Tier 3: Highly Trained/National				Tier 4: Elite/International			
	n	Sample size	Mean prevalence (%)	Range (%)	n	Sample size	Mean prevalence (%)	Range (%)	n	Sample size	Mean prevalence (%)	Range (%)
Primary Amenorrhoea	4	337	2.6	0 – 6.4	3	157	4.5	0 – 10.0	2	152	11.7	1.3 – 22.1
Secondary Amenorrhoea	7	768	12.8	2.4 – 30.8	7	468	15.9	0 – 61.5	4	186	17.2	0 – 43.9
Oligomenorrhoea	5	463	18.1	8.3 – 34.6	4	308	23.8	13.8 – 36.3	1	22	31.8	-

466 *Note:* n = number of studies

467

468 **3.8. Assessment Methods Used to Identify Menstrual Cycle Disorders and Menstrual**
469 **Cycle-Related Symptoms**

470 All 60 studies included in this review were qualitatively synthesized to evaluate the tools used to
471 identify/diagnose MC disorders in athletes (Online Resource 3). Studies examining MC disorders (n=56)
472 were assessed according to the tiered ranking system that grades the standard of methodological
473 control considering menstrual status and/or menstrual irregularities [33]. Only two studies (3.6%) [70,77]
474 were placed into the 'Gold' tier (i.e., menstrual irregularities diagnosed by a medical professional as
475 part of the study). One study (1.8%) was placed into the 'Silver' tier (i.e., menstrual irregularities
476 diagnosed by a medical professional not as part of the study but self-reported or via medical records).
477 As the remaining 53 studies did not explicitly state that the MC disorders were diagnosed by a medical
478 professional, they were considered as 'Bronze' tier (i.e., self-reported condition without a medical
479 diagnosis). While seven studies [45,47,61,70,75,77,88] included serum hormone measurements, only
480 two studies [70,75] used hormone levels for the purpose of determining menstrual function.

481

482 Fifty-four out of sixty studies (90.0%) used retrospective self-report in the form of self-developed, non-
483 validated questionnaires (n=45) [46–49,51,54–56,58,60,62,64,66–69,72–76,78–84,86,87,89–91,91–
484 94,96–104], interviews (n=2) [50,63,95], Low Energy Availability in Females Questionnaire (LEAF-Q)
485 (n=2) [70,88], and medical/gynaecological screening (n=3) [45,61,77]. While one study [85] reported
486 the use of a validated questionnaire to assess the presence of MC disorders, its validation could not be
487 verified [105]. Six out of 60 studies (10.0%) used prospective self-report/charting methods through daily
488 training logs (n=2) [65,71], daily temperature logs (n=1) [59], and daily symptoms logs (n=3) [52,53,57].
489 Among the five studies which reported MC-related symptoms, four studies used retrospective self-
490 report questionnaires [69,79,91,103] while one study tracked symptoms prospectively using training
491 logs [71].

492

493 Overall, an evaluation based on a combination of the (1) tiered ranking system which is underpinned
494 by best practice to assess menstrual function, (2) study design (i.e., prospective or retrospective) and/or
495 (3) tool (i.e., validation) revealed that the assessment method used to assess MC disorders was
496 considered to be valid and reliable in only one study (5.0%) which used transvaginal ultrasound

497 examination to assess menstrual function [52,53,70]. While two studies used the standard diagnostic
498 criteria for PMS/PMDD including daily prospective assessment of PMS and PMDD symptoms over two
499 successive MCs, a clinical/psychiatric diagnosis to exclude other mental disorders which is crucial in
500 diagnosing PMS/PMDD, was not performed within the study.

501

502 **4. Discussion**

503 This review aimed to establish the prevalence of MC disorders and MC-related symptoms in female
504 athletes of reproductive age and included the following conditions: amenorrhea (primary and
505 secondary), anovulation, dysmenorrhea (primary and secondary), HMB, LPD, oligomenorrhea, PMS
506 and PMDD. A wide range of prevalence was reported across all MC disorders and the pooled
507 prevalence results indicated that dysmenorrhea was the most prevalent MC disorder. Studies that
508 reported MC-related symptoms mainly explored symptoms which occurred during the premenstrual and
509 menstruation phases, with affective symptoms appearing more prevalent than physical symptoms. Most
510 of the studies identified MC disorders using retrospective self-report in the form of non-validated
511 questionnaires, and in most studies, it was unclear whether a formal diagnosis was made by a medical
512 professional. Of the 60 studies identified, no studies achieved a high methodological quality rating, with
513 the majority presenting as moderate quality. The observed variation in assessment methods and
514 definitions introduces significant heterogeneity across studies. Therefore, findings should be interpreted
515 under the context of these limitations.

516

517 **4.1. Definitions of Menstrual Disorders**

518 Regardless of the type of MC disorder, a wide range of prevalence was reported within the literature.
519 This could be due to heterogeneity across studies considering methodology, assessment method, and
520 definition of MC disorders. Indeed, this review process highlighted a lack of uniformity in defining MC
521 disorders within the athletic population and several studies were excluded during the screening phase
522 as the type of MC disorders examined was not defined, or non-specific definitions were used. This is
523 particularly pertinent in studies that examined MC disorders associated with abnormalities in bleeding
524 frequency (e.g., amenorrhea, oligomenorrhea) where terms such as 'irregular cycle', 'menstrual
525 irregularities', and 'short cycle' were used without specifying a frequency and/or time frame.

526

527 While studies examining amenorrhea and/or oligomenorrhea were only included in the quantitative
528 analysis if accurate definitions were used, there were still slight variations in the units of time used. For
529 example, the units of time used to define secondary amenorrhea ranged from the absence of
530 menstruation for 'six or more *months* or the length of three *cycles*' in nine studies to 'previous three
531 *months* or 90 *days*' in 15 studies. Further, a threshold age of 15 years was used to define primary
532 amenorrhea in eight studies, and the other six studies utilised 16 years. Arguably, these differing
533 measurements may be somewhat equivalent and interchangeable. However, to reduce ambiguity within
534 the literature, it would be beneficial to standardise the definitions of MC disorders using a consistent
535 time frame and unit of time. This is especially relevant to MC disorders such as amenorrhea and
536 oligomenorrhea, as these conditions can be established by the clinical observation of cycle length
537 (oligomenorrhea) or the absence of menstruation over a prolonged period (amenorrhea) [22].

538

539 Varying prevalence were also observed in other MC disorders including dysmenorrhea, HMB, PMS and
540 PMDD. The broad definitions used across studies and the subjective nature of these conditions, in
541 which the occurrence and severity are largely determined by an individual's perception, could account
542 for the wide variation in prevalence reported. Given the pre-existing diagnostic criteria by the
543 International Society for Premenstrual Disorders and DSM-5 to identify PMS and PMDD [44,106], a
544 standardised definition of PMS and PMDD, while useful, may be less relevant. All things considered,
545 the lack of uniformity in defining MC disorders within the athletic population highlights the need to
546 identify standardised definitions, alongside valid assessment methods, that could be used consistently
547 by researchers and practitioners to reduce ambiguity and enhance the comparability of data within the
548 literature.

549

550 **4.2. Prevalence of Menstrual Cycle Disorders and Menstrual Cycle-Related Symptoms**

551 ***Dysmenorrhea, Premenstrual Syndrome and Premenstrual Dysphoric Disorder***

552 Dysmenorrhea was the most common MC disorder, with a mean pooled prevalence of 29.4%, indicating
553 that, from the competitive female athletes surveyed, more than one in four self-reported that they were
554 suffering from painful menstruation. A high prevalence of PMS was also observed, with prevalence
555 ranging from 48.9% to 59.6% in studies that adopted the gold standard method to assess PMS (i.e.,
556 prospective charting of symptoms).

557

558 The prevalence of dysmenorrhea reported in this review concurs with findings in general population-
559 based studies that reported dysmenorrhea as the most common gynaecological condition in women
560 [107]. In three studies [54,74,104], the prevalence of dysmenorrhea was lower in female athletes as
561 compared to non-athletes, with one study [74] also reporting a lower prevalence of PMS in athletes.
562 These findings suggest a possible beneficial effect of exercise training against primary dysmenorrhea
563 and PMS, supporting the results of a systematic review by Armour et al [108] which demonstrated that
564 regular exercise, regardless of intensity, may reduce the intensity of primary dysmenorrhea by lowering
565 the amount of prostaglandins synthesised or released [109,110]. However, more research is warranted
566 due to the current low-quality evidence [108]. Conversely, PMDD was reported to be the least common
567 MC disorder in athletes (range 1.3% - 13.3%). This finding is consistent with that of the general
568 population which typically reported a PMDD prevalence of 2% to 10%, likely due to the more rigorous
569 diagnostic criteria for PMDD in accordance with the DSM-5 as a mental disorder [111].

570

571 ***Menstrual Cycle-Related Symptoms***

572 While PMS and PMDD are characterised by negative premenstrual symptoms, it is important to
573 distinguish these MC disorders (which diagnoses require standard clinical diagnostic criteria) from the
574 physiological premenstrual symptoms experienced by many women that are often transient and do not
575 cause significant impairment [7,106]. Most studies in this review examined MC-related symptoms
576 occurring before and during the onset of menstruation and findings suggest that during these phases,
577 affective symptoms are more prevalent than physical symptoms. The most common affective symptoms
578 among athletes during these phases were insomnia/hypersomnia, anger/irritability and anxiety/tension,
579 while the most prevalent physical symptom was abdominal cramps. It should be noted that more than
580 200 premenstrual symptoms have been described in the literature [42] and therefore, the list of
581 symptoms reported in this review is by no means exhaustive. While MC-related symptoms are common
582 in the athletic population including elite athletes [112], this is of concern as perceived reductions in
583 training and/or competition performances have been widely reported even at the highest competition
584 levels [20,113]. Therefore, athletes and practitioners should recognise the impact that MC-related
585 symptoms can have on readiness to train/compete, and implement symptom tracking as part of MC
586 monitoring alongside periodic screening for menstrual pain [26,112]. However, the challenge lies in the

587 management approach and treatment strategies of these symptoms, which will likely be highly
588 individualised. Specifically, the efficacy of medical and non-pharmacological treatments in mitigating
589 the occurrence and severity of MC-related symptoms among athletes has yet to be elucidated [19]. To
590 that end, further research to evaluate the effectiveness of interventions to manage MC-related
591 symptoms in athletes should be undertaken.

592

593 Interestingly, despite using general search terms such as 'Menstrual Cycle Symptom' in the search
594 strategy, included studies in this review focused solely on the negative MC-related symptoms
595 experienced by female athletes during the premenstrual and menstruation phases. Indeed, research
596 on MC-related symptoms seems to reflect a negative bias that does not fully encompass the
597 complexities of the physical and emotional changes across the entirety of the MC [114]. For example,
598 previous research has demonstrated that some athletes, albeit a small minority, perceived that aspects
599 of their training were positively affected during menstruation [20]. Furthermore, a recent study by
600 McNamara et al. [112] indicated that while 41.5% of Australian Olympic/Paralympic athletes preferred
601 to compete 'just after (their) period', 4% desired to compete during the premenstrual phase and 1%
602 'during their period'. Therefore, future research and monitoring practices should consider examining the
603 occurrence and interference of both negative and positive effects and experiences across the entire
604 MC. This could not only change the narrative that all symptoms associated with the MC are barriers to
605 performance but from a practical standpoint, help athletes to develop and understand their individual
606 MC profiles and identify performance trends (if any). By managing any negative symptoms and
607 leveraging on positive symptoms throughout the MC, the MC could then be viewed as an informative
608 tool, rather than a limiting factor, to ideally help athletes optimise performance on any day of the MC.

609

610 ***Oligomenorrhea and Amenorrhea***

611 The mean pooled prevalence of primary amenorrhea, secondary amenorrhea and oligomenorrhea were
612 7.1%, 16.0% and 22.7%. The findings are proportionate to but higher than that of the general population,
613 where the prevalence of oligomenorrhea and amenorrhea have been reported to be 13.5% and 3% to
614 4% [35,115]. This supports the well-established viewpoint that athletes are more susceptible to
615 oligo/amenorrhea, especially that of hypothalamic origin, as compared to non-athletes [8,116].

616

617 Among these abnormalities in bleeding frequency, oligomenorrhea was most common among female
618 athletes. In reference to the Triad, MC disorders lie along a continuum with varying degrees of severity
619 which ranges from subclinical/subtle disturbances (i.e., LPD, anovulation) to clinical/severe
620 disturbances (i.e., oligomenorrhea and secondary amenorrhea) [117,118]. Therefore, the lower severity
621 of oligomenorrhea compared to secondary amenorrhea might help explain in part why the prevalence
622 of oligomenorrhea is relatively higher. This also demonstrates that athletes who experience
623 abnormalities in bleeding frequency would more commonly experience a reduced cycle frequency,
624 rather than the extreme outcome of the complete cessation of menses (i.e., secondary amenorrhea).
625 Nonetheless, as amenorrhea is associated with lifelong negative skeletal health consequences, any
626 disruption in bleeding frequency requires priority and consideration. Specifically, low energy availability
627 and amenorrhea suppress bone formation and upregulate bone resorption, resulting in low bone mass
628 and poor bone geometry [12,119]. This can predispose female athletes to bone stress injuries ranging
629 from stress fractures to osteoporosis at a younger age, which not only reduces sports performance but
630 importantly, compromises athlete health [29,120,121]. Therefore, as oligomenorrheic athletes can
631 progress along the continuum of menstrual disturbances to secondary amenorrhea, timely identification
632 and appropriate management of oligomenorrheic athletes is crucial.

633

634 ***Luteal Phase Deficiency and Anovulation***

635 There is a paucity of information on the prevalence of anovulatory and LPD cycles in competitive female
636 athletes, as highlighted by the absence of eligible studies in this review. This could be explained in part
637 by the methodological rigour of identifying subtle MC disorders [22,118]. As compared to MC disorders
638 which can be clearly recognised through clinical observation (e.g., oligomenorrhea, amenorrhea), the
639 gold standard method to identify LPD and anovulation requires daily measures of luteinising hormone
640 and ovarian hormones, which are relatively costly and time-consuming. Nevertheless, more studies
641 examining subtle MC disorders in competitive athletes are necessary and warranted, especially since
642 approximately half of recreationally physically active women were reported to experience LPD and
643 anovulation [22]. Given that LPD and anovulation exist on the continuum of menstrual disturbances
644 described in the female athlete triad as 'milder' forms of ovarian suppression that can progress to the
645 complete cessation of menses, it is crucial that these conditions are not overlooked and inaccurately
646 perceived as eumenorrhea. This lack of consideration for LPD and anovulatory cycles in research is

647 also reflected in applied practice, where anecdotally, MC length is commonly used as a sole marker to
648 establish menstrual function without confirmation of ovulation by urinary luteinising hormone surge or
649 serum hormone concentrations. The sole reliance on MC length fails to detect subtle menstrual
650 disturbances and overlooks apparently regular yet abnormal cycles [22]. Such inadequate MC
651 monitoring practices could be, in part, due to the lack of knowledge and awareness of subtle MC
652 disorders and how to detect them, especially in a context where much of the discussions have
653 surrounded the more severe presentations of menstrual disturbances in athletes such as
654 oligomenorrhea and amenorrhea. Given that LPD and anovulation are in fact early warning signs of
655 ovarian suppression, athletes and practitioners need to be aware of these subtle MC disorders and
656 where possible, implement appropriate detection methods such as urinary ovulation tests and a mid-
657 luteal blood sample for the determination of progesterone [15]. Indeed, recent methodological
658 recommendations suggesting the replacement of daily hormone assessments with urinary ovulation
659 detection kits, followed by a single luteal phase serum progesterone measurement may reduce previous
660 logistical and knowledge barriers and potentially increase the capacity for appropriate MC monitoring
661 and high-quality research investigating anovulation and LPD [15,38].

662

663 ***Heavy Menstrual Bleeding***

664 The prevalence of HMB in athletes in this review ranged from 3.4% to 42.1%. This review demonstrates
665 that athletes may also experience other MC disorders beyond those identified within the Triad and given
666 the potential for an increased risk of iron deficiency and anaemia in women with HMB, it is crucial that
667 the condition does not remain underdiagnosed and undertreated [122]. The impact of HMB would be
668 particularly concerning for endurance athletes, as low serum ferritin levels can reduce oxygen-carrying
669 capacity, which is crucial for performance [123]. It is significant to note that currently there is no
670 universally accepted definition of HMB [124]. Hence, inconsistent definitions and the subjective nature
671 of the condition hinder the comparability of prevalence across studies and populations.

672

673 ***Prevalence of Menstrual Cycle Disorders by Sports Classifications***

674 It has been well established that the prevalence of MC disorders varies widely with sport, and greater
675 susceptibility to MC disorders has been observed in sports that emphasise leanness [10]. Due to the
676 small number of studies eligible for inclusion in this subgroup analysis, comparisons of prevalence

677 across sports were avoided and instead descriptively presented in Table 6. Nonetheless, it is apparent
678 that most studies included in this review were conducted on endurance/long distance (e.g., cross-
679 country running), precision/skill-dependent (e.g., gymnastics) and team sport athletes. Of concern,
680 there is a lack of studies examining MC disorders in combat/weight-making sports, highlighting a crucial
681 gap in the literature. Aggressive weight reduction strategies are still highly prevalent in combat/weight-
682 making sports despite the well-documented negative implications of rapid weight loss on health [125].
683 For female combat sport athletes, severe weight reduction behaviour is a risk factor for the Triad [126].
684 However, research in this area remains scarce. Given that weight loss is one of the three main
685 underlying causes of functional hypothalamic amenorrhea [127], there is an urgent need for future
686 research to examine the effects of rapid weight loss and weight fluctuations on menstrual function in
687 female combat/weight-making athletes.

688

689 ***Prevalence of Oligomenorrhea and Amenorrhea by Athletic Calibres***

690 The subgroup analysis by athletic calibres highlighted an observable upward trend in the prevalence of
691 MC disorders from trained/developmental athletes to elite/international level athletes. Considering that
692 only a small number of studies were included within each athletic calibre and MC disorder in this
693 subgroup analysis, caution is necessary when interpreting these findings. Nonetheless, this upward
694 trend in prevalence for oligomenorrhea and primary and secondary amenorrhea seems to suggest that
695 MC disorders with a hypothalamic origin may be more prevalent in higher-calibre athletes than lower-
696 calibre athletes. Indeed, previous research has demonstrated that elite/international female athletes
697 may be at a greater risk of low energy availability and menstrual disturbances compared to
698 trained/developmental [128] and recreationally active athletes [129]. This is likely mediated by the
699 higher training frequencies and volumes that higher-calibre athletes are exposed to, which induce
700 elevated exercise energy expenditure that may contribute to low energy availability if unmatched with
701 sufficient energy intake [32,130]. However, evidence in this area is conflicting as previous research has
702 also demonstrated that higher-calibre athletes were not at increased risk of low energy availability and
703 associated health outcomes [131]. Further, it is also important to recognise that the aetiologies of MC
704 disorders vary widely beyond low energy availability. For example, a previous study that examined the
705 menstrual status of female Olympic athletes highlighted that the most common endocrinological
706 mechanism underlying oligomenorrhea and/or anovulation was polycystic ovary syndrome (PCOS)

707 rather than low energy availability [132]. As the differential clinical diagnosis of MC disorders is beyond
708 the scope of this review, further investigations on the prevalence of endocrinological mechanisms
709 underlying MC disorders (e.g., PCOS, low energy availability, hyperprolactinemia) across different
710 athletic calibres should be undertaken. Taken together, to better risk stratify athletes, future research is
711 needed to further examine the relationship between athletic calibre, associated risks and mechanisms
712 of MC disorders in athletes.

713

714 **4.3. Assessment Methods**

715 The assessment methods used are invariably dependent on the type of MC disorders examined as well
716 as their specific objectives (e.g., diagnostic or screening tool) and while an in-depth discussion on the
717 differential clinical diagnosis and diagnostic procedures of various MC disorders is beyond the scope
718 of this review, the methodological standards in current research will be discussed. Overall, menstrual
719 irregularities were assessed retrospectively using self-report methods in 90% of studies. A majority of
720 studies used self-developed menstrual history questionnaires that were not validated, of which only
721 three studies piloted their questionnaires [55,98,100]. Menstrual history questionnaires may be
722 sufficient, as a screening tool, to detect abnormalities in cycle length and/or frequency such as
723 secondary amenorrhea and oligomenorrhea prior to a clinical evaluation. However, the use of
724 retrospective self-report methods introduces potential variability in the findings due to limitations such
725 as recall bias [133]. Previous research has also shown considerable measurement error in MC
726 parameters reported retrospectively, and a further reduction in reporting accuracy with increasing cycle
727 variability [134,135]. Furthermore, the singular use of questionnaires would not detect the presence of
728 LPD and anovulatory cycles. This suggests that the current methods used in research to identify
729 abnormalities in bleeding frequency and ovarian function should be improved. Specifically, as recent
730 methodological guidelines recommended, instead of using a singular method (e.g., calendar-based
731 counting only), a combination of methods such as calendar tracking, urinary ovulation detection kits
732 and luteal phase serum progesterone measurements would enhance overall research quality [24].
733 Further, a shift towards prospective monitoring instead of retrospective self-report would further
734 strengthen the level of evidence [134].

735

736 Menstrual history questionnaires were also used to identify HMB in the included studies, with
737 questionnaire items reflecting a specific definition/criterion of HMB. In recent times, consistent with the
738 ACOG definition of HMB, the impact of HMB on an individual's quality of life has been used to guide its
739 diagnosis and treatment in clinical practice [136]. The presence of blood clots and period lengths greater
740 than 7 days are also recognised as independent predictors for HMB [137]. Given that there is no
741 standardised definition of HMB within the literature, it is unsurprising that definitions/criteria varied
742 across studies. However, it is interesting to note that none of these studies used a pictorial blood loss
743 assessment chart (PBAC) [138] to identify athletes with HMB. PBACs are widely used as a semi-
744 objective method to assess menstrual blood loss by considering the number of sanitary items used and
745 the extent to which items are soiled with blood, clots and flooding [139,140]. Several PBAC tools, which
746 are self-administered and cost-effective, have been validated against the gold-standard alkaline
747 hematin technique [141]. It is also important to consider contextual factors when assessing HMB to
748 prevent an overestimation of prevalence. For instance, the occurrence of menstrual leaking/overflow
749 as a criterion to identify HMB should be considered within the context of how often the sanitary products
750 have been changed (e.g., higher risk of leaking/overflow with less frequent changing of sanitary
751 products). Similarly, while the passing of blood clots can be a useful marker for HMB, it should be noted
752 that postpartum women can pass blood clots in the weeks following delivery and this may not be
753 indicative of HMB [142]. To that end, the findings of this review reinforce the need for a universally
754 accepted definition of HMB to be identified and future studies examining HMB in athletes should
755 consider the use of PBACs to determine menstrual blood loss while taking into consideration relevant
756 contextual factors.

757
758 Prospective daily monitoring of symptoms for two consecutive MCs has been considered the gold
759 standard method to assess PMS/PMDD [42,143]. This assessment method, in the form of a daily
760 symptoms log, was used in three out of seven studies [52,53,57] examining PMS, while the remaining
761 studies [89,90,97,102] identified the condition retrospectively using cross-sectional questionnaires.
762 While the core element of diagnosing PMS/PMDD is daily, prospective self-report, a clinical evaluation
763 by a trained medical professional is crucial. Indeed, an evaluation by a medical professional to diagnose
764 MC disorders is part of the criteria to achieve the 'gold' tier for menstrual irregularities studies as outlined
765 by a ranking system established to assess studies with female participants [33]. While medical doctors

766 were involved in 59.6% of the included studies as implied by authorship, only two studies explicitly
767 reported that the conditions were diagnosed through medical/gynaecological screening by a medical
768 professional within the study and met the 'gold' tier. In the remaining studies that were categorised
769 within the 'bronze' tier, conditions were either self-reported without a medical diagnosis, or it was not
770 specified if/how the condition was diagnosed. To reduce ambiguity, it would be beneficial for future
771 studies to explicitly state the confirmation of a condition from a medical professional, where appropriate.

772

773 It is also important to highlight that only seven studies reported that participants were confirmed as non-
774 HC users for at least three months prior to participation. This suggests that the majority of the studies
775 may not have considered the timescale (i.e., up to three months) needed for eumenorrhea to be re-
776 established in past HC users [15], which may result in an overestimation of the prevalence of MC
777 disorders as the naturally irregular cycles of non-HC users (who have recently ceased HC use) may be
778 misinterpreted as a dysfunction.

779

780 Taken together, the evaluation of methods used in studies investigating MC disorders seems to echo
781 recent statements on the inadequate and varied methodological practices associated with MC-based
782 research in sport and exercise science [15,24]. Apart from adopting the recommended methodological
783 guidelines, a focus should be on developing and validating a fit-for-purpose tool to screen for MC
784 disorders (beyond those associated with the Triad) among female athletes that could be used
785 consistently within research and applied settings.

786

787 **4.4. Limitations**

788 While this systematic review was the first to examine the prevalence of MC disorders in competitive
789 female athletes, several limitations were identified. Firstly, the definitions included in some of the studies
790 of this review were slightly varied or broad but were deemed acceptable as they did not substantially
791 deviate from the standard definitions. Furthermore, the exclusion of these studies may compromise the
792 representation of potentially relevant results. For example, excluding studies which used a threshold
793 age of 16 years could lead to a misestimation of the prevalence of primary amenorrhea. Therefore, to
794 achieve a balance between accuracy and representation, it was determined that the inclusion of these

795 studies was appropriate. Nonetheless, it is acknowledged that their inclusion may partly explain the
796 between-study heterogeneity in prevalence observed.

797

798 Considering the small number of studies within each MC disorder and the methodological variation
799 across studies, meaningful findings based on sub-group comparisons could not be produced. This
800 includes factors that may influence menstrual function or the reporting of MC disorders, such as age,
801 sport and culture. For example, age-group differences in MC disorders prevalence may be observed
802 due to reasons such as 1) the immaturity of the HPO axis leading to infrequent menstruation during the
803 initial menarcheal years in adolescents [144,145], and 2) a wider range of causes for amenorrhea, such
804 as endometrial hyperplasia, being associated with adults as compared to adolescents [1,146]. As
805 findings were not readily comparable between age groups, this review was not able to provide insights
806 on potential age-group differences in the prevalence of MC disorders. In addition, studies included in
807 this review were conducted across different countries, in which the reporting of MC disorders would
808 depend on the societal ideas of menstruation and what constitutes 'normal' within the culture. With
809 menstruation remaining a taboo topic in certain cultures, along with the societal normalisation of
810 menstrual pain, the prevalence of MC disorders may have been underreported [147–150].

811

812 As the use of HCs results in a significantly different hormonal milieu from that of eumenorrheic athletes
813 this review did not include HC users (i.e., HCs override the naturally occurring endogenous ovarian
814 hormone levels and their usage results in an altered endogenous and exogenous hormonal profile).
815 The exclusion of HC users could therefore lead to potential biases. Beyond contraceptive purposes,
816 HCs are also used to manage MC disorders such as HMB, irregular MCs and negative premenstrual
817 symptoms [151–153]. Given that the reason for HC use could be due to MC disorders itself, and that
818 the use of HCs is likely to mask any presence of MC disorders, an exclusion of HC users might lead to
819 a general underrepresentation of female athletes with MC disorders. Moreover, evidence suggests that
820 almost 50% of female athletes are HC users and may also experience side effects/symptoms [69].
821 Therefore, future research including female athletes using HCs would be beneficial for the female
822 athletic population at large.

823

824 **5. Conclusions**

PREVALENCE OF MENSTRUAL CYCLE DISORDERS AND SYMPTOMS IN ATHLETES

825 Research on MC disorders in competitive female athletes has often focused on conditions associated
826 with the Triad and Relative Energy Deficiency in Sport (RED-S) such as oligomenorrhea and
827 amenorrhea. The results of this systematic review indicate that other MC disorders are also
828 commonplace among female athletes, particularly dysmenorrhea, highlighting the need for more
829 research examining MC disorders beyond that of the Triad and RED-S. The current review also
830 identified shortcomings in methodological standards in studies examining MC disorders, specifically the
831 lack of uniformity in definitions and methods used to assess MC disorders among female athletes, which
832 should be improved upon considering recent methodological recommendations. From a practical
833 standpoint, the logistical hurdles of in-depth clinical/laboratory evaluation by a medical doctor and blood
834 sampling cannot be underestimated as athletes/sports teams may not have access to these resources.
835 To that end, it would be valuable for coaches, sports practitioners and athletes to have a working
836 knowledge of medical issues specific to the female athlete, including the ability to recognise abnormal
837 MC profiles. To support coaches and sports practitioners on the field, a fit-for-purpose tool to screen for
838 a range of MC disorders among female athletes should be developed and administered periodically
839 alongside other athlete monitoring measures, including MC tracking. This review further supports expert
840 recommendations for athletes to monitor their individual MCs prospectively using a combination of
841 calendar counting, urinary ovulation test kits, and a mid-luteal blood sample for the determination of
842 progesterone, such that anovulatory and LPD cycles can be determined [15]. Notably, given the
843 prevalence of dysmenorrhea and MC-related symptoms among female athletes, it would be meaningful
844 for athletes to track the occurrence and severity of MC-related symptoms, and how their perceived and
845 actual ability to train and perform may be impacted at various points of their MC. Future studies on MC
846 disorders should examine the prevalence and impact of both negative symptoms and positive effects
847 across the MC, as well as the influence of rapid weight loss and weight fluctuations on menstrual
848 function in female combat/weight-making sports athletes. Further, as oligomenorrhea and amenorrhea
849 appeared to be more prevalent in higher-calibre athletes than lower-calibre athletes within this review,
850 future research examining the relationship between athletic calibre, associated risks and underlying
851 mechanisms of MC disorders in athletes may contribute to the understanding of MC disorders in
852 sportswomen and provide better risk stratification. Finally, the absence of studies examining subtle MC
853 disorders (e.g., LPD and anovulation) means that further research is needed before robust conclusions
854 about the prevalence of MC disorders in female athletes can be drawn.

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