

1 **TITLE**

2 Fuelling performance of female soccer players: appraisal of the likelihood of between-sex
3 differences in carbohydrate requirements

4
5 **RUNNING TITLE**

6 Appraisal of sex differences in fuelling requirements

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53 **ABSTRACT**

54 Skeletal muscle glycogen depletion is considered one of the key contributing factors
55 toward fatigue and associated performance decrements in soccer. Ensuring adequate
56 carbohydrate availability before and during training and matches is widely-advised as a priority
57 for players, and for that reason, carbohydrate-based fuelling guidelines for soccer have been
58 developed. However, the lack of female-specific research used to inform the development of
59 these guidelines raises important questions regarding their applicability for female soccer
60 players. This review critically appraises the likelihood of there being between-sex differences
61 in carbohydrate requirements for soccer performance. Males and females exhibit differences in
62 substrate utilisation during exercise, substrate storage capacity, relative quantities of fat and
63 fat-free mass, and running demands of soccer match-play, but the extent to which these
64 differences translate into practically-meaningful differences in carbohydrate requirements for
65 soccer performance remains unclear. Based on current evidence, we consider it premature to
66 suggest that female players will require sex-specific guidelines in relation to carbohydrate-
67 based fuelling strategies for performance in soccer. However, intervention studies in female
68 players investigating carbohydrate-based fuelling strategies before and during soccer match-
69 play or simulated soccer protocols are warranted.

70

71 **KEYWORDS**

72 football; muscle glycogen; running demands; substrate utilisation; team sport;

73 **Introduction**

74 Carbohydrate (CHO) is the primary fuel source for skeletal muscle during high-
75 intensity exercise (Anderson et al., 2022; Hargreaves & Spriet, 2020; Vigh-Larsen et al., 2021).
76 Soccer is characterised by repeated high intensity activities interspersed with prolonged periods
77 of low- to moderate-intensity activity, resulting in a substantial reliance on glycogen stores in
78 both skeletal muscle and the liver (Krustrup et al., 2022; Krustrup et al., 2006; Vigh-Larsen et
79 al., 2021). For example, skeletal muscle glycogen concentrations can decline by ~40% from
80 pre- to post-match (Krustrup et al., 2022; Krustrup et al., 2006), with 80% type I fibres and
81 69% type II fibres being observed to be “almost empty” or “completely empty” of glycogen in
82 one study of female soccer players (Krustrup et al., 2022). Therefore, muscle glycogen stores
83 are inherently limited in their capacity relative to the fuel demands of training and matches
84 (Krustrup et al., 2006; Vigh-Larsen et al., 2021). Consequently, ensuring adequate CHO
85 availability before and during training and matches is often advised as a key priority for players
86 to support performance (Collins et al., 2021). Current guidelines for acute intake of CHO before
87 and during soccer performance are shown in **Table 1**.

88

89 **Table 1.** Current guidelines on carbohydrate-based fuelling strategies for before and during
90 soccer performance (Collins et al., 2021)

91

Strategy/time period	Current recommendations
Carbohydrate loading	6 to 8 g/kg BM on MD-1
MD	6 to 8 g/kg BM on MD
Pre-match	1 to 3 g/kg BM 3 to 4 hours before kick-off
During exercise	~30 to 60 g after warm-up and at half-time

92 Abbreviations: g/kg BM, grams of carbohydrate per kilogram of body mass; MD, match day.

93

94 Although these guidelines are intended to apply to both male and female players, the
95 authors acknowledge that the data on female soccer players is sparse (Collins et al., 2021).
96 Indeed, our recent audit revealed that females accounted for less than 4% of participants in the
97 evidence base that informed the development of these guidelines (McManus et al., 2025). The
98 lack of female-specific research in this area raises important questions regarding the
99 applicability of existing CHO guidelines for female soccer players (McManus et al., 2025;
100 Randell et al., 2021), given that there are various physiological and metabolic differences
101 between males and females that could impact CHO-based fuelling requirements for soccer
102 performance. Males and females exhibit differences in substrate utilisation during exercise
103 (Lundsgaard & Kiens, 2014), muscle fibre composition and capillarisation (Haizlip et al.,
104 2015), substrate storage capacity (Tarnopolsky et al., 1995), relative quantities of fat and fat-
105 free mass (FFM) (Randell et al., 2021; Sebastiá-Rico et al., 2023), and physical demands of
106 soccer match-play (Datson et al., 2017; Ju et al., 2025) which may influence dietary needs,
107 nutrient-exercise interactions, and performance outcomes. Therefore, this review critically
108 appraises whether these differences are likely to result in between-sex differences in CHO-
109 based requirements for fuelling soccer performance, as well as considering the extent to which
110 any differences could translate into practically-meaningful differences in CHO requirements
111 between male and female soccer players.

112

113 **Differences in substrate utilisation between males and females**

114 Developing nutrition recommendations for soccer players necessitates a comprehensive
115 understanding of the types and quantities of energy substrates utilised during match-play. To
116 date, no study has directly compared substrate utilisation between male and female athletes
117 during soccer match-play or a simulated soccer protocol. More broadly, between-sex
118 comparisons of substrate metabolism during any intermittent-type running protocol remain

119 scarce in the literature. Consequently, findings from continuous running protocols, and
120 continuous and intermittent cycling protocols must be relied upon to assess potential between-
121 sex differences in substrate utilisation during exercise. Evidence from controlled laboratory
122 settings consistently shows that when matched for training status and $\dot{V}O_2\text{max}$, and when
123 exercising at the same relative intensity, females tend to exhibit greater reliance on fatty acids
124 during prolonged submaximal aerobic exercise compared with males (Carter et al., 2001;
125 Montero et al., 2018; Roepstorff et al., 2006; Tarnopolsky et al., 1990). A whole-room
126 calorimeter study demonstrated that, relative to FFM, females exhibited 32% and 90% greater
127 rates of fat oxidation than males during low (40% $\dot{V}O_2\text{max}$) and high-intensity (70% $\dot{V}O_2\text{max}$)
128 exercise, respectively (Melanson et al., 2002). Even when controlling for differences in diet,
129 training status, and menstrual cycle phase, males have been shown to utilise 25% more muscle
130 glycogen, and exhibit higher respiratory exchange ratios than females during prolonged
131 exercise at $\sim 65\%\dot{V}O_2\text{max}$ (Tarnopolsky et al., 1990). Other studies have reported similar
132 observations throughout a range of submaximal exercise intensities up to $\sim 85\%\dot{V}O_2\text{max}$
133 (Chenevière et al., 2011; Venables et al., 2005), and this pattern persists across other modes of
134 exercise, such as resistance exercise (Sarafian et al., 2016). Results from sprint cycling
135 protocols corroborate these findings, with the exercise-induced reduction in muscle glycogen
136 being smaller by $\sim 40\%$ in type I muscle fibres in females compared to males in response to
137 single or repeated 30 s sprints, whereas the reduction in type II muscle fibres was similar across
138 sexes (Esbjörnsson-Liljedahl et al., 2002; Esbjörnsson-Liljedahl et al., 1999). Between-sex
139 differences in substrate utilisation have been proposed as being largely oestrogen-mediated
140 (Tarnopolsky, 2008), and therefore, may be influenced by menstrual cycle phase and hormonal
141 contraceptive use. Broad differences between sexes such as greater intramuscular triglyceride
142 storage, and cell membrane and mitochondrial fatty acid transport proteins, as well as β -
143 oxidation enzymes (e.g., medium chain fatty acyl-CoA dehydrogenase) and pyruvate

144 dehydrogenase kinase 4, are likely regulated by oestrogen acting through oestrogen receptor α -
145 mediated mechanisms, which results in greater skeletal muscle fatty acid oxidation capacity
146 (Campbell & Febbraio, 2001; Cano et al., 2022; Oosthuysen & Bosch, 2012). Additionally,
147 elevated concentrations of oestrogen, as observed during the late follicular and mid-luteal
148 phases of menstrual cycle, can reduce reliance on CHO utilisation due to reduced hepatic
149 glycogen utilisation and insulin-mediated storage, therefore shifting metabolism towards
150 greater contribution from fat through free fatty acid mobilisation and oxidation (Cano et al.,
151 2022; Friedlander et al., 1999). Hormonal contraception typically reduces endogenous
152 concentrations of 17-beta oestradiol and progesterone (compared with the mid-luteal phase of
153 the menstrual cycle), which may have implications for substrate utilisation (Elliott-Sale &
154 Hicks, 2018). Yet, the independent effect of progesterone on substrate utilisation during
155 exercise is still uncertain. However, one study involving eumenorrhoeic women compared the
156 effects during 60 min of submaximal exercise ($\sim 60\% \dot{V}O_{2\max}$) pharmacologically-
157 administered oestrogen alone versus a combination of oestrogen and progesterone (D'Eon et
158 al., 2002). The results showed that total CHO oxidation was higher when both hormones were
159 administered (oestrogen: 1.05 ± 0.02 g/min vs. oestrogen plus progesterone: 1.27 ± 0.04 g/min),
160 and the greater muscle glycogen utilisation also observed was associated with lower free-fatty
161 acid concentrations. These data suggest that progesterone may attenuate the lipolytic effects of
162 oestrogen (D'Eon et al., 2002).

163 Notably, in many studies reporting between-sex differences in substrate utilisation,
164 participants were exercised in a fasted state. In studies where participants were fed directly
165 before (0 to 3 h), or during, the exercise protocol, no clear differences in the relative
166 contribution of CHO or fat to energy provision were observed (Harger-Domitrovich et al.,
167 2007; M'Kaouer et al., 2004; Pettersson et al., 2019). Therefore, because players are likely to
168 play competitive soccer matches in a fed state, the reported differences in substrate utilisation,

169 i.e. greater reliance on fat in females during fasted exercise, is unlikely to be a relevant
170 consideration to fuelling strategies for match-play. In fact, in studies where CHO was
171 administered during exercise, the percent energy contribution from exogenous CHO sources
172 has been shown to be up to 25% higher for females compared to males (Riddell et al., 2003;
173 Tremblay et al., 2010). As such, ensuring adequate CHO intake directly before and during
174 soccer matches, consistent with current guidelines (Collins et al., 2021) (**Table 1**) may be of
175 particular importance for female players.

176 When expressed in absolute terms (g/min), similar exogenous CHO oxidation rates
177 have been observed in males and females when CHO is ingested during exercise (Wallis et al.,
178 2006). From a practical perspective, this observation implies that the current CHO-based
179 fuelling guidelines for during matches (i.e. ~30 to 60 g CHO after the warm-up, and at half-
180 time) (Collins et al., 2021) may be adequate for both male and female soccer players. However,
181 a recent study observed that larger athletes (BM >70 kg; 45±13 g/h) had a higher rates of
182 exogenous glucose oxidation (mean difference: 13 g/h) than smaller athletes (BM <70 kg; 33±8
183 g/h) during 120 min of cycling at 95% of lactate threshold (Ijaz et al., 2024). This difference
184 may have been partly due to differences in absolute exercise intensity as much as differences
185 in BM itself, and is somewhat in contradiction to a previous analysis that showed no
186 relationship between BM and rates of exogenous CHO oxidation (Jeukendrup, 2010). Despite
187 those caveats, a BM-dependent effect on exogenous CHO oxidation would translate as female
188 players requiring smaller amounts of CHO during exercise given their typically smaller body
189 sizes, but this contention would need be explored specifically in the context of soccer training
190 and match-play.

191 Notably, the CHO intake practices of elite female soccer players during match-play
192 have been reported as considerably less than current recommendations. For example, only 3
193 out of 19 Spanish first division players consumed CHO during a competitive match, with an

194 average group intake of just 0.9 g/h (Tarnowski et al., 2022). In comparison, average group
195 intakes of 32 g/h and 17 g/h have been observed during competitive matches in male English
196 Premier League players (Anderson et al., 2017; Kasper et al., 2024). Possible explanations for
197 the low intakes in female players may be the fear, or actual experience, of gastrointestinal
198 problems during matches. Largely due to their smaller stomachs (Cox, 1945), females are likely
199 to exhibit greater postprandial fullness and frequency and/or severity of gastrointestinal
200 symptoms following consumption of a given food volume (Tiller et al., 2021). However,
201 female athletes accustomed to feeding during exercise are able to tolerate greater intakes of
202 food and fluid during training and competition (Martinez et al., 2023). Accordingly,
203 implementing targeted nutrition ‘training’ of the gut (Jeukendrup, 2017), by ensuring repeated
204 exposure to CHO intake in training and matches, may help improve tolerance and optimise
205 fuelling practices during matches in this population.

206

207 **Differences in substrate storage between males and females**

208 On the day prior to a match (“MD-1”), a strategy known as “CHO loading” that targets
209 CHO intakes as high as 6 to 8 g/kg body mass (BM) is recommended to elevate glycogen stores
210 in skeletal muscle and liver (Collins et al., 2021) (**Table 1**). These guidelines are suggested to
211 be applicable for both male and female soccer players, but no study has yet investigated the
212 impact of CHO loading on muscle glycogen concentrations or performance in female soccer
213 players (McManus et al., 2025), although CHO loading is generally recognised to benefit
214 exercise performance more broadly (Larrosa et al., 2025). A difference between sexes in the
215 ability to store muscle glycogen when following a CHO loading protocol was previously
216 reported (Tarnopolsky et al., 1995). After 3 days of CHO loading, which increased CHO intake
217 from 55% to 75% of habitual energy intake (equating to an average CHO intake of 6.6 g/kg in
218 females and 8.4 g/kg in males during the protocol), a ~150 mmol/kg dw difference in glycogen

219 concentration in the vastus lateralis muscle of the thigh was observed between male
220 (565 mmol/kg dw) and female (409 mmol/kg dw) athletes (Tarnopolsky et al., 1995).
221 However, the same authors later demonstrated that when the males and females consumed a
222 similar absolute quantity of CHO (males: 600 g CHO equating to 8 g/kg BM vs. females: 540 g
223 CHO equating to 9 g/kg BM), no differences in muscle glycogen concentration were evident
224 (Tarnopolsky et al., 2001). Furthermore, when using a higher intake of CHO, muscle glycogen
225 concentrations were found to increase to a similar extent in both males (~68% increase) and
226 females (~79% increase) following a 3 day CHO loading protocol (12 g/kg lean body mass,
227 equating to ~10.5 g/kg BM for males and ~ 9.9 g/kg BM for females) (James et al., 2001).
228 Overall, these findings suggest that the capacity to store glycogen is not different between the
229 sexes once the intake of CHO is sufficient, with 8-10 g/kg BM of CHO often being referred to
230 as the “CHO loading threshold” (Bergström et al., 1967; Burke & Hawley, 1999; Wismann &
231 Willoughby, 2006). These interpretations should, however, be considered in light of there being
232 a limited number of studies, generally small sample sizes, and the general lack of
233 standardisation or monitoring of menstrual cycle phase in these studies.

234 Nonetheless, if such a threshold exists, it remains uncertain whether CHO intakes on
235 the lower end of the MD-1 CHO guidelines for soccer (i.e. 6 to 8 g/kg BM) would be sufficient
236 to achieve the ergogenic benefits of CHO loading in female players. In studies examining the
237 effects of CHO loading in male soccer players, all protocols have included a daily intake of at
238 least 7 g/kg BM of CHO (Abt et al., 1998; Balsom et al., 1999; Bangsbo et al., 1992; Hiromatsu
239 et al., 2023; Kazemi et al., 2023; Souglis et al., 2013). Furthermore, the majority of CHO
240 loading studies to date have used 2 to 4 day protocols under controlled conditions with limited
241 exercise, a pattern that does not reflect the typical training and fuelling protocols in professional
242 soccer players (Anderson et al., 2022; McHaffie et al., 2024; Morehen et al., 2022). For
243 example, match-day preparations in soccer often involve only one day of high CHO intake

244 (Anderson et al., 2022). Observations in female soccer players show markedly lower pre-match
245 resting muscle glycogen concentrations (~400 mmol/kg dw) (Krustrup et al., 2022) compared
246 to values previously reported in extended CHO loading studies in well-trained females, even
247 when CHO intake has only been moderate e.g. ~600 mmol/kg dw after 4.7 g/kg/d for 4 days
248 (Walker et al., 2000). Whether the short duration increase in CHO intake undertaken by soccer
249 players in practice would result in meaningfully different pre-match muscle glycogen
250 concentrations between male and female players is unknown.

251 In male soccer players, CHO loading strategies have been shown to delay fatigue and
252 enhance the capacity for intermittent high-intensity exercise in some studies (Balsom et al.,
253 1999; Bangsbo et al., 1992; Kazemi et al., 2023; Souglis et al., 2013), though not all have
254 reported such benefits (Abt et al., 1998; Hiromatsu et al., 2023). A key limitation of several
255 CHO loading studies in male soccer is the absence of a standardised pre-exercise meal. Given
256 the established impact of pre-exercise CHO intake on performance (Burke & Hawley, 1999),
257 absence of a standardised meal or major differences in CHO content of the pre-exercise meal
258 introduces a confounder and a question whether any effects observed from these studies can be
259 attributed to the CHO loading protocol alone. Furthermore, none of the existing CHO loading
260 studies in soccer reported including CHO ingestion during exercise (Abt et al., 1998; Balsom
261 et al., 1999; Bangsbo et al., 1992; Hiromatsu et al., 2023; Kazemi et al., 2023; Souglis et al.,
262 2013), despite UEFA guidelines recommending 30 to 60 g of CHO immediately before and
263 during soccer performance (Collins et al., 2021). Notably, a previous study in cycling (100 km
264 time trial) reported negligible performance differences following a CHO loading protocol (9
265 vs. 6 g/kg/d for 3 days; muscle glycogen, 572 ± 107 vs. 485 ± 128 mmol/kg dw) when participants
266 consumed CHO during exercise (1 g/kg BM), suggesting that CHO intake during exercise may
267 mitigate the effects of commencing exercise with lower pre-exercise glycogen concentrations,
268 or alternatively that the difference in muscle glycogen achieved with 9 g/kg/d compared 6

269 g/kg/d was not enough to impact performance (Burke et al., 2000). Taken together, these
270 observations highlight the need for better-controlled CHO loading studies in soccer across both
271 sexes.

272 One issue when determining the CHO requirements of female soccer players around
273 competition is the lack of studies investigating the contribution of muscle glycogen to energy
274 provision during soccer match-play or simulated soccer protocols in this population.
275 Furthermore, the glycogen cost of soccer match-play has not yet been directly compared
276 between male and female players. However, results from separate investigations demonstrate
277 similar reductions in muscle glycogen concentrations between pre and post-match in males and
278 females ($\Delta 194$ mmol/kg dw vs. $\Delta 173$ mmol/kg dw, respectively), with largely similar resting
279 glycogen concentrations observed (449 ± 22 vs 409 ± 62 mmol/kg dw) (Krustrup et al., 2022;
280 Krustrup et al., 2006). Although previous investigations have reported lower resting muscle
281 glycogen concentrations in females compared to males (~ 300 vs. ~ 500 mmol/kg dw) (Impey
282 et al., 2020), albeit not in soccer players specifically, this difference likely reflects the lower
283 habitual CHO intakes in female athletes (~ 3 to 4 vs. ~ 4 to 6 g/kg BM/d) (de Sousa et al., 2022;
284 García-Rovés et al., 2014). In studies that control for daily CHO intake and physical activity
285 levels, similar resting muscle glycogen concentrations have been observed between sexes
286 (James et al., 2001; Tarnopolsky et al., 2001). Furthermore, the lack of between-sex differences
287 in GLUT-4, hexokinase and glycogen synthase expression and activities also suggests limited
288 differences in muscle glucose delivery and storage between the sexes (Lundsgaard & Kiens,
289 2014).

290 Lastly, an important question specific to female players is whether the capacity to store
291 muscle glycogen is altered during specific phases of the menstrual cycle. Previous studies
292 suggest that females can supercompensate muscle glycogen consequent to CHO loading in
293 either phase of the menstrual cycle, at least when CHO is high (>8 g/kg BM) (McLay et al.,

294 2007; Nicklas et al., 1989), but resting glycogen concentration has been observed to be lower
295 during the follicular phase compared to the luteal phase (443 vs. 391 mmol/kg dw, respectively)
296 (Hackney, 1990). Practically, this observation may mean that female players should aim to
297 ingest the upper limits of recommendations (~8 g/kg BM) prior to matches, and pay particular
298 attention to doing so during the follicular phase of their menstrual cycle (Moore et al., 2022),
299 but this suggestion is tentative based on limited investigations to date as well the lack of
300 evidence for or against the performance benefits of CHO loading in female soccer players.
301 Similar recommendations may also be relevant for oral contraceptive pill users, particularly
302 during the 'inactive' pill phase, which is typically 4 to 7 days of pills without active hormones
303 that are taken following a 21 day course of 'active' pills, when circulating oestrogen and
304 progesterone concentrations resemble those of the early follicular phase (Cano et al., 2022).

305

306 **Differences in proportion of body mass as fat-free mass between males and females**

307 While evidence suggests minimal between-sex differences in the muscle's capacity to
308 store a given amount of glycogen per unit of muscle mass (James et al., 2001; Lundsgaard &
309 Kiens, 2014), one consideration could be whether variations in FFM as a proportion of total
310 BM could influence CHO requirements between male and female soccer players. CHO
311 recommendations for exercise are often expressed as grams per kilogram of BM (Collins et al.,
312 2021; Thomas et al., 2016). However, because FFM incorporates the lean tissue mass in the
313 form of skeletal muscle as the primary user of CHO during exercise, and in the form of skeletal
314 muscle and liver as the main storage sites for CHO, requirements for CHO intake could be
315 more closely related to FFM than to overall BM. By analogy, there is increasing discussion of
316 protein requirements being expressed on a per kg FFM basis (Dekker et al., 2022; Geisler et
317 al., 2016).

318 Dual-energy X-ray absorptiometry (DXA) is a widely-accepted and reliable non-
319 invasive method of estimating FM and FFM (Tewari et al., 2018). Average values for %FM
320 and FFM in elite male soccer players by DXA typically range from ~8% to ~13% and ~55 kg
321 to ~71 kg, respectively (Milanese et al., 2015; Milsom et al., 2015; Sebastián-Rico et al., 2023).
322 Data on elite female soccer players are more scarce, but average values for %FM and FFM of
323 between ~14.5% to ~22% and ~42.5 kg to ~49.5 kg, respectively, have been observed using a
324 combination of skinfold and bioelectrical impedance analysis methods (Petri et al., 2024;
325 Randell et al., 2021), while values of ~18% to ~23% and ~47 kg to ~49 kg were reported using
326 DXA (Morehen et al., 2023; Moss et al., 2020; Savolainen et al., 2023). Theoretically, the
327 lower proportion of BM as FFM in female players, on average, suggests reduced CHO
328 requirements in this population when expressed relative to BM.

329 **Table 2** illustrates this concept by considering the requirements applied to players with
330 BM of 60 and 80 kg, and %FM of 10%, 17.5% and 25%. The analysis suggests that compared
331 to a male of 10%FM, a female of 25%FM who consumes the same target of 7 g/kg BM of CHO
332 will consume 20% more CHO per kg FFM (7.8 vs. 9.3 g/kg FFM). If instead the target CHO
333 intake is based on g/kg FFM, that same female would only require 5.8 g/kg BM in order to
334 match the 7.8 g/kg FFM provided by the 7 g/kg BM.

335 These calculations illustrate that CHO requirements expressed per kg of BM may
336 overestimate the needs of female players relative to their glycogen storage capacity, and
337 support the rationale for using FFM-based requirements to more accurately individualise
338 nutrition strategies across sexes. Yet, in practical terms, the change in the example above would
339 mean consuming 348 g rather than 420 g of CHO as an absolute amount, with the 72 g
340 difference being the equivalent of a large CHO-based snack, or small CHO-based meal.
341 Therefore, whether these differences are physiologically-meaningful, or practically-actionable
342 would warrant further exploration, such that when combined with the inherent inaccuracies in

343 estimating %FM, there may be limited added value in routinely applying FFM-based
 344 recommendations over the more practical BM-based recommendations when working with
 345 female players.

346

347 **Table 2.** Sample calculations illustrating differences in carbohydrate requirements relative to
 348 body mass and fat-free mass as a function of different levels of body fat
 349

Body mass (kg)	Fat mass (%)	FFM (kg)	CHO (g/kg BM)	CHO (g)	CHO (g/kg FFM)
60	10.0	54.0	7.0	420	7.8
60	17.5	49.5	7.0	420	8.5
60	25.0	45.0	7.0	420	9.3
60	25.0	45.0	5.8	348	7.7
80	10.0	72.0	7.0	560	7.8
80	17.5	66.0	7.0	560	8.5
80	25.0	60.0	7.0	560	9.3
80	25.0	60.0	5.8	464	7.7

350 Note: The value of 7 g/kg BM is based on the recommendation that for the day prior to a match
 351 (MD-1), CHO intake should be at least 6 to 8 g/kg BM to elevate muscle and liver glycogen
 352 stores (Collins et al., 2021). Abbreviations: BM, body mass; FFM, fat-free mass; CHO,
 353 carbohydrate.

354

355 **Differences between males and females in physical performance and running demands of**
 356 **soccer match-play**

357 The question of whether CHO requirements differ between male and female soccer
 358 players is intrinsically linked to potential differences in the physical demands of match-play
 359 between men's and women's soccer. Efforts to quantify the metabolic demands of soccer
 360 match-play, particularly those related to CHO metabolism, are constrained by the limitations
 361 of existing measurement technologies. The most commonly reported metrics used to assess
 362 physical demands include total distance covered and distance covered within fixed speed or

363 velocity zones derived from wearable tracking technologies such as global positioning systems
364 (GPS) technology, or from semi-automated optical tracking technologies used at the elite level
365 (Harkness-Armstrong et al., 2022; Miguel et al., 2021).

366 Across international and domestic competitions, elite women's soccer players cover on
367 average ~9,500 to 9,900 m per match, or ~103 to 110 m/min (Harkness-Armstrong et al., 2022;
368 Pérez Armendáriz et al., 2024). Compared to men's soccer, female players cover less total
369 distance during elite competition at both international and domestic levels (Bradley et al., 2014;
370 Ju et al., 2025). While these differences are statistically significant, the effect sizes (ES) are
371 small-to-moderate (ES=0.5-0.8), equating to an average difference of ~400 m over 90 minutes,
372 or ~4.4 m/min (~0.26 km/h). Interpreted solely as average speed, total distance suggests a
373 relatively low overall movement intensity. However, the physiological demands of match-play
374 are not captured by average speed alone, but rather by how this distance is accumulated,
375 through repeated high-speed efforts, accelerations, decelerations, and directional changes,
376 which collectively underpin the metabolic demands and CHO requirements of match-play
377 (Paul et al., 2015).

378 One common issue in the comparison of the high-speed activity between men's and
379 women's soccer is the application of identical absolute speed zones to both male and female
380 players, despite well-documented sex differences in maximal running speed (Djaoui et al.,
381 2017; Hallam & Amorim, 2022; Haugen et al., 2012; Ju et al., 2025; McClelland & Weyand,
382 2022). As male players generally achieve higher average maximal running speeds by ~8 to
383 10% (Bradley & Vescovi, 2015; Ju et al., 2025), males are more likely to accumulate greater
384 distances in higher speed categories. This method skews the interpretation of high-speed efforts
385 in favour of male players, irrespective of relative exercise intensity (Bradley & Vescovi, 2015;
386 Gualtieri et al., 2023). To address this, some studies have implemented downward-adjusted
387 thresholds for female players (Harkness-Armstrong et al., 2022). These adjusted thresholds are

388 often arbitrarily selected, and there is currently no standardised or validated framework for this
389 approach (Gualtieri et al., 2023). As a result, speed zone definitions vary widely across studies,
390 limiting both cross-study and between-sex comparisons (Gualtieri et al., 2023; Harkness-
391 Armstrong et al., 2022). When adjusted thresholds are applied, the proportion of total distance
392 categorised as high speed is largely similar between men's and women's soccer (Bradley et al.,
393 2014; Ju et al., 2025). The integration of acceleration and deceleration metrics into match-play
394 analyses remains limited by inconsistent definitions, poor inter-unit reliability, and variability
395 introduced by differing proprietary data processing methods (Harkness-Armstrong et al., 2022;
396 Malone et al., 2017; Sweeting et al., 2017; Thornton et al., 2019; Whitehead et al., 2018). These
397 issues must be addressed before firmer conclusions as to the differences in the physical
398 demands of men's and women's soccer can be made.

399

400 **Assessing the metabolic demands of soccer match-play and the implications for** 401 **carbohydrate requirements**

402 While GPS-derived metrics are useful for general quantification of running demands,
403 they provide only a partial representation of the physical work performed (Bourdon et al.,
404 2017). Specifically, these metrics fail to capture the complex, multidirectional movement
405 patterns characteristic of soccer, including jumping, landing, rapid changes of direction and
406 high-intensity actions embedded within gameplay scenarios such as physical duels or pressing
407 (Malone et al., 2017; Reilly, 1997). Although most modern player tracking technologies are
408 technically capable of differentiating movement types, particularly GPS that incorporate
409 inertial measurement units (Beato et al., 2018; Whitehead et al., 2018), most analyses do not
410 distinguish between biomechanically distinct movement types, such as forward, lateral, and
411 backward running, each of which likely imposes different energetic demands (Flynn et al.,

412 1994). As a result, analyses based solely on linear, speed-based thresholds may underestimate
413 the true metabolic demands of match-play.

414 Accurately quantifying substrate utilisation, particularly CHO oxidation, during soccer
415 remains a methodological challenge. Indirect calorimetry, a gold standard for assessing energy
416 expenditure and substrate utilisation in steady-state exercise, is unsuitable in soccer due to the
417 intermittent and stochastic nature of match-play. The dynamic variation in movement intensity,
418 combined with the frequent transitions between low- and high-speed activity, precludes steady-
419 state conditions required for accurate calorimetric measurement (Jeukendrup & Wallis, 2005).
420 Indirect calorimetry also lacks the temporal sensitivity to capture the brief, high-intensity bursts
421 of activity that often exceed the velocity of $\dot{V}O_2\text{max}$, which are an inherent part of elite-level
422 match-play. These methodological constraints limit the capacity to measure real-time energy
423 expenditure and, by extension, the specific contribution of CHO oxidation during key phases
424 of performance.

425 Alternative methods commonly used to assess the demands of match-play, such as heart
426 rate monitoring and blood lactate sampling, also lack the sensitivity to differentiate between
427 movement types or accurately quantify energetic cost. Heart rate remains elevated for much of
428 match-play in both men's and women's soccer, typically averaging 80 to 90% of maximal heart
429 rate across both halves, even during periods of relatively low physical work (Bangsbo et al.,
430 2006; Krstrup et al., 2006). As a result, heart rate offers limited insight into the variability of
431 physical work or the intensity of discrete movement patterns. Similarly, blood lactate
432 concentration is a poor proxy for the metabolic demands of match-play, as measurements are
433 highly dependent on the intensity and type of activity performed immediately prior to sampling
434 (Bangsbo et al., 2006). This temporal limitation likely contributes to the wide range of values
435 reported at the end of playing halves (i.e. 1 to 12 mmol/L), making it an unreliable indicator of
436 the overall metabolic demands of match-play (Bangsbo et al., 2006; Krstrup et al., 2010).

437 While muscle biopsy studies have demonstrated substantial reductions in glycogen
438 content across both type I and II fibres by the end of match-play in men's and women's soccer
439 (Krustrup et al., 2022; Krustrup et al., 2006), it remains unclear to what extent these reductions
440 impair soccer performance (Bangsbo et al., 2006). In women's soccer, declines in sprint
441 performance were significantly correlated to reductions in muscle glycogen (Krustrup et al.,
442 2022), whereas no association was found in men's soccer despite similar reductions in muscle
443 glycogen content (Krustrup et al., 2006). Match-tracking data often show that high-speed
444 running and sprint outputs are greater during the first half of match-play (Di Salvo et al., 2009),
445 with the most demanding periods also occurring in the early stages of a game (Bradley &
446 Noakes, 2013; Harkness-Armstrong et al., 2022). This temporal decline in physical output is
447 often interpreted as fatigue, though this view likely oversimplifies the underlying causes, which
448 may also reflect other factors such as tactical pacing, substitution patterns, or in-game nutrition
449 practices (Carling, 2013). Without real-time markers of substrate utilisation, it is difficult to
450 isolate the role of CHO availability from other performance determinants.

451 Further complicating this picture is the lack of in vivo data on hepatic glucose output
452 and the limited understanding of glycogen compartmentalisation within the skeletal muscle.
453 Glycogen pools localised near mitochondria or sarcoplasmic reticulum may serve functional
454 roles beyond general energy provision, particularly in maintaining excitation-contraction
455 coupling during prolonged or repeated efforts (Vigh-Larsen et al., 2021). However, current
456 methodologies do not allow for the in-field assessment of these mechanisms, leaving a critical
457 gap in our understanding of the local metabolic environment within working muscle during
458 match-play. Collectively, the above limitations underscore the need for more nuanced and
459 context-sensitive tools to assess metabolic demands in soccer. Without methodologies capable
460 of capturing the true energetic cost of movement, particularly with respect to CHO utilisation,

461 our understanding of fuelling strategies, fatigue mechanisms, and recovery processes remains
462 incomplete.

463

464 **Concluding remarks**

465 There have been increasing calls to develop, or at least systematically evaluate, the need
466 for female-specific nutrition guidelines for soccer (Holtzman & Ackerman, 2021; Martinho et
467 al., 2024; Moore et al., 2022). While such efforts have the potential to support player health
468 and performance, it must be acknowledged that, in practice, elite players have been consistently
469 reported to not adhere to the existing evidence-based recommendations (Morehen et al., 2022;
470 Renard et al., 2021). For example, average CHO intakes around training and match play that
471 have been reported in female soccer players (3-4 g/kg BM) (Gravina et al., 2022; Martinho et
472 al., 2024; Morehen et al., 2022) are well below current recommendations (6-8 g/kg BM)
473 (Collins et al., 2021). Of relevance are recent data from elite youth female players in England
474 that indicate that higher CHO consumption is often perceived as contributing to increases in
475 fat mass and negative body image (McHaffie et al., 2022), which highlights a need to better
476 understand the psychosocial and environmental determinants of dietary behaviours in female
477 soccer players more broadly. From a practitioner perspective, challenges in eliciting
478 behavioural change among players, uncertainty regarding the scientific basis of existing CHO
479 guidelines, limited time and support, and a lack of autonomy over the nutrition service
480 provision, were all identified as barriers to implementing nutrition recommendations in elite
481 men's soccer (Costello et al., 2025). Similar research involving practitioners in the women's
482 game is necessary, particularly given that effective implementation of CHO guidelines requires
483 coordinated efforts between players, coaching staff, and nutritionists.

484 Although some between-sex differences in physiology, metabolism, and running
485 demands among soccer players exist, the currently-available evidence, although somewhat

486 limited, does not suggest that CHO-based requirements differ between male and female players
487 in obvious ways that would translate into practically-meaningful differences in fuelling
488 strategies being employed. However, variations in FFM as a proportion of total BM and its
489 potential implications for CHO storage and intake may warrant further investigation as an
490 influencing factor. Nonetheless, on the basis of the current evidence, we consider it premature
491 to suggest that female players will require sex-specific guidelines in relation to CHO-based
492 fuelling strategies for performance in soccer. Because of the inter-individual variability that
493 exists regardless of biological sex, an individualised approach to fuelling remains the
494 cornerstone to sports nutrition practice. However, given the dearth of sports nutrition research
495 in female athletes, intervention studies in female players investigating CHO-based fuelling
496 studies before and during soccer match-play or simulated soccer protocols are warranted.

497

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