Effect of a short-term low fermentable oligosaccharide, disaccharide, monosaccharide and polyol (FODMAP) diet on exercise-related gastrointestinal symptoms

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Abstract

Background: Research has demonstrated that low fermentable oligosaccharide, disaccharide, monosaccharide and polyol (FODMAP) diets improve gastrointestinal (GI) symptoms in irritable bowel syndrome sufferers. Exercise-related GI issues are a common cause of underperformance, with current evidence focusing on the use of FODMAP approaches with recreationally competitive or highly trained athletes. However, there is a paucity of research exploring the potential benefit of FODMAP strategies to support healthy, recreational athletes who experience GI issues during training. This study therefore aimed to assess whether a short-term LOWFODMAP diet improved exercise-related GI symptoms and the perceived ability to exercise in recreational runners.

Methods: Sixteen healthy volunteers were randomly assigned in a crossover design manner to either a LOWFODMAP (16.06 ± 1.79 g·d⁻¹) or HIGHFODMAP (38.65 ± 6.66 g·d⁻¹) diet for 7 days, with a one week washout period followed by a further 7 days on the alternate diet. Participants rated their gastrointestinal symptoms on an adapted version of the Irritable Bowel Syndrome-Severity Scoring System (IBS-SSS) questionnaire before and at the end of each dietary period. Perceived ability to exercise (frequency, intensity and duration) in relation to each dietary period was also rated using a visual analogue scale. Resting blood samples were collected prior to and on completion of each diet to determine plasma intestinal fatty acid binding protein (I-FABP) as a marker of acute GI injury.

Results: Overall IBS-SSS score significantly reduced in the LOWFODMAP condition from 81.1 ± 16.4 to 31.3 ± 9.2 (arbitrary units; \(P = 0.004\)). Perceived exercise frequency (\(z = 2.309, P = 0.02\)) and intensity (\(z = 2.687, P = 0.007\)) was significantly improved following a short-term LOWFODMAP approach compared to HIGHFODMAP. No significant differences were reported between dietary conditions for plasma I-FABP (\(P > 0.05\)).

Conclusions: A short-term LOWFODMAP diet under free-living conditions reduced exercise-related GI symptoms and improved the perceived ability to exercise in otherwise healthy, recreational runners. These findings may be explained by a reduction in indigestible carbohydrates available for fermentation in the gut. The therapeutic benefits of LOWFODMAP diets in recreational and trained athletes during sustained training periods warrants further investigation.

Keywords: FODMAP diet, Gastrointestinal symptoms, Nutrition, Recreational athletes

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**Introduction**

Fermentable oligosaccharides, disaccharides, monosaccharaides and polyols (FODMAPs) are short-chain carbohydrates that are widespread in the diet in foods such as fruits, vegetables, dairy, wheat, grains, legumes, and are commonly added to processed foods to improve palatability. The major types of FODMAPs known to be problematic are fructose, lactose, oligosaccharides and polyols, each of which has a distinct mechanism of action.

Fructose is absorbed in the small intestine by two carrier protein transporters, GLUT2 (in the presence of glucose) and GLUT5, which facilitate fructose diffusion across cell membranes [1, 2]. In some individuals, the limited availability of the GLUT5 transporter results in fructose malabsorption when it is present in excess of glucose [3–7]. Fructose malabsorption is commonly reported both within irritable bowel syndrome (IBS) sufferers (45%) and healthy individuals (34%) [5]. Lactose malabsorption occurs when there is insufficient lactase to break lactose down into its component sugars glucose and galactose [5].

Oligosaccharides are generally poorly absorbed, resulting in the undigested carbohydrates being fermented by gut bacteria [5, 8, 9]. This results in gas production and flatulence in both healthy and hypersensitive individuals which may instigate adverse symptoms [10]. Polyols do not have an associated active transport system and are thought to be absorbed by diffusion [6], which is variable across the intestine and between individuals [11]. If the polyol is too large for diffusion, malabsorption may occur [8], resulting in fermentation or an increased osmotic load leading to fluid retention in the small intestine [12, 13].

Cumulatively, the malabsorption of these short chain carbohydrates as part of a habitual diet may result in increased small intestinal water volume which can affect gut motility [6, 8]. It has previously been established that altered gut motility is associated with symptoms which are analogous to IBS and exercise including nausea, diarrhea and urge to defecate during exercise [14, 15]. A LOWFODMAP diet has been established as an evidence-based approach to reduce symptoms in approximately 75% of patients diagnosed with IBS [16]. Sixty to 70% of patients report a worsening of IBS symptoms after habitual meals [17], and consequently will eliminate foods that they believe trigger their symptoms [18, 19].

It is possible that hypersensitive individuals are more susceptible to an adverse reaction to a HIGHFODMAP diet as a result of the mechanistic changes during exercise [14, 15, 20, 21] which can ultimately impact on training and/or performance. Increasing the intensity and duration of exercise corresponds with slower gastric emptying and potential for structural epithelial damage, tight-junction disruption and transient luminal permeability, as demonstrated through acute elevated levels of plasma intestinal-fatty acid binding protein (I-FABP) [22–24]. It is therefore relevant to consider whether a short-term LOWFODMAP diet impacts on habitual levels of I-FABP, or indeed whether markers of intestinal damage are exacerbated or sustained as a result of a HIGHFODMAP diet.

It has been reported that 30-50% of athletes cite GI issues as one of the most common causes of underperformance in endurance events [15]. This likely explains why athletes may eliminate food groups they believe cause GI distress [25–28], with a gluten free diet becoming one of the most common approaches reported [27]. However, there is little evidence that removal of gluten has any performance benefits for non-celiac athletes and it has been suggested that gluten may not be a specific trigger of GI symptoms once dietary intake of FODMAPs are reduced [29]. Recent evidence has focused on the use of FODMAP approaches to support recreationally competitive or highly trained athletes. However, there is a paucity of research exploring the potential benefit of FODMAP strategies to support healthy, recreational athletes who experience GI issues during endurance training. Therefore, the purpose of this randomised, crossover trial was to investigate the effect of altering FODMAP intake upon the GI symptoms reported by recreational athletes in free-living conditions during habitual training. It was hypothesised that a short-term LOWFODMAP diet would improve GI symptoms and the perceived ability to exercise.

**Materials/methods**

**Study design**

This study employed a randomised, crossover design. The study was conducted in accordance with the Declaration of Helsinki, and ethical approval was granted by the Faculty of Science and Technology Ethics Committee, Anglia Ruskin University (Project Number: FST/FREP/15/567). All participants provided written informed consent prior to study inclusion. All monitoring procedures took place in the Cambridge Centre for Sport and Exercise Sciences, Anglia Ruskin University under controlled conditions.

Participants were required to be healthy, recreationally active runners (training a minimum of 3 days per week with at least 3 months habitual experience, and satisfactorily complete a health screen questionnaire) and prepared to comply with study requirements. Ineligible participants were those with a known health condition (including persistent non-exercise related GI issues), current injury, or recent viral infection. Participants were required to only eat foods in conjunction with the lists provided for each 7 day period, and be prepared to weigh food and keep a detailed food log. All participants reported no known or diagnosed gut disorders, were not currently following a LOWFODMAP ketogenic or calorie restricted diet, were not currently taking antibiotics or probiotics, and had no known blood disorders or allergies.

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Participants
An *a priori* power calculation was undertaken based on the primary end point being the difference in IBS–SSS before and after the LOW\(_{\text{FODMAP}}\) diet. It was estimated that 11 participants were needed per dietary condition to have an 80% power to detect a difference within group of > 1 SD of IBS–SSS score using a paired t-test with a one-sided \(\alpha\) of 0.05 based on previous data [6]. Participants (\(n = 19\)) were recruited through personal contacts with local running clubs. One participant withdrew due to the burden of keeping a weighed food diary and two participants were excluded from the final analysis due to dietary non-compliance. Sixteen participants completed all aspects of the study (10 female, 6 male; age: 44 ± 10 years, height: 1.70 ± 0.78 m, body-mass: 69.2 ± 8.8 kg).

Eating plans and dietary intake
According to previous research which categorized carbohydrates as low or high FODMAP [30–33], two separate food lists were devised for this study containing either high or low FODMAP foods. To protect the integrity of the study, participants were instructed to follow the two diets (with an explanation that the type of carbohydrate was different) with no specific reference to FODMAPs in pre study information. For the purposes of protocol blinding lists were named A and B, although complete blinding was not feasible. In order to mimic dietary choice that athletes make in free-living conditions participants were free to select foods from the list and were individually advised to match their typical dietary and calorie intake and record via a weekly weighed food diary. Participants were provided with example diaries and individually instructed in diary completion, with emphasis on meal breakdown, portion size/weight and weighing procedure. Dietary analyses were undertaken by the same researcher for standardisation using Nutritics Professional Dietary Analysis software (Nutritics Limited, Dublin).

Experimental procedures
Participants attended the laboratory prior to and immediately after each dietary period, and were requested to be rested (no exercise) in the 24-hour period prior to all laboratory measures. For all visits, on arrival, participants rested for 5 minutes prior to fasted blood sample collection, and were then required to complete a symptom questionnaire (see below). Participants were randomised using a pseudo-random number generator (www.randomizer.org) to start on either the low or high FODMAP condition for 7 days based upon previously reported research [34]. All participants undertook a one week washout period between conditions (in a similar manner to previously reported research [10]) and were requested to return to their normal eating patterns during this period before undertaking the opposing dietary condition. Prior to starting, and throughout the study, participants were requested to continue their normal training routine.

Blood sampling and analysis
Upon arrival, a venous whole blood sample was collected from participants by a qualified phlebotomist into duplicate 4 ml K3EDTA vacutainers (Greiner Bio-One GmbH, Kremsmunster, Austria). Samples were centrifuged for 10 minutes at 3000 rpm, with aliquotted plasma pipetted into sterile, non-pyrogenic, polypropylene cyrovials (Fisherbrand, Fisher Scientific, Loughborough, UK) and immediately frozen at −80 °C for later assessment of I-FABP using an ELISA kit (Hycheck Biotechnology, Uden, the Netherlands; analytical measurement range: 47 to 3000 pg·ml\(^{-1}\); intra-assay variance: 3.2% at 360 pg·ml\(^{-1}\), 5.4% at 557 pg·ml\(^{-1}\) and 6.6% at 809 pg·ml\(^{-1}\)). Reagents were prepared in accordance with the manufactures instructions at room temperature. Duplicate plasma samples were thawed to room temperature (22 °C) and diluted 10-fold using the sample dilution buffer. I-FABP was extracted from the plasma samples by the addition of the following reagents to ELISA kit in the following order: diluted plasma samples; diluted tracer; diluted streptavidin-peroxidase. In between additions the tray was covered with foil, incubated at room temperature (22 °C) for 1 hour before washing. Finally tetramethylbenzidine (TMB) substrate was added to each well and the tray was incubated for 30 minutes at room temperature. The reaction was stopped with the addition of the stop solution and gently mixed. Samples were read on a spectrophotometer at an absorbance of 450 nm (Victor 3 multilabel plate reader, PerkinElmer Inc., Llantrisant, UK) and referenced against a calibration curve (logarithmic scale).

Gastrointestinal symptom monitoring
Prior to, and following each dietary period participants rated individual GI symptoms (i.e. bloating, abdominal pain, flatulence, belching, nausea, diarrhoea, defecation, urge to defecate and constipation) against a standardised 0–100 visual analogue scale (VAS) questionnaire (arbitrary units (au)), with no interference from the research team. Global IBS symptom severity scores (IBS-SSS) were based on accumulated results. Clinically significant change of symptoms was defined as > 20 au on the VAS scale [35]. Participants were also requested to rate their perception of their ability to exercise over the week in which each dietary period occurred. Having recorded their training, participants rated their exercise intensity, duration and frequency based on a category scale (no change 0, improved 1, worsened 2) in comparison to a typical training week. Following this, food diaries were collected and inspected for accuracy, detail and
compliance using a second pass interview approach between the researcher and participant.

**Statistical analysis**
Statistical analyses were performed using SPSS (IBM, Version 24.0). Normality of data were verified by the Shapiro-Wilks test. Outliers were identified by inspection of box plots > 1.5 IQR in SPSS. A repeated measures ANOVA was used to compare effects of dietary interventions (i.e. nutritional intake, IBS-SSS, 1-FABP) with Bonferroni post-hoc assessment where applicable. Where sphericity was violated a Greenhouse Geisser correction was applied. A dependent samples t-test was carried out to assess relative differences between diets where pertinent. Ability to exercise data was analysed using a Wilcoxon signed rank test. An alpha level of $P \leq 0.05$ was considered statistically significant for all tests. Data are presented as mean ± SE.

**Results**
**Dietary intake**
No significant differences in mean caloric intake were reported between dietary conditions, or in comparison to habitual intake ($F = 2.921, P = 0.07, \eta^2 = 0.173$; Table 1). When normalised for body-mass, mean habitual caloric intake ($34.12 \pm 2.48$ kcal·kg$^{-1}$·d$^{-1}$) was comparable with both LOWFODMAP ($29.04 \pm 1.88$ kcal·kg$^{-1}$·d$^{-1}$) and HIGHFODMAP ($32.53 \pm 2.08$ kcal·kg$^{-1}$·d$^{-1}$) conditions ($F = 3.053, P = 0.063, \eta^2 = 0.179$). For carbohydrate intake, a significant main effect was observed ($F = 7.091, P = 0.003, \eta^2 = 0.336$), with participants reporting consuming less total (Table 1) and relative carbohydrate intake during the LOWFODMAP condition compared with the HIGHFODMAP condition ($2.79 \pm 0.30$ g·kg$^{-1}$·d$^{-1}$ and $3.91 \pm 0.36$ g·kg$^{-1}$·d$^{-1}$ respectively, $P = 0.003$).

Total FODMAP intake (including relative to body-mass) was also statistically different between conditions ($F = 10.354, P < 0.0001, \eta^2 = 0.425$), with post-hoc analysis demonstrating the expected reduction with LOWFODMAP ($15.75 \pm 1.91$ g·d$^{-1}$) compared with both HIGHFODMAP ($38.59 \pm 6.48$ g·d$^{-1}$, $P = 0.004$) and habitual conditions ($28.04 \pm 4.33$ g·d$^{-1}$, $P = 0.045$). There were no significant differences reported for dietary fat (habitual: $1.48 \pm 0.17$ g·kg$^{-1}$·d$^{-1}$; LOWFODMAP $1.29 \pm 0.15$ g·kg$^{-1}$·d$^{-1}$; HIGHFODMAP $1.21 \pm 0.08$ g·kg$^{-1}$·d$^{-1}$; $F = 1.446, P = 0.253$) or protein intake (habitual: $1.34 \pm 0.10$ g·kg$^{-1}$·d$^{-1}$; LOWFODMAP $1.35 \pm 0.70$ g·kg$^{-1}$·d$^{-1}$; HIGHFODMAP $1.36 \pm 0.08$ g·kg$^{-1}$·d$^{-1}$; $F = 0.142, P = 0.798$) between experimental conditions or in comparison to habitual intake (total or relative to body-mass).

**Gastrointestinal symptom scores (overall)**
Mean gastrointestinal symptom scores (IBS-SSS) did not differ between conditions prior to each FODMAP diet ($P > 0.05$), although a wide variance of responses was noted between participants (mean: $66.1 \pm 16.3$ au; range 0–206 au). A significant diet x time interaction effect was found for IBS-SSS ($F = 6.98, P = 0.02, \eta^2 = 0.32$), with post-hoc analysis indicating a significant reduction in scores from $81.1 \pm 16.4$ au (pre) to $31.3 \pm 9.2$ au (post) with LOWFODMAP ($P = 0.004$; Fig. 1). Although a non-significant increase in IBS-SSS was reported with HIGHFODMAP from $51.1 \pm 15.7$ au (pre) to $104.0 \pm 25.0$ au (post; $P = 0.08$); overall end-point scores were significantly different between dietary conditions ($P = 0.007$). Expressed as relative change (Fig. 2), a significant difference was also reported between dietary conditions (mean difference $= -102.7 \pm 38.9$ au; $t = -2.64, P = 0.02$) in favour of an improvement in responses following a LOWFODMAP approach. Individual responses indicated that 69% of participants (11/16) reported positive effects of the LOWFODMAP diet, in contrast to 25% (4/16) on the HIGHFODMAP diet.

**Table 1** Mean dietary intake under habitual and FODMAP conditions

| Variable | Category | Habitual | LOWFODMAP | HIGHFODMAP |
|----------|----------|----------|-----------|------------|
| Total EI | (kcal·d$^{-1}$) | $2355.86 \pm 197.10$ | $1999.23 \pm 138.43$ | $2269.14 \pm 162.11$ |
|          | (kcal·kg$^{-1}$·d$^{-1}$) | $34.12 \pm 2.48$ | $29.04 \pm 1.88$ | $32.53 \pm 2.08$ |
| Carbohydrates | (g·d$^{-1}$) | $245.77 \pm 19.05$ | $193.53 \pm 21.56^a$ | $272.28 \pm 26.23$ |
|          | (g·kg$^{-1}$·d$^{-1}$) | $3.55 \pm 0.22$ | $2.79 \pm 0.30^b$ | $3.91 \pm 0.36$ |
| FODMAPs  | (g·d$^{-1}$) | $28.04 \pm 4.33$ | $15.75 \pm 1.91^a$ | $38.59 \pm 6.48$ |
|          | (g·kg$^{-1}$·d$^{-1}$) | $0.42 \pm 0.07$ | $0.23 \pm 0.03^a$ | $0.56 \pm 0.09$ |
| Protein  | (g·d$^{-1}$) | $92.13 \pm 7.82$ | $94.34 \pm 5.81$ | $94.64 \pm 6.54$ |
|          | (g·kg$^{-1}$·d$^{-1}$) | $1.34 \pm 0.10$ | $1.35 \pm 0.07$ | $1.36 \pm 0.08$ |
| Fat      | (g·d$^{-1}$) | $101.95 \pm 12.74$ | $88.48 \pm 9.88$ | $83.92 \pm 5.82$ |
|          | (g·kg$^{-1}$·d$^{-1}$) | $1.48 \pm 0.17$ | $1.29 \pm 0.15$ | $1.21 \pm 0.08$ |

Table 1 outlines mean habitual dietary intake and between dietary conditions with data expressed in total amounts per day. EI = energy intake. * denotes significant difference between LOWFODMAP and HIGHFODMAP conditions only ($P < 0.004$). + denotes significant difference to habitual diet ($P < 0.045$).
Gastrointestinal symptom scores (individual)

Table 2 demonstrates mean responses to individual GI symptoms across both dietary conditions. A significant diet x time interaction effect was found for pain ($F = 6.861$, $P = 0.019$, $\eta^2 = 0.314$) with post hoc analyses indicating that end-point scores were significantly different between dietary conditions (4.13 ± 2.52 au for LOW FODMAP and 22.50 ± 6.35 au for HIGH FODMAP respectively, $P = 0.003$), which coincided with significant within-group changes for both LOW FODMAP ($P = 0.031$) and HIGH FODMAP ($P = 0.028$). A significant main effect (time) was reported for flatulence ($F = 4.428$, $P = 0.05$, $\eta^2 = 0.228$), as well as a main effect (diet) for belching ($F = 5.686$, $P = 0.03$, $\eta^2 = 0.275$), although post-hoc analyses were not significant. A significant main effect (diet) was reported for bloating ($F = 6.186$, $P = 0.025$, $\eta^2 = 0.292$), with post-hoc analyses indicating that end-point scores were significantly different between dietary conditions (1.25 ± 0.72 au for LOW FODMAP and 12.69 ± 4.53 au for HIGH FODMAP respectively, $P = 0.021$). All symptoms except constipation and defeation recorded a decrease in score on the LOW FODMAP whilst all symptoms except flatulence recorded an increase in score on the HIGH FODMAP diet, however no other significant findings were reported.

FODMAP intake and self-reported ability to exercise

There was a statistically significant median difference in the perceived exercise frequency ($z = 2.309$, $P = 0.02$) and intensity ($z = 2.687$, $P = 0.007$) between a LOW FODMAP and HIGH FODMAP diet. No significant median difference was reported in perceived exercise duration on a LOW FODMAP and HIGH FODMAP diet ($z = 1.414$, $P = 0.157$). Participants were more likely to report that the ability to exercise improved on a LOW FODMAP diet (frequency (4/16), intensity (6/16)), and deteriorated on a HIGH FODMAP diet (frequency (4/16), intensity (9/16)).

FODMAP intake and intestinal fatty acid binding protein (I-FABP) levels

The effect of both FODMAP diets on I-FABP levels under resting conditions is shown in Fig. 3. A non-significant increase from 206.93 ± 7.27 pg·ml$^{-1}$ to 219.46 ± 10.42 pg·ml$^{-1}$ was noted for LOW FODMAP remaining within expected limits. I-FABP for HIGH FODMAP remained comparable across the intervention (218.21 ± 10.93 pg·ml$^{-1}$ to 222.60 ± 13.08 pg·ml$^{-1}$; $P > 0.05$). No significant interaction effects were reported ($P > 0.05$) between dietary conditions.

Discussion

This study aimed to investigate the perceived effect of acute FODMAP intake on GI symptom severity and ability to exercise in recreational athletes under free-living conditions. Whilst the clinical effectiveness of a LOW FODMAP diet in treating IBS is established in the literature [16, 28], research into the potential therapeutic effects in otherwise healthy, recreational athletes is limited. The main findings from the current study revealed that short-term LOW FODMAP intake significantly improved exercise-related GI symptoms in 69% of participants. These results support recent observations [36] demonstrating reductions in daily GI symptoms in trained athletes on a short-term (6-day) LOW FODMAP diet, as well as reduced GI symptom severity in case studies of a male [37] and female [38] runner. Consistency between these findings infers that both recreational and more trained athletes may benefit from self-prescribed LOW FODMAP approaches in the short-term, providing there is compliance with food choices. This may have implications for...
longer term FODMAP strategies during sustained training periods, which may provide adjunct nutritional support in maintaining training volume and/or intensity [20] particularly in symptomatic individuals who suffer from GI distress with exercise.

Whilst a LOW FODMAP approach appeared to result in improved scores for most individual symptoms, only responses to perceived pain and bloating were significantly different between conditions following the dietary interventions. This suggests that whilst an improvement in overall GI symptom severity may reflect accumulated reductions in individual symptom responses, the effects of a short-term LOW FODMAP diet may in fact be specific. The reported improvement in perceived pain, in conjunction with reduced experiences of bloating whilst on a LOW FODMAP diet is likely explained by a reduction in intestinal water volume and gas production [6, 10, 39]. Strategies to reduce or minimise such GI symptoms may be important for recreational athletes, especially considering the reported negative impact on exercise training and/or performance [15, 40, 41]. However, based on the wide inter-individual responses observed across conditions, such findings should be interpreted with caution.

An interesting observation from the current study was the improved perception of exercise frequency and intensity from participants whilst undertaking the LOW-FODMAP approach. Although this only reflected perceived changes in the short-term (7 days), this may have implications for sustained approaches where training routines may be disrupted (including volume and intensity) due to GI-related issues. Participants were requested to maintain their typical training routine throughout the study to assess whether perceived changes (in frequency, training duration or intensity) were related to the dietary intervention. Whilst a significant effect was observed for improved perception of exercise frequency and intensity following a LOW FODMAP approach, this only occurred in 25 and 38% of the participants respectively. Only one other study [36] appears to have attempted to standardise training sessions (albeit 2 sessions in a 6-day period) whilst participants undertook an acute LOW FODMAP or HIGH FODMAP diet. In this study, daily GI symptoms for flatulence, urge to defecate and diarrhoea were reportedly improved in the LOW FODMAP condition [36]. However, assessment of participants’ perception of training session ability in relation to the dietary approach was not considered. Further research to establish training related benefits of a LOW FODMAP strategy, particularly with symptomatic individuals, is therefore warranted.

Moderate to high-intensity exercise impacts on gastric emptying, GI transit and intestinal absorption due to GI hypoperfusion and ischemia [15, 22]. Provocation of luminal tight junction proteins (e.g. caludin and occludin) through increased expression of phosphorylation enzymes, reactive oxygen species (ROS) activity and cytokine mediators may lead to acute GI permeability

Table 2 Mean responses to individual GI symptoms across both dietary conditions

| Symptom           | LOW FODMAP | POST | HIGH FODMAP | POST | Interaction |
|-------------------|------------|------|-------------|------|-------------|
| Nausea            | 4.75 ± 2.86| 0.88 ± 0.45| 1.56 ± 0.63| 4.94 ± 2.22| 0.096       |
| Pain              | 19.75 ± 6.75| 4.13 ± 2.52^a| 7.63 ± 3.15| 2.50 ± 6.35^abc| 0.019*      |
| Belching          | 3.38 ± 1.92| 2.25 ± 1.45| 3.56 ± 1.91| 7.50 ± 2.99| 0.218       |
| Flatulence        | 14.75 ± 3.66| 8.50 ± 2.84| 17.38 ± 5.29| 13.81 ± 3.57| 0.705       |
| Constipation      | 3.75 ± 2.72| 4.00 ± 2.82| 0.56 ± 0.34| 4.19 ± 2.71| 0.369       |
| Diarrhoea         | 12.19 ± 5.70| 3.06 ± 1.83| 3.75 ± 3.75| 13.13 ± 5.72| 0.149       |
| Defecation        | 2.50 ± 1.88| 2.88 ± 2.23| 0.37 ± 0.32| 8.75 ± 5.21| 0.247       |
| Urge to defecate  | 12.19 ± 4.92| 5.13 ± 2.67| 7.69 ± 4.96| 17.06 ± 6.70| 0.155       |
| Bloating          | 7.88 ± 4.15| 1.25 ± 0.72| 8.94 ± 4.96| 12.69 ± 4.53^b| 0.236       |

Table 2 displays mean data for individual GI symptoms (arbitrary units) across both dietary interventions. * denotes significant interaction effect (P < 0.02). ^ denotes significant change within-group (P < 0.03). ^b denotes significant post-diet difference in comparison to LOW FODMAP (P < 0.02).

Fig. 3 Mean plasma I-FABP concentrations at rest in response to FODMAP conditions. The figure demonstrates the average I-FABP (pg·ml⁻¹) before and after the LOW FODMAP and HIGH FODMAP intervention. No significant differences reported (P > 0.05)
[22, 23, 42] and paracellular transport. Although transient, acute GI disruption may exacerbate nutrient malabsorption, as well as provoke delayed systemic immune responses. Increased residual \( \text{HIGH}_{\text{FODMAP}} \) GI content as a result of malabsorption [43], along with increased delivery of fluid to the colon and reduced GI motility could synergistically impact on perceived severity of symptoms, including acute or transient pain. This may limit the intensity of exercise training, particularly in symptomatic or hypersensitive individuals. The reduction in pain observed in this study, along with improved perception of flatulence and bloating symptoms within-group, indicates that lowering FODMAPs in the diet may support exercise training. Mechanistically, a reduction in fluid re-uptake across the GI epithelia, leading to less fluid and gas build-up pre- or during exercise in response to daily or more habitual \( \text{LOW}_{\text{FODMAP}} \) approaches may assist with sustained exercise tolerance.

In connection with perceived symptom changes, this study also investigated whether a short-term FODMAP approach impacted on basal GI damage via assessment of I-FABP. Whilst it was hypothesised that a \( \text{HIGH}_{\text{FODMAP}} \) diet may lead to an elevated residual level of I-FABP following the short-term intervention, no significant differences were observed within or between conditions. Therefore, even though a \( \text{HIGH}_{\text{FODMAP}} \) approach may have resulted in increased perception of symptom severity, disruption of the epithelial barrier in response to dietary modifications was not evident. Previous research has demonstrated that splanchnic hypoperfusion in response to acute, moderate exercise resulted in elevated I-FABP from 309 ± 46 pg·ml\(^{-1}\) to 615 ± 118 pg·ml\(^{-1}\) in healthy, male volunteers [22], which rapidly returned to baseline concentrations within minutes of recovery. I-FABP is a sensitive marker of small intestinal cell damage. However, rapid changes as observed in the previous study [22] indicate that GI damage is highly transient, and possibly only in response to exercise-based hypoperfusion, which may explain the lack of significant residual findings under resting conditions in the current study.

Assessment of I-FABP and/or GI permeability (e.g. urinary lactulose: rhamnose evaluation) in response to daily bouts of exercise in conjunction with a FODMAP approach may, however, provide mechanistic understanding of the potential benefits of a \( \text{LOW}_{\text{FODMAP}} \) diet.

A limitation to the current study observed when implementing a \( \text{LOW}_{\text{FODMAP}} \) diet in free-living conditions was that participants tended to consume fewer calories compared to both habitual and \( \text{HIGH}_{\text{FODMAP}} \) intakes, albeit non-significant. Taking into consideration methodological constraints in maintaining a weighed food diary, this observation was supported by a significant reduction in carbohydrate intake to achieve \( \text{LOW}_{\text{FODMAP}} \) adherence. This finding is consistent with a case study of a female athlete competing in a Multi-stage Ultramarathon [38], which reported that whilst following a \( \text{LOW}_{\text{FODMAP}} \) approach total daily energy intake did not meet estimated energy requirements. Upon further investigation suboptimal carbohydrate intake rather than protein and fat was observed.

Similar findings have been reported elsewhere [44], in which 29% of participants reported acute weight loss whilst on a \( \text{LOW}_{\text{FODMAP}} \) approach in free-living conditions. Many carbohydrate-rich foods typically consumed by active individuals (e.g. pasta, cereals, bread, energy drinks) are \( \text{HIGH}_{\text{FODMAP}} \) whereas alternative food sources (e.g. rice, corn) may be less palatable or more difficult to substitute. Indeed, in the previous study [44], participants cited that \( \text{LOW}_{\text{FODMAP}} \) approaches were either too complicated, expensive, or did not enjoy the overall taste as reasons for not sustaining the diet. The potentially restrictive or limiting nature of food choices on a \( \text{LOW}_{\text{FODMAP}} \) diet could therefore outweigh GI symptom benefits in the longer term due to weight loss, lethargy, fatigue, perceived effort, cost and/or enjoyment. Furthermore, a sustained energy/carbohydrate reduction in the longer term could also impact on training maintenance and recovery adaptations, and lead to unintended reduced nutrient availability.

However, the finding that acute dietary FODMAP manipulation positively impacted on overall GI symptom severity has pertinent implications for active individuals, particularly those more symptomatic or hypersensitive. Future research should consider whether there is a threshold of symptom severity in the context of exercise above which individuals may benefit from a \( \text{LOW}_{\text{FODMAP}} \) nutritional approach. There is also the need to establish the minimum intervention length required to alleviate GI symptoms in both recreational and trained athletes; as well as how long interventions can be sustained or indeed whether a FODMAP approach can impact on prolonged training periods. This is especially important considering the finding that carbohydrate intake was reduced on the \( \text{LOW}_{\text{FODMAP}} \) diet in free-living conditions, and the known importance of carbohydrates in fuelling regular exercise. Finally, whilst a \( \text{LOW}_{\text{FODMAP}} \) diet is known to significantly affect the gut microflora composition [45–47], the consequences of prolonged \( \text{LOW}_{\text{FODMAP}} \) intake on other types of physical activity, particularly those of a high-intensity or intermittent nature, has yet to be determined.

**Conclusion**

In conclusion, this study provides evidence that recreational athletes implementing a short-term \( \text{LOW}_{\text{FODMAP}} \) diet under free-living conditions may experience benefits in exercise-related GI symptoms and perceived
improvements in exercise intensity and frequency. However, caution is warranted to minimise unnecessary reductions in total caloric and/or carbohydrate intake that may impact on nutritional quality. Further studies are warranted to investigate the impact of a LOWFODMAP diet on sustained training strategies in healthy, recreationally active individuals and trained athletes.

Abbreviations
ANOVA: analysis of variance; ELSA: Enzyme-linked immunosorbent assay; FODMAP: Fermentable oligosaccharide, disaccharide, monosaccharide and polyol diet; GI: Gastrointestinal; IBS: Irritable bowel syndrome; IBS-SSS: Irritable bowel syndrome-symptom severity score; I-FABP: Intestinal fatty acid binding protein; ROS: Reactive oxygen species

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Availability of data and materials
The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions
All authors were involved in the study. JR and MW conceived and designed the study; MW was centrally involved with data collection; data were analysed by MW and JR with statistical support from LS; MW and JR constructed the manuscript. All authors reviewed the manuscript and approved the final version prior to submission.

Ethics approval and consent to participate
This study was conducted in accordance with the Declaration of Helsinki, and the protocol was approved by the Faculty of Science and Technology Ethics Committee, Anglia Ruskin University (PST/REF/15/667). Written informed consent was obtained from all individual participants included in the study.

Consent for publication
As part of the written informed consent procedure, all participants were duly made aware (as part of both the study briefing and information sheet) that the study results may be published. As such, consent for publication was included as part of this process.

Competing interests
The authors declare that they have no competing interests.

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