Original research

Repeated muscle glycogen supercompensation with four days’ recovery between exhaustive exercise

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**A R I C L E   I N F O**

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**A B S T R A C T**

Objectives: To determine if a 4 d period of high carbohydrate intake can supercompensate muscle glycogen and exercise work capacity on back-to-back occasions.

Design: Seven trained cyclists (6 male, VO\textsubscript{peak}: 57 ± 4 mL.kg\textsuperscript{-1} min\textsuperscript{-1}) completed a 9-d experimental period, consisting of three intermittent exhaustive cycling trials on days 1 (trial 1), 5 (trial 2) and 9 (trial 3). Following trial 1 cyclists were fed a high carbohydrate diet (10 g.kg\textsuperscript{-1} day\textsuperscript{-1}) for eight days to assess their capacity to repeatedly supercompensate muscle glycogen with 4 d recovery.

Methods: A resting muscle biopsy was obtained prior to each trial consisting of 2 min work intervals (90–60% peak power output) interspersed with 2 min recovery (40% peak power output) repeated until exhaustion. Each 72-h period between trial days included two days of low volume cycling and a rest day. Resting muscle glycogen and total work completed was determined for each trial day.

Results: Baseline muscle glycogen on day 1 (583.6 ± 111.0 mmol kg\textsuperscript{-1} dry mass) was supercompensated on day 5 (835.1 ± 112.8 mmol kg\textsuperscript{-1} dry mass; p = 0.04, d = 2.25) and again on day 9 (848.3 ± 111.4 mmol kg\textsuperscript{-1} dry mass; p = 0.01, d = 2.38). Total cycling work capacity increased from trial 1 to trial 2 (+8.7 ± 5.4 kJ kg\textsuperscript{-1}; p = 0.01; d = 1.41); a large effect was observed in trial 3 compared to trial 1 (+6.4 ± 6.8 kJ kg\textsuperscript{-1}; p = 0.10; d = 1.10).

Conclusions: A 4 d high carbohydrate feeding strategy is sufficient to repeatedly supercompensate muscle glycogen content following exhaustive exercise and results in enhanced work capacity.

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**Practical implications**

- It is possible to repeatedly supercompensate muscle glycogen with 4 d of high CHO feeding between consecutive bouts of exhaustive exercise.
- Improved high intensity exercise capacity (25–30%) was maintained during “back-to-back” exercise performances, demonstrating beneficial effects associated with effective repeated CHO loading.
- Practitioners and athletes should carefully consider the dietary and recovery strategy that enables the maintenance of exercise capacity when the competition schedule is congested with limited recovery between matches or events.

**1. Introduction**

The relationship between dietary carbohydrate (CHO) intake and skeletal muscle glycogen content has been recognised for over half a century. Pioneering work from the 1960’s demonstrated that glycogen depletion concomitant with a high CHO diet over subsequent days resulted in an enhanced glycogen storage.\textsuperscript{1} These data were the foundation for studies elucidating the relationship between dietary CHO intake and muscle glycogen concentrations,\textsuperscript{2} exercise capacity\textsuperscript{3} and performance.\textsuperscript{4} The use of stable isotopes in later years not only confirmed an increased reliance on carbohydrate oxidation with increasing exercise intensity, but also highlighted the crucial contribution of endogenous CHO to exercise greater than 85% of VO\textsubscript{2peak}.\textsuperscript{5,6}

Carbohydrate loading achieved through high dietary CHO intake has been shown to promote skeletal muscle glycogen supercompensation and enhance prolonged (>90 min) endurance exercise performance in laboratory\textsuperscript{7} and field\textsuperscript{8} settings. Current recommendations suggest a 36–48 h period of high CHO feeding at...
8–12 g·kg⁻¹·day⁻¹ is sufficient to supercompensate muscle glyco-
gen stores during a taper period. However, despite the potential
performance benefits of high pre-exercise CHO intake, the con-
gested competition schedules of professional athletes of many tea-
sports and cycling in various formats including one-day classics,
often limits the time between events and removes the oppor-
tunity for a traditional taper. In such a scenario, there is a paucity of
data available on the capacity to supercompensate muscle glycogen
twice in quick succession; i.e., repeatedly CHO loading for com-
petitions separated by 3–4 days.

To date, a single study has examined the efficacy of “back-to-
tack” carbohydrate loading to achieve repeated muscle glycogen
supercompensation. McInerney et al. determined the capacity of
ewell-trained male cyclists to supercompensate muscle glycogen
prior to consecutive exhaustive exercise trials separated by 48 h
recovery. They showed that with very high CHO intakes
(12 g·kg⁻¹·day⁻¹) throughout the experimental period, participants
were able to supercompensate muscle glycogen once, but not on
a second occasion when a 48 h recovery or “turnaround time” was
available. Interestingly, Bussau et al. have shown well-trained
cyclists can supercompensate muscle glycogen in just 24 h; thus,
while the mechanisms underpinning the inability to repeatedly
supercompensate muscle glycogen remain unclear, the interaction
between days of recovery following repeated exhaustive exercise,
the amount of CHO consumed, and the fate of ingested CHO appear
to dictate the physiological and performance outcomes if repeated
muscle glycogen supercompensation is desired.

Accordingly, the aim of the present study was to determine
if extending the CHO loading period to 4 d between exhaus-
tive exercise trials would enable “back-to-back” muscle glycogen
supercompensation in trained athletes. We hypothesised that a
4-day duration of higher total CHO intake between trials would
augment the capacity for glycogen storage enabling repeated mus-
cle glycogen supercompensation and enhanced performance.

2. Methods

Six male and one female cyclist completed the study (n = 7,
37 ± 6 y, VO₂peak: 56.8 ± 3.7 mL·kg⁻¹·min⁻¹, peak power output
(PPO): 323 ± 36 W, 76.1 ± 10.0 kg, 177.8 ± 6.2 cm). All participants
reported a consistent training volume (10.1 ± 3.0 h·w⁻¹) for at least
eight weeks prior to participation. The study was approved by Bond
University’s Human Research Ethics Committee and participants
provided written informed consent prior to the study.

Participants initially visited the laboratory for preliminary test-
ing and familiarisation, and provided a completed 3 d dietary
record. Height and body mass was recorded (WM204, Wedder-
burn, Australia) and mass at this time-point was used for dietary
prescription. Subsequently, a graded exercise test was conducted
to determine PPO for steady state (SS) cycling and trial workload
prescription. Testing was conducted on an Excalibur Sport cycle
ergometer (Lode, Groningen, Netherlands) and expiratory gasses
were continuously monitored on a metabolic cart calibrated to
manufacturer’s instructions (CosMed, Rome, Italy). VO₂peak was
determined as the highest rate of oxygen consumption recorded over a
30 s average. Participants then undertook a 60 min familiarisation
of the trial day protocol on the same Wattbike Pro cycling ergome-
ter that was to be used in each trial (Wattbike Ltd, Nottingham,
UK).

Seven (±2) days later participants commenced a 48 h exercise
and 24 h dietary standardisation. Two days prior to cycling trial
1, participants completed a standardised SS cycling bout between
58 and 63% PPO for 60–90 min. SS cycling bouts were also com-
pleted throughout the experimental period on days 2, 3, 6 and 7.
All SS cycling bouts during the study were undertaken on the same
habitual road training circuit matched within-participant, and a
pedal-based power meter system was fitted to each participant’s
bike for training prescription and load monitoring (Assioma Uno,
Favero Electronics, Arcade TV, Italy). One day prior to cycling trial
1, participants were provided with a diet containing 5 g·kg⁻¹·CHO.
A continuous glucose monitor (CGM; iPro2; Medtronic, Northridge,
CA, USA) was also inserted into the abdominal region, and a second
CGM was inserted on day 4 of the experimental period. Partici-
pants recorded blood glucose concentrations from capillary blood
(Abbott, IL USA) prior to each meal and sleep for CGM calibration.
Seventy-two-hour area under the curve (AUC) was determined for
the three days preceding trial 2 and 3, during CHO load 1 and 2,
respectively. Due to technical difficulties CGM data are n = 6. Fol-
lowing the 48 h standardisation period participants commenced the
9 d experimental period consisting of three cycling trials on days 1, 5 and 9 (Fig. 1).

On the morning of each cycling trial, participants reported to
the laboratory (0600) after an overnight fast and a resting skele-
tal muscle biopsy was obtained from the vastus lateralis under local anaesthetic (1% xylocaine) using a 5 mm Bergstrom needle
modified with manual suction. Body mass was measured and par-
ticipants completed a 10 min warm up at 40% PPO. Trials consisted of
2 min intervals at a 1:1 work-recovery ratio with heart rate
(HR) recorded continuously (Polar H10, Polar Electro, Finland).
Work intervals initially corresponded to a mechanical workload
of 90% PPO, and all recovery intervals were 40% PPO. Participants
completed as many repetitions as possible until they could no
longer maintain the prescribed workload (mean interval workload
less than 3 W from prescribed); mechanical workload was then
decreased by 10%. This process was repeated at workloads of 80,
70 and 60% PPO. When a workload of 60% PPO could no longer be
maintained the trial was terminated. Blood glucose was measured
immediately post-exercise, and athletes were subsequently fed a
meal containing 2 g·kg⁻¹·CHO within 15 min of exercise cessation.
Total work was calculated as the sum of all completed/attempted intervals inclusive of recovery bouts and expressed relative to body
mass recorded on that day (kg·kg⁻¹). Work done at each workload
was calculated as the sum of only fully completed intervals, exclud-
ing recovery work. All exercise testing bouts were completed under
standard laboratory conditions (22 ± 2 °C) on the same cycle ergome-
ter. At the completion of trial 1, participants consumed a high CHO
diet (10 g·kg⁻¹·day⁻¹) for the next eight days.

All meals and snacks were provided to participants and all foods
were weighed prior to packaging for estimation of consumed CHO
(Foodworks 7.0, Xyris, Australia). Participants were instructed to
consume only foods provided and complete only exercise associated with the study; both were verified by daily food checklists and training diaries, respectively. Participants were instructed to consume foods as three main meals as prescribed, with additional foods consumed ad libitum but replicated on a daily basis. Final adjustments were made to the nutrient analysis for any foods omitted by participants. Training diaries recorded session rating of perceived exertion (sRPE)\(^\text{15}\) for SS cycling bouts, and data files from participant's global positioning systems were received daily to quantify external load (kJ kg\(^{-1}\)) and verify compliance. Trials were separated by matched 72 h periods consisting of the same prescribed low intensity SS cycling bout (as in the standardisation period) for two days followed by one rest day; thus, both training and diet from days 1–4 (CHO load 1, prior to trial 2) were replicated in days 5–8 (Fig. 1).

Muscle glycogen concentration was determined according to the acid hydrolysis method\(^\text{16,17}\) with glucose concentration quantified using a commercially available kit (GLUC-HK, Randox Laboratories, Antrim, UK). Glycogen concentration (mmol kg\(^{-1}\) dry mass [DM]) was then calculated and the mean of replicates used for data analysis. The CV of glycogen replicates was 5.9 ± 6.0% (mean ± SD).

Data were analysed using one-way repeated measures analysis of variance and Tukey's multiple comparisons test was used for trial-to-trial comparisons. Paired t-tests were used to compare 72 h AUC values from interstitial glucose traces. Normality was confirmed by Shapiro–Wilks test and alpha was 0.05. Cohen's d effect sizes were used to determine the magnitude of differences with threshold values for small, moderate and large effects 0.2, 0.5 and 0.8, respectively. All statistical analyses were conducted in GraphPad Prism 7.03 (GraphPad software, CA USA) and data are mean ± standard deviation.

3. Results

Participants' reported energy intake was 151.0 ± 24.0 kJ kg\(^{-1}\), containing 3.7 ± 0.8 g kg\(^{-1}\) CHO. In the 24 h prior to cycling trial 1 participants consumed 137.4 ± 6.4 kJ kg\(^{-1}\), containing 5.2 ± 0.1 g kg\(^{-1}\) CHO in the provided standardised diet. Over the following eight days, participants consumed a daily mean energy intake of 248.2 ± 9.7 kJ kg\(^{-1}\) with mean macronutrient intakes of 10.6 ± 0.4 g kg\(^{-1}\) CHO (0.3 ± 0.1 g kg\(^{-1}\) fibre), 1.6 ± 0.1 g kg\(^{-1}\) protein and 1.0 ± 0.1 g kg\(^{-1}\) fat. Small daily variations in consumed CHO and energy intake were apparent with CHO intake higher on day 7 (10.8 ± 0.4 g kg\(^{-1}\)) compared to days 1, 2 and 4 (range of mean data: 10.4–10.5 g kg\(^{-1}\), p < 0.05). Total energy was also lower on day 1 (241.1 ± 9.4 kJ kg\(^{-1}\)) compared to days 2, 4, 6, 7 and 8 (249.3–252.1 kJ kg\(^{-1}\), p < 0.05), and on day 5 (243.2 ± 10.7 kJ kg\(^{-1}\)) compared with days 6 and 8 (252.1–252.2 kJ kg\(^{-1}\); p < 0.05).

There was a main effect of trial for muscle glycogen concentration (Fig. 2A). Baseline muscle glycogen at trial 1 (583.6 ± 111.0 mmol kg\(^{-1}\) DM) was increased prior to trial 2 (835.1 ± 112.8 mmol kg\(^{-1}\) DM, p = 0.04, d = 2.25) and trial 3 (848.3 ± 111.4 mmol kg\(^{-1}\) DM, p = 0.01, d = 2.38). There was also a main effect of trial for total work capacity (Fig. 2B) and trial/cycling duration. Work capacity and cycling duration in trial 1 (25.3 ± 6.1 kJ kg\(^{-1}\); 169.1 ± 43.8 min) was increased in trial 2 (34.0 ± 6.2 kJ kg\(^{-1}\); p = 0.01, d = 1.41; 226.6 ± 31.7 min, p < 0.01, d = 1.50); a large but non-significant effect was evident in trial 3 (31.7 ± 5.5 kJ kg\(^{-1}\); p = 0.10, d = 1.10; 208.0 ± 22.4 min, p = 0.12, d = 1.12). There was a modest decrease in cycling work capacity between trials 2 and 3 only evident as a small effect size (−2.3 ± 3.6 kJ kg\(^{-1}\); p = 0.28, d = 0.39; −18.6 ± 24.6 min, p = 0.19, d = 0.68). Body mass increased 1.0 ± 0.7% from trial 1 to trial 2 (75.6 ± 10.1 vs. 76.3 ± 10.3 kg, p = 0.02, d = 0.07), but was not different in trial 3 compared to trial 1 (0.6 ± 0.9%; 76.0 ± 10.0 kg, p = 0.25, d = 0.04).

There were no differences across trials in the amount of work completed at any intensity. However, compared to trial 1 (7.1 ± 2.3 kJ kg\(^{-1}\)) there was a small and moderate increase in high intensity work done at 90% PPO in trial 2 (8.9 ± 4.7 kJ kg\(^{-1}\); p = 0.30, d = 0.49) and 3 (9.8 ± 5.8 kJ kg\(^{-1}\); p = 0.25, d = 0.61), respectively. There was a main effect of trial for peak HR during high intensity (90% PPO) workloads. Peak HR was lower in trial 3 (170.4 ± 7.1) compared to both trial 2 (174.1 ± 7.5, p = 0.03, d = 0.51) and trial 1 (177.3 ± 7.3, p = 0.006, d = 0.96). Post-exercise blood glucose concentration was not different between trials (4.1 ± 0.6 mM).

Intestinal glucose AUC was different during matched 72 h periods spanning CHO load 1 and 2. The AUC during CHO load 1 (days 2–4; 390.2 ± 28.9 mmol L\(^{-1}\)) was greater than during CHO load 2 with a small effect size (days 6–8; 376.8 ± 31.1 mmol L\(^{-1}\); p = 0.03, d = 0.45; Fig. 3). There were no differences in external load (110.0 ± 3.4 kJ kg\(^{-1}\)) or sRPE across all SS cycling bouts (297.8 ± 111.3 AU). The mean duration of SS cycling bouts was 75.5 ± 22.0 min and the mean workload was 58.3 ± 3.4% of PPO (2.5 ± 0.2 W kg\(^{-1}\)).

4. Discussion

The main findings of this study were that: (1) repeated muscle glycogen supercompensation was achieved with consecutive 4 d periods of high CHO feeding; (2) 72 h interstitial glucose AUC was lower throughout the second CHO loading period compared to the
first CHO load, despite matched dietary intakes; and (3) successive glycogen supercompensation was accompanied by a large increase in exercise work capacity on each occasion.

Our study shows for the first time that back-to-back muscle glycogen supercompensation is possible following exhaustive exercise with 4 d of high carbohydrate feeding. The data from the present study expand on the previous work from McInerney et al.\textsuperscript{12} that show consecutive 2 d periods of CHO loading between exhaustive exercise substantially increased muscle glycogen concentration after the first 2 d CHO loading period, but failed to promote glycogen supercompensation after the second CHO loading period. Despite the difference in recovery days between studies, the initial glycogen supercompensation was in close agreement, resulting in an increased muscle glycogen content of 250–280 mmol kg\textsuperscript{-1} DM above normal resting levels. However, the contrasting effect on glycogen resynthesis between studies when the high CHO feeding strategy was immediately repeated, indicates interactions between temporal, dietary and physiological factors may affect the capacity for consecutive muscle glycogen supercompensation in relatively short timeframes.

It has been proposed that failure of muscle glycogen stores to repeatedly reach supercompensated levels may be a consequence of the cumulative effect of exhaustive exercise bouts in close succession\textsuperscript{12}. For example, acute mechanisms of glycogen resynthesis are purported to include elevated AMPK and glycogen synthase activity.\textsuperscript{18} However, glycogen synthase activity appears to be similar with repeated glycogen depletion and carbohydrate loading,\textsuperscript{19} and therefore seems unlikely to attenuate glycogen supercompensation under these conditions. The possibility exists that excessive muscle fatigue and/or a priority for muscle repair and remodelling after repeated and prolonged strenuous exercise impairs, at least in part, activation of the cellular mechanisms for glycogen resynthesis, or glucose is utilised for cellular processes rather than glycogen storage\textsuperscript{19}; such a hypothesis has been proposed following exercise inducing mechanical muscle damage,\textsuperscript{19} but is also plausible following high intensity cycling, given prolonged/intense aerobic exercise is also associated with high rates of protein turnover.\textsuperscript{20} Consequently, the 4 d between exercise to exhaustion in the present study that included two recovery days and a rest day, together with a total CHO intake of 40 g kg\textsuperscript{-1}, may be necessary to permit repeated muscle glycogen supercompensation. Whether an impairment in glycogen resynthesis is unique to exhaustive exercise bouts with limited recovery (<48 h) and whether severe glycogen depletion is a prerequisite for repeated muscle glycogen supercompensation are important questions for future research. A limitation of the present study was the omission of post-exercise muscle biopsies following trials 1 and 2; consequently, the post-exercise muscle glycogen concentrations are not reported. However, based on data from McInerney and colleagues,\textsuperscript{12} it is clear that this exercise protocol significantly reduces muscle glycogen, and post-exercise muscle glycogen concentrations in our study were likely 100 mmol kg\textsuperscript{-1} DM. Nonetheless, our data provide support for the practical utility of CHO loading to elevate muscle glycogen concentrations for enhanced performance capacity in competition,\textsuperscript{8} and also provides new information to show repeated glycogen supercompensation is achievable in a 4 d timeframe, which has important implications for dietary and recovery strategies during congested competition schedules.

Interestingly, despite muscle glycogen supercompensation in response to the consecutive CHO loading trials, there was a small effect for a lower glucose AUC during the second CHO load. There are several potential mechanisms to explain such a finding. Firstly, a reduced CHO absorption at the gastrointestinal tract may be evident, due to an interaction between repeated prolonged exercise bouts that may predispose to intestinal epithelial damage,\textsuperscript{21} and prolonged exposure to a very high CHO/energy diet. Indeed, the primary glucose transporter in enterocytes (SGLT1) has the capacity for dynamic change and is highly responsive to altered carbohydrate intake.\textsuperscript{22,23} Accordingly, the initial days of high CHO feeding following trial 1 may have promoted an immediate adaptive response that increased activity of transporters, which abated in response to the arduous exercise and dietary regimens. Second, it is possible the lower glucose AUC may be due to enhanced storage of glycogen in Type II fibres stimulated by the high intensity, exhaustive exercise during trial 1 that may not have been detectable from the glycogen assay. Finally, it’s most likely an increased clearance of circulatory glucose is due to greater rates of CHO oxidation with prolonged exposure to high carbohydrate availability. Goedecke et al.\textsuperscript{24} have shown muscle glycogen content is strongly correlated with resting respiratory exchange ratio, and together with Type I muscle fibre content (among other factors) explains 60% of the variability in resting respiratory exchange ratio. This phenomenon and the increase in CHO oxidation routinely reported at rest\textsuperscript{25} and during exercise\textsuperscript{26} with high dietary CHO intakes likely accounts for the lower whole body glycaemia during the 72 h period between trials 2 and 3.

The carbohydrate loading protocol in the present study was associated with an increase in total work capacity, and a small-moderate effect for greater work performed at 90% PPO in trials 2 and 3, respectively. Accordingly, our data support the well characterised and beneficial effect of commencing exercise with high muscle glycogen concentration on endurance performance.\textsuperscript{7} Indeed, a recent meta-analysis\textsuperscript{27} presents models of muscle glycogen utilisation during exercise of differing intensities and starting glycogen concentrations. Our data support the findings of Areta and Hopkins\textsuperscript{28} showing higher starting glycogen concentration extends time to fatigue during prolonged bouts of endurance exercise.

While there was similar high intensity work performed in trial 2 and 3, a marginal reduction in total work capacity was apparent in trial 3 despite comparable starting muscle glycogen. It may be that the repeated priming of muscle with high starting glycogen concentration prior to prolonged exercise predisposed cellular metabolism to greater rates of glycogen utilisation during trial 3,\textsuperscript{28} resulting in a minor reduction in exercise capacity. In addition, peak heart rate in trial 3 was lower than trials 2 and 1, which may be indicative of cumulative fatigue/overreaching in participants undertaking a series of exercise bouts to exhaustion.\textsuperscript{29} Regardless, in trial 3 participants completed more high intensity work compared to trial 1, and displayed a similar total work capacity to trial 2, demonstrating a beneficial effect of CHO loading to a fatigued athlete.
5. Conclusion

Our study provides novel data on the supercompensation of muscle glycogen concentration with a high CHO intake between repeated bouts of exhaustive exercise. We show that repeated muscle glycogen supercompensation is possible with 4 d of high CHO feeding between exhaustive exercise bouts, and that the CHO loading strategy we employed was associated with an increase in the amount of high intensity work performed and superior endurance capacity.

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