

Sex differences in the effects of 12 weeks sprint interval training on body fat mass and the rates of fatty acid oxidation and VO₂max during exercise

Liam Bagley,¹ Mark Slevin,¹ Steven Bradburn,¹ Donghui Liu,¹ Chris Murgatroyd,¹ George Morrissey,² Michael Carroll,¹ Mathew Piasecki,¹ William S Gilmore,¹ Jamie S McPhee¹

To cite: Bagley L, Slevin M, Bradburn S, *et al*. Sex differences in the effects of 12 weeks sprint interval training on body fat mass and the rates of fatty acid oxidation and VO₂max during exercise. *BMJ Open Sport Exerc Med* 2016;**2**:e000056. doi:10.1136/bmjsem-2015-000056

► Prepublication history for this paper is available online. To view these files please visit the journal online (<http://dx.doi.org/10.1136/bmjsem-2015-000056>).

Accepted 11 February 2016



CrossMark

¹School of Healthcare Science, Manchester Metropolitan University, Manchester, UK

²Cambridge University School of Clinical Medicine, Addenbrooke's Hospital, Cambridge, UK

Correspondence to

Liam Bagley;
liam.bagley@mmu.ac.uk

ABSTRACT

Background: The purpose of this study was to examine whether very short duration, very high intensity sprint interval training (SIT) leads to loss of body fat mass in association with improvements to VO₂max and fatty acid oxidation, and to assess the extent of sex dimorphism in these physiological responses.

Methods: A total of 24 men and 17 women (mean (SEM) age: 39 (±2) years; body mass index 24.6 (0.6)) completed measurements of the maximal rate of oxygen uptake (VO₂max) and fatty acid oxidation (FATmax). Body fat and lean mass were measured by dual emission x-ray absorptiometry, and fasting blood lipid, glucose and insulin profiles were assessed before and after training. SIT consisted of 4×20 s sprints on a cycle ergometer at approximately 175% VO₂max, three times per week for 12 weeks.

Results: Fat mass decreased by 1.0 kg, although men lost statistically significantly more fat than women both when expressed in Kg and as % body fat. VO₂max increased by around 9%, but women improved VO₂max significantly more than men. FATmax improved by around 13%, but fasting plasma glucose, insulin, total triglyceride, total cholesterol and high-density lipoprotein (HDL) did not change after training, while low-density lipoprotein decreased by 8% (p=0.028) and the HDL:Total Cholesterol ratio improved by 6%. There were no sex differences in these metabolic responses to training.

Conclusions: These results show lower body fat %, and higher rates of fatty acid oxidation and VO₂max after 12 weeks of training for just 4 min per week. Notably, women improved VO₂max more than men, while men lost more fat than women.

INTRODUCTION

Aerobic-type training of between 225 and 420 min/week is recommended to those who wish to lose fat mass by increasing physical

Summary

- In previous research, sprint interval training (SIT) has been shown to increase concentrations of enzymes of fatty acid metabolism in skeletal muscles of young adults.
- In the present study, adults from the general population performed SIT of 4 min per week for 12 weeks.
- The men and women improved maximal rate of oxygen uptake (VO₂max) by an average of 9%, rates of fatty acid oxidation during exercise by 13% and decreased body fat by 1 kg.
- Men and women responded differently to training, with women increasing VO₂max more than men, and men losing more body fat than women.

activity levels,¹ but very high intensity, short duration training might also be effective.^{2–3} ‘Sprint interval training’ (SIT) sessions require participants to perform repeated short duration (often approximately 30–60 s) bouts at very high or maximal power output, each separated by a recovery period of very low intensity exercise.^{4–5} This exercise differs from conventional aerobic training that typically involves prolonged and continuous exercise, which during cycling is equivalent to around 25% of the peak leg power output⁶ that is met principally by the aerobic energy systems.^{7–8}

Despite the intensity, duration and energetic differences between short duration, high-intensity activities and endurance activities lasting around 60 min per session, they promote similar physiological adaptations.⁹ This includes improvements to the maximal rate of oxygen uptake (VO₂max)^{9–10}

occurring in association with improvements to skeletal muscle capillarisation,^{11 12} enzymes of fat metabolism^{2 5 13–18} and improved insulin sensitivity,^{4 5 19 20} which contribute to better health-status and physical performance.

There are also reports that very high intensity, short duration training can lead to loss of total body fat mass.^{2 21–23} These studies used weekly durations of around 9 min^{2 21 22} and 30 min²³ without controlling food intake. Whether even shorter duration exercise will be effective at reducing fat mass in men and women remains unknown.

Compared with men, women have lower relative muscle mass and higher relative fat mass,²⁴ and slower skeletal muscle contractile properties;^{25 26} they oxidise fatty acids at higher relative exercise intensity²⁷ and have less reliance on muscle glycogen stores following bouts of repeated sprinting.²⁸ Therefore, men and women might not necessarily show similar adaptations to fatty acid metabolism and body composition after SIT, but such sex differences remain unknown because the overwhelming majority of research into SIT has included only young men (usually university students) who trained for between 2 and 6 weeks.

The aim of this study was to examine the hypothesis that very short duration, very high-intensity sprinting exercise (on cycle ergometers) could lead to loss of body fat mass as well as improvements to rates of fat oxidation and VO_2max after 12 weeks in men and women recruited from the general population. In the light of the sex differences in body composition and muscle metabolism, we also hypothesised that men and women would differ in their physiological responses to SIT.

DESIGN

Ethical approval and participant recruitment

Volunteers provided written informed consent prior to participation. Those with a history of cardiovascular, neuromuscular or metabolic disease were excluded as well as people who had suffered a leg fracture within the

past 2 years. A total of 41 participants completed the 12-week SIT intervention and post-training measurements. Participant characteristics are shown in table 1.

Measurements

Participants attended the research laboratory in the morning, following a 12 h overnight fast, and having avoided strenuous activity and alcohol in the 24 h prior to testing. After providing informed consent and resting for at least 15 min, a 10 mL blood sample was collected into an EDTA collection tube from a vein in the forearm and then a second sample during the same laboratory visit was collected 20 min after completion of the physiological testing. Blood was immediately centrifuged at 4°C, with the plasma separated and stored at –80°C until analysis.

Body fatness was assessed from a total-body dual-energy X-ray absorptiometry (DEXA) scan using a Lunar Prodigy Advance and off-line analysis (GE Medical; EnCore V.10.50.086). Total body fat, trunk fat and limb fat mass were analysed as previously reported.^{30 31}

Maximal rates of oxygen uptake and fatty acid oxidation were measured during cycle ergometry (Jaeger Ergocycle, Germany) with VO_2 , VCO_2 and heart rate (HR) measurements (Cortex Biophysik, Germany). Workload began at 50 W and a cadence of 70 rpm was maintained throughout. Workload was increased in increments of 50 W for men and 30 W for women every 3 min until the respiratory exchange ratio (calculated as VCO_2/VO_2) was higher than 1.0 for at least 1 min. From this point onwards, increments of 20 W were given every minute until volitional exhaustion. Off-line analysis was used to estimate rates of fatty acid oxidation, which were calculated from VO_2 and VCO_2 values collected during steady-state stages of the VO_2max test using the formula described by Frayn,³² assuming urinary nitrogen was negligible:

$$\text{Fatty acid oxidation (g/min)} = (1.67 \times \text{VO}_2) - (1.67 \times \text{VCO}_2)$$

Table 1 Body composition in men and women before and after 12 weeks of SIT

	Men (pre)	Men (post)	Women (pre)	Women (post)	Training effect	Sex effect	Sexxtraining
Age (years)	38 (2.7)		41 (3.2)				
Height (cm)	177 (1.6)		165 (1.2)				
Total body mass (kg)	80.0 (2.2)	79.2 (2.1)	62.5 (2.6)	62.1 (2.5)	0.070	<0.001	0.542
Body mass index (kg/m ²)	26.3 (0.8)	26.0 (0.7)	22.2 (0.7)	22.1 (0.6)	0.081	0.001	0.554
Total body fat (kg)	17.9 (1.3)	16.6 (1.3)	18.6 (1.4)	18.0 (1.2)	<0.001	0.556	0.052
Total Body Fat (%)	22.7 (1.7)	21.2 (1.4)	31.2 (1.7)	31.3 (1.6)	<0.001	<0.001	0.015
Trunk fat (kg)	9.7 (0.8)	9.0 (0.9)	8.1 (0.8)	8.0 (0.7)	0.014	0.313	0.046
Leg fat (kg)	5.7 (0.4)	5.4 (0.4)	7.8 (0.6)	7.0 (0.5)	<0.001	0.001	0.208
Total body lean mass (kg)	60.4 (1.6)	61.1 (1.6)	39.1 (0.9)	39.2 (1.0)	0.016	<0.001	0.162
Leg lean mass (kg)	21.6 (0.7)	21.0 (0.7)	13.6 (0.3)	13.0 (0.4)	0.185	<0.001	0.821

The bold font highlights statistically significant values ($p < 0.05$).

Data are shown as mean (SEM).

SIT, sprint interval training.

The maximal rate of fatty acid oxidation (FATmax) was estimated as the highest value occurring along the line of best fit of estimated rates of fatty acid oxidation (plotted from 50 W through to the point at which respiratory exchange ratio was higher than 1.0), adapted from Venables *et al.*²⁷

Sprint interval training

Participants were asked to maintain their usual dietary and exercise habits throughout the intervention. SIT was completed on cycle ergometers (Cateye, Japan). The training consisted of a 2 min warm-up at a self-selected moderate intensity. This was followed by four bouts of 20 s 'maximal effort' sprints at a workload that was set at 175% of the workload attained in the VO₂max test. Each of these intervals was separated by 2 min of very low intensity cycling (a workload of approximately 20% of that attained at VO₂max). Thus, each training session lasted less than 10 min and only 80 s was completed at an intensity that would be expected to improve physical fitness.

The first training session for each participant was fully supervised in the research laboratory, and participants also received clear instructions on the use of the cycle ergometers and the training regimen. Participants trained three times per week for 12 weeks using the ergonomic cycles that we provided. The training workload was increased by 5% every 2 weeks. Gym staff were fully informed of the research and training protocols, they logged the training session and were available to offer advice to research participants if needed during training sessions. Participants maintained a training-log to record workloads during training sessions.

Fasting blood lipoprotein profile, insulin and glucose

Four fasting plasma samples were collected as described above: (1) at baseline in the fasted and rested state; (2) at baseline in the fasted state but 20 min after completion of the physiological testing session; (3) after the 12-week exercise intervention in the fasted and rested state; (4) after the 12-week exercise intervention in the fasted state but 20 min after completion of the physiological testing session.

Insulin sensitivity was estimated using the Homeostatic Model of Assessment (HOMA) as described by Matthews *et al.*³³ which was calculated as:

$$\text{Fasting Plasma (Glucose [nmol/L])} \\ \times \text{Insulin } [\mu\text{mol/L}] \div 22.5$$

Biochemical markers were determined from fasting plasma samples using the RX Daytona auto analyser (Randox Laboratories, UK). High-density lipoprotein (HDL), low-density lipoprotein (LDL), total cholesterol (CHOD-PAP), triglycerides (GPO-PAP) and glucose (GOD-PAP) concentrations were determined in duplicate, and an average value calculated.

Statistical analysis

Data were analysed using SPSS (V.20 IBM). Repeated measures of ANOVA were used to assess adaptations to training and sex differences, using Greenhouse-Geisser correction when sphericity was violated. Pearson's Product Moment or Spearman's Correlations, as appropriate, were used to examine relationships between variables. Statistical significance was accepted at $p < 0.05$. Data are presented as mean \pm SEM, unless otherwise stated.

RESULTS

Body composition

Changes in body composition for men and women are shown in [table 1](#). Body fatness (%) decreased after training by 1.2% ($p < 0.001$), with men losing more total body (%) and trunk fat (kg) than women ([table 1](#)). Total body lean mass increased by 1.2% in men and 0.03% in women after training, but there was no statistically significant sex \times training interaction ($p = 0.162$), indicating that men and women showed similar changes to total lean mass after training ([table 1](#)).

Maximal oxygen uptake and fatty acid oxidation

[Table 2](#) shows data for VO₂max and FATmax in men and women. In the untrained state, men were able to utilise an absolute higher amount of fatty acids (in g/min) compared with women ($p = 0.027$), and FATmax (g/min) occurred at a higher absolute workload (W) ($p < 0.001$). Lean mass (kg) was correlated with FATmax (g/min) at baseline ([figure 1](#)) ($r = 0.366$; $p = 0.019$), however, the training-related change in total body lean mass (kg) was not statistically significantly correlated to training-related change in FATmax (g/min) ($r = 0.100$, $p = 0.536$) or when normalised to body mass (mg/kg/min) ($r = 0.065$, $p = 0.688$).

When normalised to total body mass and to lean mass (kg), FATmax did not differ between men and women ($p = 0.985$ and 0.287 , respectively). FATmax (g/min) occurred at a lower %VO₂max in men than in women ($p = 0.023$).

After 12 weeks of SIT, men and women increased VO₂max (mL/kg/min), with the percentage gains being 18.7% in women and 6.0% in men (sex \times training interaction: $p = 0.009$). Men and women showed similar gains in FATmax after training ([table 2](#)). The sex \times training interaction revealed that FATmax occurred at a statistically significantly lower %VO₂max in women after training but slightly higher %VO₂max in men after training ([table 2](#)).

Fasting plasma glucose, insulin and lipids

There were no statistically significant training effects for fasting plasma glucose ($p = 0.496$), insulin ($p = 0.708$), HOMA ($p = 0.426$), total triglyceride ($p = 0.702$), total cholesterol ($p = 0.129$) and HDL ($p = 0.332$). Training decreased LDL by 8% ($p = 0.028$) and the HDL:Total

Table 2 Maximal oxygen uptake and rates of fat oxidation measured during exercise in men and women before and after 12 weeks of SIT

	Men (pre)	Men (post)	Women (pre)	Women (post)	Training effect	Sex effect	Sex×training
VO ₂ max (L/min)	3.49 (0.13)	3.68 (0.15)	1.99 (0.11)	2.37 (0.12)	0.001	<0.001	0.049
VO ₂ max (mL/kg/min)	42.91 (1.84)	45.49 (1.48)	33.56 (2.29)	39.84 (2.44)	<0.001	0.005	0.009
HR _{max} (bpm)	177 (3)	178 (3)	175 (4)	174 (4)	0.645	0.977	0.268
RER _{max}	1.21 (0.01)	1.18 (0.02)	1.20 (0.02)	1.21 (0.02)	0.430	0.763	0.214
FATmax (g/min)	0.41 (0.03)	0.49 (0.04)	0.31 (0.03)	0.34 (0.02)	0.032	0.004	0.465
FATmax (mg/kg/min)	5.12 (0.39)	6.04 (0.40)	5.14 (0.52)	5.77 (0.41)	0.033	0.815	0.690
FATmax (mg/kg lean mass/min)	6.90 (0.50)	8.10 (0.62)	7.90 (0.76)	8.87 (0.65)	0.041	0.272	0.824
Workload at FATmax (W)	111.6 (9.81)	112.80 (7.56)	62.94 (5.43)	62.73 (6.76)	0.961	<0.001	0.834
FATmax (as %VO ₂ max)	51.43 (1.88)	51.83 (1.65)	60.26 (3.61)	53.25 (2.18)	0.229	0.066	0.047
Heart Rate at FATmax (bpm)	123 (5)	115 (3)	120 (4)	112 (4)	0.038	0.601	0.465

The bold font highlights statistically significant values ($p < 0.05$).

Data are shown as mean (SEM). All measurements were recorded during an incremental cycling test.

FATmax, maximal rate of fat oxidation; HRmax, maximal heart rate; RERmax, maximal respiratory exchange ratio; SIT, sprint interval training; VO₂max, maximal rate of oxygen uptake.

Cholesterol ratio improved by 6% ($p = 0.005$). There were no statistically significant sex×training interaction effects (all $p > 0.05$; table 3).

The responses to the acute mixed-exercise session were analysed by comparing the rested samples with those taken 20 min after exercise both at baseline and after the 12-week SIT. Total triglycerides, total cholesterol and LDL did not change in concentration between time points 1 and 2, or 3 and 4 (all $p > 0.05$). HDL increased by 4% from time point 1 to 2 ($p = 0.001$), but did not differ between 3 and 4 ($p = 0.318$). Total cholesterol:HDL ratio decreased by 2% from time point 1 to 2 ($p = 0.002$) and by 1% between time points 3 and 4 ($p = 0.013$). There were no statistically significant sex×time point interactions, indicating similar responses for men and women.

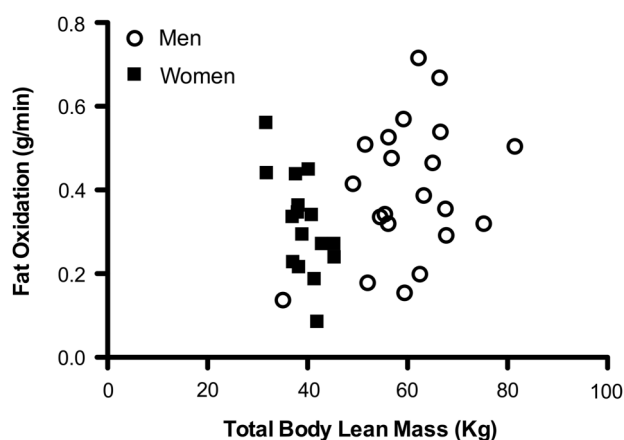


Figure 1 The relationship between total body lean mass and rates of fatty acid oxidation. In the untrained men and women, total body lean mass (kg) was correlated with rates of fatty acid oxidation (FATmax (g/min)) measured during exercise ($r = 0.366$; $p = 0.019$).

DISCUSSION

The present study showed that the SIT programme of just 4 min/week for 12 weeks decreased total body fat mass in men but not in women, while women improved VO₂max more than men. Improvements to the rates of fatty acid oxidation during submaximal exercise after the 12-week training programme were similar in men and women. These results highlight statistically significant sex-differences in physiological responses to very high-intensity training in the participants of this study who were recruited from the general population.

Body fatness

Body fat levels were within normal ranges for participants' ages at baseline³⁴ and lower body fat than their BMI related cut-offs.³⁵ Just 80 s of very intense sprint exercise per session, equal to 48 min exercise over 12 weeks, resulted in statistically significant reductions to body fat mass. As far as we are aware, the training duration of 4 min/week is the shortest reported to effectively cause fat loss without additionally restricting food intake. These results build on previous high-intensity training studies that also reported fat loss,^{2 21–23} although they utilised lower intensity and longer duration intervals than our training programme did.

The fat loss was principally due to changes seen in men, since women did not change their total body composition despite losing approximately 800 g fat from their legs. Our study is the first to show sex-differences in the changes to body composition after SIT and are in line with previous work indicating that men generally lose more fat than women after endurance training.³⁶ The reasons for the sex differences in training responses of body fatness and metabolism are not clear.³⁷ There is evidence that sex-differences in body fatness are linked to physiological actions of sex hormones,³⁸ but when

Table 3 Glucose and insulin concentrations in serum of men and women before and after 12 weeks of SIT

	Men (Pre)	Men (Post)	Women (Pre)	Women (Post)	Training effect	Sex effect	Sexxtraining
Glucose (mmol/L)	5.10 (0.16)	5.06 (0.05)	4.99 (0.11)	4.99 (0.05)	0.496	0.634	0.663
Insulin (μ U/mL)	3.50 (0.06)	3.48 (0.05)	3.35 (0.08)	3.44 (0.08)	0.708	0.995	0.241
HOMA (%S)	76 (2)	77 (1)	71 (2)	75 (2)	0.426	0.880	0.897
Triglycerides (mmol/L)	1.14 (0.27)	0.83 (0.07)	0.85 (0.06)	0.84 (0.08)	0.702	0.899	0.661
Total Cholesterol (mmol/L)	4.97 (0.24)	5.18 (0.20)	4.68 (0.20)	5.09 (0.19)	0.129	0.307	0.456
High-density Lipoprotein (mmol/L)	1.39 (0.06)	1.81 (0.14)	1.43 (0.07)	1.83 (0.14)	0.332	0.008	0.811
Low-density Lipoprotein (mmol/L)	2.75 (0.18)	2.55 (0.20)	2.51 (0.13)	2.42 (0.21)	0.042	0.562	0.523
Total Cholesterol:HDL ratio	3.60 (0.16)	3.01 (0.21)	3.29 (0.09)	2.91 (0.21)	0.010	0.039	0.161

The bold font highlights statistically significant values ($p < 0.05$).

Data are shown as mean (SEM).

HOMA, homeostatic model of assessment; SIT, sprint interval training.

energy balance was more closely regulated throughout an endurance training programme, overweight and obese men and women had similar fat loss.³⁹ The sex differences in fat loss might also be related to the sex differences in relative muscle mass,²⁴ skeletal muscle contractile^{25 26} and metabolic characteristics.^{27 28}

SIT sessions have very low energy consumption, of around 200 kJ/week, compared with endurance exercise of around 2000 kJ/week.¹³ The direct energy expenditure therefore cannot explain the fat loss occurring after 12 weeks of SIT. Other contributing factors might include an increase in post exercise energy expenditure or overall shift towards greater fatty acid oxidation during habitual activities throughout the day, as occurs after endurance training⁴⁰: the results showing increased fatty acid oxidation during low and moderate intensity activity are consistent with this. It is also possible that metabolic rate and lipid oxidation remain elevated after each training session, and there are reports that such effects are pronounced in men and negligible in women,^{40 41} which might help to explain our observed loss of fat in men but not in women. Women may also have a lower energy expenditure during exercise, due to the lower absolute workloads, despite working at similar relative exercise intensities, allowing men to reduce fat mass more than women after training.⁴² Treuth *et al*⁴³ found that resting metabolic rate remained around 15% higher in participants performing an acute session of SIT compared to those who completed endurance training. Higher intensity exercise is also associated with prolonged suppression of appetite and hunger.⁴⁴ However, others have reported increased energy intake in participants completing high-intensity training compared to non-exercising controls and those performing lower intensity exercise.⁴⁵ The combination of a shift towards greater fatty acid oxidation and elevated resting energy expenditure could lead to a negative calorie balance and thus loss of body fat.

Maximal rate of oxygen uptake:

Our results showed an overall 9% increase in VO_2 max after 12 weeks of SIT. This increase is similar to previous

reports from younger participants after SIT⁹ and of similar magnitude to VO_2 max gains after conventional endurance training programmes in which training sessions lasted around 1 h.^{13 18} Men and women both improved VO_2 max, but the gains in women were greater than those in men. It is not clear why disparity between sexes would occur in VO_2 max SIT responses. Men have been shown to have higher gains in VO_2 max following conventional endurance exercise,⁴⁶ but results from SIT studies are mixed. Scalzo *et al*⁴⁷ showed young women had similar gains in VO_2 max to young men, but other studies reported that men did not increase VO_2 max⁴⁸ while women showed large increases after SIT.⁴⁹

VO_2 max is an indicator of overall cardiopulmonary fitness and is dependent on the transportation of oxygen through the respiratory, cardiovascular and muscle systems to supply oxygen to the mitochondria for oxidative metabolism. A higher relative amount of lean mass in men compared to women, coupled with a higher relative body fat mass in women compared to men, may go some way in explaining the differences between men and women in maximal oxygen consumption.⁵⁰ However, the supply of oxygen to the working skeletal muscles is thought to be a limiting factor in VO_2 max,⁵¹ so the higher VO_2 max response in women might point to higher adaptations of oxygen supply than those in men following SIT, but more focused studies examining cardiac output, blood volume, haematocrit and blood flow distribution are needed to clarify this finding. Conversely, after regular endurance training, men had higher gains in VO_2 max compared with women.⁴⁶ It is possible that the training volume (higher in endurance) and training intensity (higher in SIT) lead to disparate adaptations between men and women in the oxygen carrying capacity of blood (eg, total blood volume, haemoglobin or cardiac output) or local vasculature, but physiological mechanisms driving such responses are unclear.

Rates of fatty acid oxidation

In the untrained state, women have higher relative oxidative capacity than men,⁵² and higher relative rates of

fatty acid oxidation during prolonged exercise,⁵³ slower muscle phenotype with proportionally more type I fibres⁵⁴ and lower mitochondrial fractional synthetic rate after training.⁵⁵ Despite these sex differences in muscle metabolism and contractile characteristics, men and women in the present study showed similar gains in rates of fatty acid oxidation at all workloads from 30% to 60% VO_2max after training (figure 2). This metabolic adaptation occurred independently of changes to VO_2max and was not associated with the changes to body composition. Ours was the first study to look for sex differences in FATmax responses after SIT, but increased fat oxidation has previously been reported,^{2 3 16} and could be related to improved muscle oxidative capacity and enzymes of fatty acid oxidation.^{56–58} The underlying metabolic pathways coordinating these adaptations seem to be similar in SIT and conventional endurance exercise, with increased metabolic enzymes and capillarisation^{13 7 19} regulated in part by PGC-1 α , AMPk and CAMk.^{13 20} Elevated post exercise oxygen consumption may reflect increased rates of fat utilisation after high-intensity exercise.^{59 60} The elevated rates of fatty acid oxidation are associated with improved muscle metabolism⁶¹ and, alongside changes to blood lipid profiles, cardiopulmonary fitness and body composition, it is a defining feature of health status.⁶²

Fasting plasma glucose, insulin and lipid profiles

Levels of circulating total triglycerides in fasting plasma samples remained unchanged after training, while LDL and the cholesterol:HDL ratio improved. A reduction of total circulating triglycerides has been associated with regular aerobic exercise,^{63–65} although it is not a consistent finding in high-intensity training programmes,^{66 67} as noted in a recent review.⁶⁸ HDL increased after the first mixed exercise session, but did not change over the 12-week training. Fasting plasma glucose, insulin and the insulin sensitivity estimated using HOMA,³³ did not

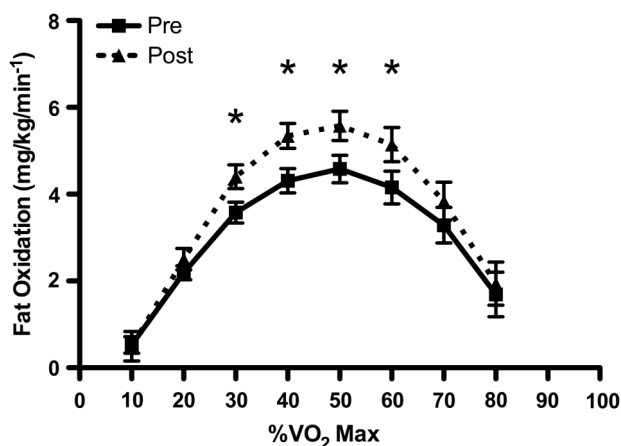


Figure 2 Maximal rates of fat oxidation in men and women across different submaximal exercise intensities in the untrained state and after 12 weeks of SIT. *Indicates statistically significant increase from baseline ($p < 0.05$). SIT, sprint interval training.

change significantly with training. Nor was change in fasting plasma glucose and insulin reported in a study of 16 healthy men after 6 SIT sessions, although those participants did improve glucose tolerance when measured using an oral glucose tolerance test.⁴ Hood *et al*⁶⁹ reported 35% improvement in HOMA in seven middle-aged men ($n=4$) and women ($n=3$) after six interval training sessions performed at lower intensity than that used in the present study. Richards *et al*⁷⁰ used the hyperinsulinaemic euglycaemic glucose clamp to show improved insulin sensitivity in 12 participants after 6 SIT sessions, and Whyte *et al*¹⁶ showed improved insulin sensitivity in 10 young, sedentary men after 6 SIT sessions. These previous reports outlining positive effects of SIT on glucose metabolism contrast with those from the present study.

Limitations

The semisupervised design of the training programme gave exercise volunteers more control, and although this is the case in real-life situations, it may confer less commitment or obligation to training compared with typical fully supervised laboratory-based programmes. We were not able to control for physical activities outside of the training programme and dietary intake was not monitored throughout the training programme. Instead, participants were asked to maintain their usual patterns of food and drink consumption. Finally, we did not control for menstrual cycle variations.

CONCLUSION

A total of 4 min exercise per week over 12 weeks improved cardiorespiratory, metabolic and body composition profiles of men and women recruited from the general population. Sex differences were observed, with men losing more fat mass than women and women showing higher VO_2max adaptations than men.

Acknowledgements The investigators would like to thank the participants for their involvement in this study, and The Midland Hotel (operated by Q Hotels) and gym staff for the use of their training facilities.

Contributors LB and JSM made contributions to the design, acquisition, analysis and interpretation of data; and drafted and critically revised the submitted work. MS made contribution to the conception and design of the study and interpretation of data; and critically revised the submitted work. SB made contribution to the acquisition, analysis and interpretation of data; and critically revised the submitted work. DL and MP made contributions to the acquisition of data; and critically revised the submitted work. CM made contribution to the design and interpretation of data; and critically revised the submitted work. GM made contribution to the acquisition of data; and critically revised the submitted work. MC made contribution to the design and interpretation of data; and critically revised the submitted work. WSG made contribution to the conception and design of the study; and critically revised the submitted work.

Funding This study was funded by Manchester Metropolitan University. JMcP is supported by UK Medical Research Council (MR/K025252/1).

Competing interests None declared.

Ethics approval Manchester Metropolitan University Local Research Ethics Committee (application number: SE111216).

Provenance and peer review Not commissioned; externally peer reviewed.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

REFERENCES

- Swift DL, Johannsen NM, Lavie CJ, *et al.* The role of exercise and physical activity in weight loss and maintenance. *Prog Cardiovasc Dis* 2014;56:441–7.
- Trapp EG, Chisholm DJ, Freund J, *et al.* The effects of high-intensity intermittent exercise training on fat loss and fasting insulin levels of young women. *Int J Obes (Lond)* 2008;32:684–91.
- Shepherd SO, Cocks M, Tipton KD, *et al.* Improvements in insulin sensitivity and whole-body fat oxidation after a period of high-intensity interval training. *Br J Sports Med* 2010;44:i11.
- Babraj JA, Vollaard NB, Keast C, *et al.* Extremely short duration high intensity interval training substantially improves insulin action in young healthy males. *BMC Endocr Disord* 2009;9:3.
- Little JP, Gillen JB, Percival ME, *et al.* Low-volume high-intensity interval training reduces hyperglycemia and increases muscle mitochondrial capacity in patients with type 2 diabetes. *J Appl Physiol* 2011;111:1554–60.
- Zoladz JA, Rademaker AC, Sargeant AJ. Human muscle power generating capability during cycling at different pedalling rates. *Exp Physiol* 2000;85:117–24.
- Talanian JL, Galloway SD, Heigenhauser GJ, *et al.* Two weeks of high-intensity aerobic interval training increases the capacity for fat oxidation during exercise in women. *J Appl Physiol* 2007;102:1439–47.
- Garber CE, Blissmer B, Deschenes MR, *et al.* American College of Sports Medicine. American College of Sports Medicine position stand. Quantity and quality of exercise for developing and maintaining cardiorespiratory, musculoskeletal, and neuromotor fitness in apparently healthy adults: guidance for prescribing exercise. *Med Sci Sports Exerc* 2011;43:1334–59.
- Gist NH, Fedewa MV, Dishman RK, *et al.* Sprint interval training effects on aerobic capacity: a systematic review and meta-analysis. *Sports Med* 2014;44:269–79.
- Sloth M, Sloth D, Overgaard K, *et al.* Effects of sprint interval training on VO₂max and aerobic exercise performance: a systematic review and meta-analysis. *Scand J Med Sci Sports* 2013;23:e341–52.
- Jensen L, Bangsbo J, Hellsten Y. Effect of high intensity training on capillarization and presence of angiogenic factors in human skeletal muscle. *J Physiol (Lond)* 2004;557:571–82.
- Iaia FM, Hellsten Y, Nielsen JJ, *et al.* Four weeks of speed endurance training reduces energy expenditure during exercise and maintains muscle oxidative capacity despite a reduction in training volume. *J Appl Physiol* 2009;106:73–80.
- Burgomaster KA, Howarth KR, Phillips SM, *et al.* Similar metabolic adaptations during exercise after low volume sprint interval and traditional endurance training in humans. *J Physiol (Lond)* 2008;586:151–60.
- Little JP, Safdar A, Bishop D, *et al.* An acute bout of high-intensity interval training increases the nuclear abundance of PGC-1 α and activates mitochondrial biogenesis in human skeletal muscle. *Am J Physiol Regul Integr Comp Physiol* 2011;300:R1303–10.
- Perry CG, Heigenhauser GJ, Bonen A, *et al.* High-intensity aerobic interval training increases fat and carbohydrate metabolic capacities in human skeletal muscle. *Appl Physiol Nutr Metab* 2008;33:1112–23.
- Whyte LJ, Gill JM, Cathcart AJ. Effect of 2 weeks of sprint interval training on health-related outcomes in sedentary overweight/obese men. *Metabolism* 2010;59:1421–8.
- Stensvold D, Tjønnå AE, Skaug EA, *et al.* Strength training versus aerobic interval training to modify risk factors of metabolic syndrome. *J Appl Physiol* 2010;108:804–10.
- Tremblay A, Simoneau JA, Bouchard C. Impact of exercise intensity on body fatness and skeletal muscle metabolism. *Metabolism* 1994;43:814–18.
- Shepherd SO, Cocks M, Tipton KD, *et al.* Sprint interval and traditional endurance training increase net intramuscular triglyceride breakdown and expression of perilipin 2 and 5. *J Physiol (Lond)* 2013;591:657–75.
- Gibala MJ, Little JP, Macdonald MJ, *et al.* Physiological adaptations to low-volume, high-intensity interval training in health and disease. *J Physiol (Lond)* 2012;590:1077–84.
- Macpherson RE, Hazell TJ, Olver TD, *et al.* Run sprint interval training improves aerobic performance but not maximal cardiac output. *Med Sci Sports Exerc* 2011;43:115–22.
- Hazell TJ, Hamilton CD, Olver TD, *et al.* Running sprint interval training induces fat loss in women. *Appl Physiol Nutr Metab* 2014;39:944–50.
- Gillen JB, Percival ME, Ludzki A, *et al.* Interval training in the fed or fasted state improves body composition and muscle oxidative capacity in overweight women. *Obesity (Silver Spring)* 2013;21:2249–55.
- Bijlsma AY, Meskers MC, Molendijk M, *et al.* Diagnostic measures for sarcopenia and bone mineral density. *Osteoporos Int* 2013;24:2681–91.
- McPhee JS, Maden-Wilkinson TM, Narici MV, *et al.* Knee extensor fatigue resistance of young and older men and women performing sustained and brief intermittent isometric contractions. *Muscle Nerve* 2014;50:393–400.
- Simoneau JA, Lortie G, Boulay MR, *et al.* Skeletal muscle histochemical and biochemical characteristics in sedentary Male and female subjects. *Can J Physiol Pharmacol* 1985;63:30–5.
- Venables MC, Achten J, Jeukendrup AE. Determinants of fat oxidation during exercise in healthy men and women: a cross-sectional study. *J Appl Physiol* 2005;98:160–7.
- Esbjörnsson-Liljedahl M, Sundberg CJ, Norman B, *et al.* Metabolic response in type I and type II muscle fibers during a 30-s cycle sprint in men and women. *J Appl Physiol* 1999;87:1326–32.
- Association WM. World Medical Association Declaration of Helsinki: ethical principles for medical research involving human subjects. *JAMA* 2013;310:2191–4.
- McPhee JS, Hogrel JY, Maier AB, *et al.* Physiological and functional evaluation of healthy young and older men and women: design of the European MyoAge study. *Biogerontology* 2013;14:325–37.
- Maden-Wilkinson TM, Degens H, Jones DA, *et al.* Comparison of MRI and DXA to measure muscle size and age-related atrophy in thigh muscles. *J Musculoskelet Neuronal Interact* 2013;13:320–8.
- Frayn KN. Calculation of substrate oxidation rates in vivo from gaseous exchange. *J Appl Physiol Respir Environ Exerc Physiol* 1983;55:628–34.
- Matthews DR, Hosker JP, Rudenski AS, *et al.* Homeostasis model assessment: insulin resistance and beta-cell function from fasting plasma glucose and insulin concentrations in man. *Diabetologia* 1985;28:412–19.
- Kelly TL, Wilson KE, Heymsfield SB. Dual energy X-Ray absorptiometry body composition reference values from NHANES. *PLoS ONE* 2009;4:e7038.
- Heo M, Faith MS, Pietrobelli A, *et al.* Percentage of body fat cutoffs by sex, age, and race-ethnicity in the US adult population from NHANES 1999–2004. *Am J Clin Nutr* 2012;95:594–602.
- Donnelly JE, Smith BK. Is exercise effective for weight loss with ad libitum diet? Energy balance, compensation, and gender differences. *Exerc Sport Sci Rev* 2005;33:169–74.
- Devries MC. Sex-based differences in endurance exercise muscle metabolism: impact on exercise and nutritional strategies to optimize health and performance in women. *Exp Physiol* 2016;101:243–9.
- Tamopolsky MA. Sex differences in exercise metabolism and the role of 17- β -estradiol. *Med Sci Sports Exerc* 2008;40:648–54.
- Caudwell P, Gibbons C, Hopkins M, *et al.* No sex difference in body fat in response to supervised and measured exercise. *Med Sci Sports Exerc* 2013;45:351–8.
- Henderson GC, Alderman BL. Determinants of resting lipid oxidation in response to a prior bout of endurance exercise. *J Appl Physiol* 2014;116:95–103.
- Henderson GC. Sexual dimorphism in the effects of exercise on metabolism of lipids to support resting metabolism. *Front Endocrinol (Lausanne)* 2014;5:162.
- Donnelly JE, Hill JO, Jacobsen DJ, *et al.* Effects of a 16-month randomized controlled exercise trial on body weight and composition in young, overweight men and women: the Midwest Exercise Trial. *Arch Intern Med* 2003;163:1343–50.
- Treuth MS, Hunter GR, Williams M. Effects of exercise intensity on 24-h energy expenditure and substrate oxidation. *Med Sci Sports Exerc* 1996;28:1138–43.
- Thompson DA, Wolfe LA, Eikelboom R. Acute effects of exercise intensity on appetite in young men. *Med Sci Sports Exerc* 1988;20:222–7.
- Pomerleau M, Imbeault P, Parker T, *et al.* Effects of exercise intensity on food intake and appetite in women. *Am J Clin Nutr* 2004;80:1230–6.
- Bouchard C, An P, Rice T, *et al.* Familial aggregation of VO₂(max) response to exercise training: results from the HERITAGE Family Study. *J Appl Physiol* 1999;87:1003–8.

47. Scalzo RL, Peltonen GL, Binns SE, *et al.* Greater muscle protein synthesis and mitochondrial biogenesis in males compared with females during sprint interval training. *FASEB J* 2014;28:2705–14.
48. Allemeier CA, Fry AC, Johnson P, *et al.* Effects of sprint cycle training on human skeletal muscle. *J Appl Physiol* 1994;77:2385–90.
49. Talanian JL, Holloway GP, Snook LA, *et al.* Exercise training increases sarcolemmal and mitochondrial fatty acid transport proteins in human skeletal muscle. *Am J Physiol Endocrinol Metab* 2010;299:E180–8.
50. Ogawa T, Spina RJ, Martin WH, *et al.* Effects of aging, sex, and physical training on cardiovascular responses to exercise. *Circulation* 1992;86:494–503.
51. Saltin B, Calbet JA. Point: in health and in a normoxic environment, VO₂ max is limited primarily by cardiac output and locomotor muscle blood flow. *J Appl Physiol* 2006;100:744–5.
52. Maughan RJ, Harmon M, Leiper JB, *et al.* Endurance capacity of untrained males and females in isometric and dynamic muscular contractions. *Eur J Appl Physiol Occup Physiol* 1986;55:395–400.
53. Carter SL, Rennie C, Tarnopolsky MA. Substrate utilization during endurance exercise in men and women after endurance training. *Am J Physiol Endocrinol Metab* 2001;280:E898–907.
54. Simoneau JA, Lortie G, Boulay MR, *et al.* Human skeletal muscle fiber type alteration with high-intensity intermittent training. *Eur J Appl Physiol Occup Physiol* 1985;54:250–3.
55. Karakelides H, Irving BA, Short KR, *et al.* Age, obesity, and sex effects on insulin sensitivity and skeletal muscle mitochondrial function. *Diabetes* 2010;59:89–97.
56. Holloszy JO, Coyle EF. Adaptations of skeletal muscle to endurance exercise and their metabolic consequences. *J Appl Physiol Respir Environ Exerc Physiol* 1984;56:831–8.
57. Hoppeler H, Howald H, Conley K, *et al.* Endurance training in humans: aerobic capacity and structure of skeletal muscle. *J Appl Physiol* 1985;59:320–7.
58. Gollnick PD, Armstrong RB, Saltin B, *et al.* Effect of training on enzyme activity and fiber composition of human skeletal muscle. *J Appl Physiol* 1973;34:107–11.
59. Børsheim E, Bahr R. Effect of exercise intensity, duration and mode on post-exercise oxygen consumption. *Sports Med* 2003;33:1037–60.
60. LaForgia J, Withers RT, Gore CJ. Effects of exercise intensity and duration on the excess post-exercise oxygen consumption. *J Sports Sci* 2006;24:1247–64.
61. Jeukendrup AE, Saris WH, Wagenmakers AJ. Fat metabolism during exercise: a review. Part I: fatty acid mobilization and muscle metabolism. *Int J Sports Med* 1998;19:231–44.
62. Wolfe RR. The underappreciated role of muscle in health and disease. *Am J Clin Nutr* 2006;84:475–82.
63. Plaisance EP, Mestek ML, Mahurin AJ, *et al.* Postprandial triglyceride responses to aerobic exercise and extended-release niacin. *Am J Clin Nutr* 2008;88:30–7.
64. Durstine JL, Grandjean PW, Davis PG, *et al.* Blood lipid and lipoprotein adaptations to exercise: a quantitative analysis. *Sports Med* 2001;31:1033–62.
65. Crouse SF, O'Brien BC, Grandjean PW, *et al.* Effects of training and a single session of exercise on lipids and apolipoproteins in hypercholesterolemic men. *J Appl Physiol* 1997;83:2019–28.
66. Tsekouras YE, Magkos F, Kellas Y, *et al.* High-intensity interval aerobic training reduces hepatic very low-density lipoprotein-triglyceride secretion rate in men. *Am J Physiol Endocrinol Metab* 2008;295:E851–8.
67. Bellou E, Magkos F, Kouka T, *et al.* Effect of high-intensity interval exercise on basal triglyceride metabolism in non-obese men. *Appl Physiol Nutr Metab* 2013;38:823–9.
68. Kessler HS, Sisson SB, Short KR. The potential for high-intensity interval training to reduce cardiometabolic disease risk. *Sports Med* 2012;42:489–509.
69. Hood MS, Little JP, Tarnopolsky MA, *et al.* Low-volume interval training improves muscle oxidative capacity in sedentary adults. *Med Sci Sports Exerc* 2011;43:1849–56.
70. Richards JC, Johnson TK, Kuzma JN, *et al.* Short-term sprint interval training increases insulin sensitivity in healthy adults but does not affect the thermogenic response to beta-adrenergic stimulation. *J Physiol (Lond)* 2010;588:2961–72.