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## **A comparison of meal-related appetite, food reward and eating behaviour traits in people with and without spinal cord injury**

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1 **Title: A comparison of meal-related appetite, food reward and eating behaviour traits in people**  
2 **with and without spinal cord injury**

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36 **Abstract**

37 Persons with a spinal cord injury (SCI) are at a heightened risk of obesity. However, little is known  
38 about the effect of SCI on factors that influence energy intake. This study compared measures of food  
39 reward, eating behaviour traits, and appetite perceptions between adults with and without SCI. Twenty  
40 wheelchair dependent persons with chronic (>1 year) SCI (C1-T12) and twenty non-SCI individuals  
41 matched for BMI, age and sex participated. Following a familiarisation visit, participants consumed a  
42 breakfast meal, normalised for resting metabolic rate (RMR), and provided subjective appetite  
43 perceptions every 30 min for 4 h. Subsequently, energy intake was determined via an *ad libitum* lunch  
44 meal. Explicit liking, explicit wanting, implicit wanting and relative preference were assessed in a  
45 hungry and fed state via the Leeds Food Preference Questionnaire prior to and following the lunch meal.  
46 Eating behaviour traits were assessed via the Adult Eating Behaviour Questionnaire, Control of Eating  
47 Questionnaire, Reasons Individuals Stop Eating Questionnaire, and Three Factor Eating Questionnaire  
48 Revised 18-item version. Sweet appeal bias was greater for explicit liking, explicit wanting, and relative  
49 preference in the group with SCI compared to the non-SCI group ( $p \leq 0.024$ ). The group with SCI also  
50 reported higher levels of cognitive restraint and satiety responsiveness ( $p \leq 0.029$ ). No group differences  
51 in postprandial appetite perceptions ( $p \geq 0.690$ ) or energy intake relative to RMR were seen ( $p = 0.358$ ).  
52 However, the group with SCI demonstrated a trend toward a lower absolute energy intake ( $p = 0.063$ ).  
53 In conclusion, food reward for sweet foods was greater in the group with SCI. Further, our findings  
54 suggest that acute appetite perceptions, including satiety profiles, are not different between persons with  
55 and without SCI.

56 **Keywords:** paraplegia; food preference; satiety; energy intake.

57 **Abbreviations:** BMI, body mass index; LFPQ, Leeds Food Preference Questionnaire; RMR, resting  
58 metabolic rate; SCI, spinal cord injury; VAS, visual analogue scale.

59 **1. Introduction**

60 A spinal cord injury (SCI) is a medical condition wherein damage to the spinal cord impairs motor,  
61 sensory and autonomic function below the level of injury. Paralysis and increased reliance on a  
62 wheelchair for locomotion results in a substantial reduction in physical activity energy expenditure.  
63 Further, a decrease in resting metabolic rate (RMR), due to the loss of metabolically active tissue, such  
64 as skeletal muscle mass, prompt substantial reductions in total daily energy expenditure, predisposing  
65 this population to obesity (Gater, 2007; Gorgey et al., 2014). Research in non-SCI populations  
66 demonstrates that people with low daily energy expenditure exhibit signs of impaired appetite  
67 regulation, with difficulty matching daily energy intake and expenditure (Hopkins & Blundell, 2016).  
68 Given physical limitations, it is possible that appetite may become desensitised in people with SCI;  
69 however, few studies have investigated this issue.

70 Appetite and eating behaviour are influenced by interacting physiological and psychological factors  
71 (Hopkins et al., 2016). Specifically, a network of appetite-related hormones is known to modulate  
72 feelings of hunger, satiation, and satiety; with acute (meal-to-meal) and chronic influences. In the  
73 context of SCI, evidence suggests that circulating concentrations of acylated ghrelin, peptide YY, and  
74 glucagon-like peptide-1 are not different between persons with or without SCI, in a fasted state (Fenton  
75 et al., 2021; Saltzstein et al., 1995; Wang et al., 2005). In our recent comparative study, we also observed  
76 no differences between groups with and without SCI in the postprandial trends of these hormones  
77 following an isocaloric preload over 2.5 h (Fenton et al., 2021). However, visual analogue scales (VAS)  
78 revealed accentuated satiety responses in the group with SCI in the first hour postprandially. This  
79 outcome may reflect undefined population differences in gastrointestinal physiology, and/or meal-  
80 related sensory perceptions / cognitive processes. Alternatively, in our study, the *relatively* larger meal  
81 size consumed by those with SCI may be explanative. Further experiments are needed to confirm  
82 whether SCI is associated with enhanced satiety responsiveness and to explore the potential contribution  
83 of psychological mediators.

84 Within the discipline of biological psychology, ‘food reward’ considers the hedonic aspects of appetite  
85 and eating behaviour. Food reward is underpinned by momentary “liking”, the subjective pleasure of a  
86 food, and “wanting”, the motivational value of a food (Finlayson & Dalton, 2012; Oustric et al., 2020),  
87 and is commonly assessed using the Leeds Food Preference Questionnaire (LFPQ) (Finlayson et al.,  
88 2007). Evidence suggests that people who are inactive and living with overweight/obesity often exhibit  
89 greater liking and wanting scores for energy-dense foods, compared to active individuals with a healthy  
90 body weight (Finlayson & Dalton, 2012; Horner et al., 2016; Oustric et al., 2018). These hedonic  
91 characteristics have been shown to encourage overconsumption, thus amplifying the rate of weight gain  
92 (Beaulieu et al., 2018). Given that excess adiposity and physical inactivity are common in people with

93 SCI (Spungen et al., 2003; van den Berg-Emons et al., 2010), it is possible that disrupted food reward  
94 is apparent in this population. However, this possibility has not been investigated.

95 Other psychological factors, such as eating behaviour traits, influence food choice and portion size  
96 selection. For example, positive associations between ‘food approach’ traits, such as uncontrolled eating  
97 and appetite-related disinhibition, have been reported in persons with higher adiposity and lower levels  
98 of physical activity (Myers et al., 2017; Shook et al., 2015). Likewise, ‘food avoidance’ traits, such as  
99 satiety responsiveness and slowness in eating, are associated with a lower body mass index (BMI)  
100 (Mallan et al., 2017). Although non-ambulatory persons with SCI often exhibit low physical activity  
101 energy expenditure and excess adiposity SCI (Spungen et al., 2003; van den Berg-Emons et al., 2010),  
102 a quantitative assessment of eating behaviour traits has not been conducted in the general SCI  
103 population.

104 Based on the aforementioned evidence, this study sought to compare meal-related appetite perceptions,  
105 measures of satiety profile, and food reward responses between persons with and without SCI. Potential  
106 differences in eating behaviour traits were also examined. We hypothesised that no group differences  
107 in meal-related perceptions of hunger, satiation, and satiety would be apparent when the energy content  
108 of a test meal was scaled to individuals’ daily energy requirements. Moreover, people with SCI would  
109 display greater motivation and preference for energy dense foods (sweet and fatty) than those without  
110 SCI. Finally, persons with SCI would exhibit more ‘food approach’ eating behaviour traits, and fewer  
111 ‘food avoidance traits compared to the non-SCI group.

## 112 **2. Methods**

### 113 *2.1. Ethical approval and participants*

114 Following approval from the Loughborough University Ethics Review Sub-Committee (project ID:  
115 5574), 20 participants with chronic (>1-year post-injury) SCI and 20 age-, sex- and BMI-matched non-  
116 SCI individuals provided written informed consent to participate. Inclusion criteria were non-smoking  
117 men and women, aged 18-64 years, metabolically, weight stable (body mass  $\pm$  3 kg for the past 3  
118 months), and habitual breakfast and lunch consumers. Trained athletes, persons following vegan or  
119 vegetarian diets, and those with allergies, intolerances, or a dislike to any of the foods included in the  
120 study were excluded. All female participants reported being eumenorrheic and not pregnant.  
121 Classification of SCI was self-reported, with motor complete SCI defined as a complete loss of motor  
122 function below the lesion level, and incomplete SCI defined as partial loss of motor and sensory  
123 function. All participants with incomplete SCI were wheelchair dependent for daily ambulation.  
124 Characteristics of the study participants are shown in Table 1. Given the exploratory nature of this study,  
125 a formal power calculation was not conducted. Instead, sample size was determined from previous  
126 studies using similar study designs (Beaulieu et al., 2020; Horner et al., 2016).

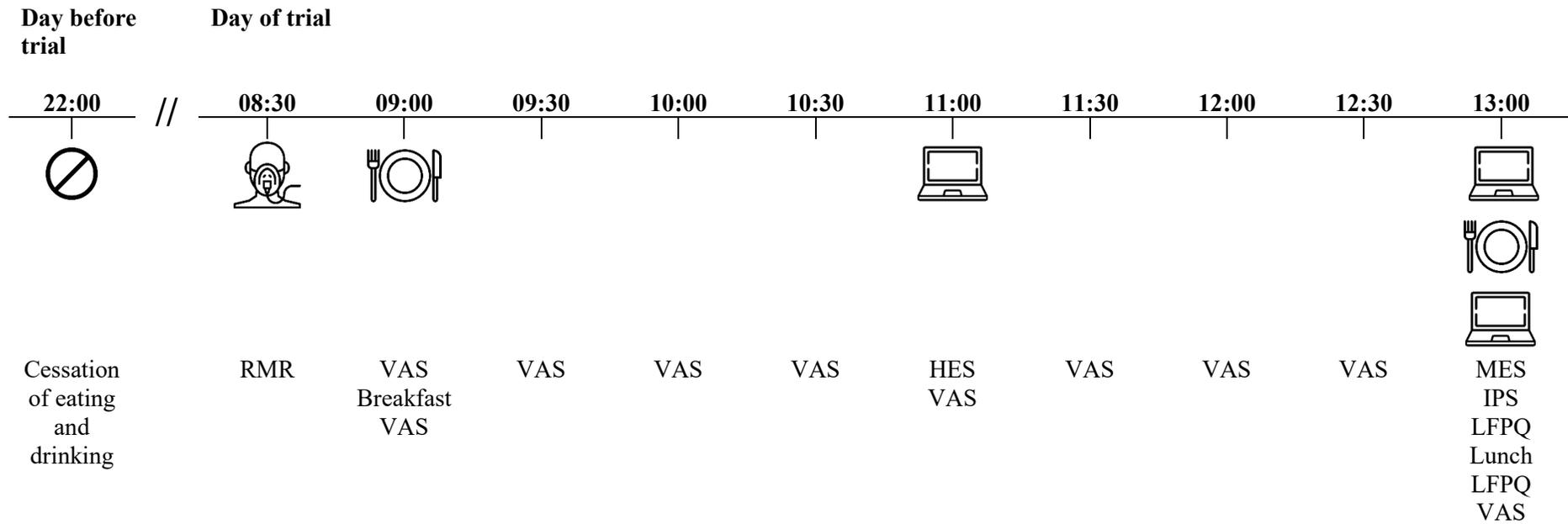
127 2.2. *Pre-assessment and familiarisation*

128 Participants attended the laboratory on two occasions for a familiarisation visit and the experimental  
129 trial. In the familiarisation visit body mass of the participants with SCI was measured using a wheelchair  
130 weighing scale (DETECTO 6550, Missouri, USA). Height and SCI-related information (time since  
131 injury, injury level and injury completeness) was self-reported. Supine waist and hip circumferences  
132 were measured using an inelastic polyfibre tape measure (Hokanson, Washington, USA) for all  
133 participants. In the non-SCI group, measures of stature and body mass were performed using a portable  
134 stadiometer and digital scale (Seca Ltd, Hamburg, Germany). All measures were made in duplicate and  
135 to the nearest 0.1 cm or 0.1 kg. Participants were then familiarised with the study protocol, the LFPQ,  
136 the 100 mm visual analogue scale(s) (VAS) used for subjective appetite measures (Flint et al., 2000),  
137 and the *ad libitum* lunch meal.

138 Skinfold thickness was measured on the biceps, triceps, subscapular, iliac crest, supraspinale,  
139 abdominal, front thigh and medial calf on the right-hand side of the body in accordance with the  
140 International Society for the Advancement of Kinanthropometry (ISAK), using Harpenden skinfold  
141 callipers (HaB Direct, Southam, UK) (Norton, 2019), as previously used by our group (Goosey-Tolfrey  
142 et al., 2021; Sutton et al., 2009). To permit group comparisons, for all participants, skinfolds were  
143 measured whilst sat in a chair/wheelchair.

144 2.3. *Study design*

145 Participants were instructed to abstain from strenuous exercise, caffeine, and alcohol in the 24 h prior  
146 to arriving at the laboratory for the main trial and were asked to stop eating and drinking (other than  
147 water) from 22:00. Participants arrived at 08:00 for the experimental trial (Figure 1) and were asked to  
148 rest on a bed in a supine position whilst RMR was measured. Participants were then provided with  
149 breakfast. Appetite rating VAS were completed immediately before and after breakfast, and every 0.5  
150 h thereafter for 4 h. During this period, participants were asked to remain seated in a quiet room and  
151 were permitted to undertake sedentary tasks. Participants were also asked to complete several  
152 questionnaires related to their eating behaviour during this time. At 2 h post-breakfast, participants  
153 completed the hypothetical expected satiety task. After 4 h, participants underwent the momentary  
154 expected satiety task, the ideal portion size task, and the pre-meal LFPQ. They were then asked to  
155 complete a VAS for perceived appetite and consume the *ad libitum* lunch meal. This was succeeded by  
156 the post-meal LFPQ and a final VAS.



157

158 **Figure 1.** Schematic illustration of the experiment protocol (HES, hypothetical expected satiety; IPS, ideal portion size; LFPQ, Leeds food preference  
 159 questionnaire; RMR, resting metabolic rate; VAS, visual analogue scale).

#### 160 2.4. Study foods

161 Breakfast consisted of a ground oat porridge meal (Ready Brek, UK), whole milk and honey (56%  
162 carbohydrate, 28% fat, 14% protein), with an energy content calculated relative to the estimated daily  
163 energy requirements of each participant (20% of participants' estimated energy requirements) (Clayton  
164 et al., 2016). Given the sedentary nature of the protocol, estimated energy requirements were determined  
165 by multiplying RMR by a sedentary activity factor of 1.2 (Collins et al., 2010; Jetté et al., 1990;  
166 Nightingale et al., 2017).

167 The lunch meal consisted of cheese (medium cheddar), passata, fusilli pasta and olive oil (all Tesco,  
168 UK). Participants were presented with the pasta in a large bowl, served at room temperature, and in  
169 excess of predicted consumption (6188 kJ, 44% carbohydrate, 41% fat, 14% protein) and were  
170 instructed to eat until comfortably full. The meals were consumed in isolation (other than the  
171 participants with SCI that required assistance) and the use of electronic devices during the eating period  
172 was prohibited (Braude & Stevenson, 2014). The mass of food consumed was determined by subtracting  
173 the mass of food remaining from that initially presented. Participants were offered a 150 ml glass of  
174 water with the study foods and were told to drink *ad libitum* throughout the day.

#### 175 2.5. Leeds Food Preference Questionnaire

176 The LFPQ is a validated laptop-based questionnaire that assesses different components of food  
177 preference and reward (Finlayson et al., 2007). A detailed description of how the LFPQ is conducted  
178 and analysed has been published (Dalton & Finlayson, 2014). To summarise, sixteen food items from  
179 four different categories (high-fat savoury, high-fat sweet, low-fat savoury, low-fat sweet) were utilised  
180 in the questionnaire. Explicit liking and explicit wanting were assessed by asking participants how much  
181 they would 'like to taste', and 'want to eat', respectively, some of the food presented in front of them  
182 at that moment in time using a 100 mm VAS. A score of 0 represented "Not at all", and a score of 100  
183 represented "Extremely". Relative preference was assessed by presenting participants with two foods  
184 and asking them to choose which food they 'most want to eat now'. Scores for relative preferences were  
185 ascertained by the mean frequency of selection for each food type. Implicit wanting was determined by  
186 the number of times a certain category of food was and was not chosen, as well as the time taken to  
187 choose between the two foods. 'Fat appeal bias' and 'sweet appeal bias' scores were determined by  
188 deducting the low-fat scores from the high-fat scores, and the savoury scores from the sweet scores,  
189 respectively.

#### 190 2.6. Eating behaviour traits

191 Four eating behaviour trait questionnaires were implemented in this study. These included the Adult  
192 Eating Behaviour Questionnaire (Hunot et al., 2016), the Control of Eating Questionnaire (Dalton et

193 al., 2015), the Reasons Individuals Stop Eating Questionnaire (Cunningham et al., 2021), and the Three-  
194 Factor Eating Questionnaire – Revised 18-item version (de Lauzon et al., 2004).

### 195 *2.7. Appetite perceptions*

196 Ratings of perceived hunger, satisfaction, fullness, and prospective food consumption were assessed  
197 using a 100 mm VAS (Flint et al., 2000). Pleasantness and satisfaction of the breakfast and lunch meals  
198 were also rated immediately after participants had finished eating. Each scale defined 0 mm as “not at  
199 all” and 100 mm as “totally / a lot / very”, and participants were asked to put a vertical line at the point  
200 corresponding to their appetite at that point in time.

### 201 *2.8. Expected satiety and ideal portion size*

202 Expected satiety measures and ideal portion size were determined using the method of adjustment,  
203 which has been described previously (Brunstrom & Rogers, 2009; McLeod et al., 2022). To assess  
204 hypothetical expected satiety participants were presented with a laptop which displayed a ‘standard’  
205 food (1255 kJ portion of plain pilau rice) and one of six ‘comparison’ foods (porridge, granola, chicken  
206 chow mein, cheese and tomato pasta, crisps, or chocolate). Participants were instructed to change the  
207 portion size of the comparison foods so that both foods would stave off hunger to the same extent. For  
208 momentary expected satiety and ideal portion size, participants were presented with an image of cheese  
209 and tomato pasta that could be adjusted for portion size. Participants were asked to select a portion that  
210 would stop them feeling hungry until their next meal in 5 hours’ time, and a portion that they would eat  
211 right now if it was the only food available until their next meal in 5 hours’ time, respectively. Portion  
212 sizes for each comparison food ranged from 0-4184 kJ.

### 213 *2.9. Energy expenditure and physical activity status*

214 Resting metabolic rate was determined using a mobile spiroergometric device (Metamax 3B, Cortex,  
215 Germany) and MetaSoft Studio software (Cortex, Germany) in accordance with best practise guidelines  
216 (Compher et al., 2006). Participants lay in a supine position for 30 min whilst the device measured  
217 oxygen consumption ( $\dot{V}O_2$ ), carbon dioxide output ( $\dot{V}CO_2$ ) and the volume of expired air on a second-  
218 by-second basis, with the last 10 min used for analysis. Carbohydrate and fat oxidation were determined  
219 from the volume of  $\dot{V}O_2$  and  $\dot{V}CO_2$ , with the assumption that protein oxidation at rest was negligible  
220 (Frayn, 1983). Energy expenditure was then calculated using a caloric equivalent for carbohydrate and  
221 fat oxidation (Ferrannini, 1988) and averaged over the last 10 min. This value was extrapolated over 24  
222 h to determine RMR. Physical activity status was assessed using the Leisure Time Physical Activity  
223 Questionnaire for People with Spinal Cord Injury (Ginis et al., 2012), and the International Physical  
224 Activity Questionnaire – Short Form (Craig et al., 2003) for the non-SCI group.

### 225 *2.10. Statistical analysis*

226 Data were analysed using the Statistical Package for Social Sciences (SPSS) software version 27 for  
227 Windows (IBM Corporation, USA). Preceding analysis, tests of normality and homogeneity were run  
228 to assess parametric assumptions. For the LFPQ data, two-way mixed ANOVAs were used to determine  
229 group (SCI vs. non-SCI) time (pre- vs. post-lunch) effects for relative preference, implicit wanting,  
230 explicit liking and explicit wanting. A two-way mixed ANOVA was also used to explore the interaction  
231 between momentary expected satiety and ideal portion size, as well as between group effects. For  
232 hypothetical expected satiety, a repeated measures ANOVA was used to explore the effect of group and  
233 food as fixed factors. Similarly, differences in appetite perceptions over time were assessed using  
234 repeated measures ANOVAs with group and time. Post-hoc Bonferroni corrected pairwise comparisons  
235 were conducted for all general linear models when a main effect was detected. Given the anticipated  
236 differences in lean mass between groups, absolute energy intake was also analysed relative to RMR  
237 ((energy intake / RMR) \* 100). Independent samples t-tests and Mann-Whitney U tests were used to  
238 compare all other parametric and non-parametric data between groups, respectively. Test-specific effect  
239 sizes (Cohen's  $d$ ) were calculated to supplement each parametric statistical test, with large, moderate,  
240 small, and trivial effect sizes defined as 0.8, 0.5, 0.2, and  $< 0.2$ , respectively (Cohen, 1988). For mixed  
241 ANOVAs, effect sizes were determined using partial eta squared ( $\eta_p^2$ ), with 0.14, 0.06 and 0.01 defined  
242 as large, medium, and small effect sizes, respectively. For non-parametric data, effect sizes were  
243 calculated using Rosenthal's  $r$  (Rosenthal, 2011), with large, medium, and small effects classified as  
244 0.5, 0.3, and 0.1 respectively. Mean  $\pm$  SD was reported for all parametric data, whereas the median  
245 (IQR) was reported for non-parametric data. Statistical significance was defined as  $p < 0.05$ .

### 246 **3. Results**

#### 247 *3.1. Demographic information*

248 Demographic data for the 40 participants are reported in Table 1. There were no differences in age,  
249 body mass, height, BMI, or supine hip circumference between the SCI and non-SCI groups ( $p \geq 0.092$ ;  
250  $d \leq 0.56$ ). However, supine waist circumference ( $t(38) = -2.465$ ;  $p = 0.018$ ;  $d = 0.78$ ) and supine waist-  
251 to-hip ratio ( $t(38) = 2.437$ ;  $p = 0.020$ ;  $d = 0.77$ ) were greater in the group with SCI, with  $\Sigma$ SF8  
252 approaching between-group significance ( $t(38) = 1.948$ ;  $p = 0.059$ ;  $d = 0.62$ ). RMR was also lower in  
253 the group with SCI compared to the non-SCI group ( $t(38) = -2.188$ ;  $p = 0.035$ ;  $d = 0.69$ ).

**Table 1**

Participant demographic information.

Group	SCI, n = 20	Non-SCI, n = 20
Sex	Male, n = 14	Male, n = 14
	Female, n = 6	Female, n = 6
Age (yr)	37 (28-49)	30 (26-48)
Body mass (kg)	79.2 ± 17.4	73.8 ± 14.5
Height (cm)	176.0 ± 0.1	173.0 ± 0.1
BMI (kg.m <sup>-2</sup> )	25.5 ± 5.2	24.6 ± 3.2
Resting metabolic rate (kJ.day <sup>-1</sup> )	6736 ± 1084*	7719 ± 1703
Hip circumference - supine (cm)	104.5 ± 16.0	97.8 ± 5.6
Waist circumference - supine (cm)	92.7 ± 15.3 *	82.7 ± 9.5
Waist-to-hip ratio - supine	0.91 ± 0.1 *	0.84 ± 0.1
Sum of eight skinfold sites (mm)	152.8 ± 56.4	121.7 ± 43.6
Time spent sitting (min.week <sup>-1</sup> ) <sup>+</sup>		3465 (2940-4200)
Mild exercise / walking (min.week <sup>-1</sup> ) <sup>+</sup>	140 (23-360)	225 (203-420)
Moderate exercise (min.week <sup>-1</sup> ) <sup>+</sup>	120 (30-180)	120 (60-203)
Heavy / vigorous exercise (min.week <sup>-1</sup> ) <sup>+</sup>	99 ± 105	155 ± 135
Injury duration (yr)	11.5 ± 9.9	
Injury category (AIS)	A, n = 7	
	B, n = 4	
	C, n = 9	
Injury level	Paraplegia (T2-T12), n = 8	
	Tetraplegia (C1-C7), n = 12	

Mean ± SD; Median (IQR).

\* Significant difference between groups ( $p < 0.05$ ); <sup>+</sup> Different questionnaires were used to assess physical activity status in these groups, meaning results were not directly comparable.

AIS, American Spinal Injury Association Impairment Scale; BMI, body mass index; C, cervical; SCI, spinal cord injury; T, thoracic.

254

255 *3.2. Food preference and reward*

256 For sweet appeal bias scores, there were significant group effects for explicit liking, explicit wanting,  
 257 and relative preference ( $p \leq 0.024$ ;  $\eta_p^2 \geq 0.127$ ), with implicit wanting approaching significance ( $p =$   
 258  $0.059$ ;  $\eta_p^2 = 0.091$ ) (Table 2). Both groups demonstrated a shift in sweet appeal bias for explicit liking,  
 259 explicit wanting, implicit wanting and relative preference from pre- to post-meal which inferred a  
 260 stronger bias for sweet foods following the lunch meal ( $p < 0.001$ ;  $\eta_p^2 \geq 0.569$ ). However, no interaction

261 effects were found for any sweet appeal bias metric ( $p \geq 0.501$ ;  $\eta_p^2 \leq 0.01$ ). For fat appeal bias scores,  
262 explicit liking, explicit wanting, implicit wanting and relative preference were not different between  
263 pre-lunch and post-lunch time points ( $p \geq 0.127$ ;  $\eta_p^2 \leq 0.060$ ). Similarly, no group or interaction effects  
264 were seen for these any of these metrics ( $p \geq 0.181$ ;  $\eta_p^2 \leq 0.047$ ).

**Table 2**

Measures of explicit liking, explicit wanting, implicit wanting and relative preference assessed immediately before and after an *ad libitum* lunch meal in persons with a spinal cord injury (SCI; n = 20) and non-SCI (n = 20) (no interaction main effects were found ( $p \geq 0.501$ ;  $\eta_p^2 \leq 0.01$ ) and data were omitted for clarity).

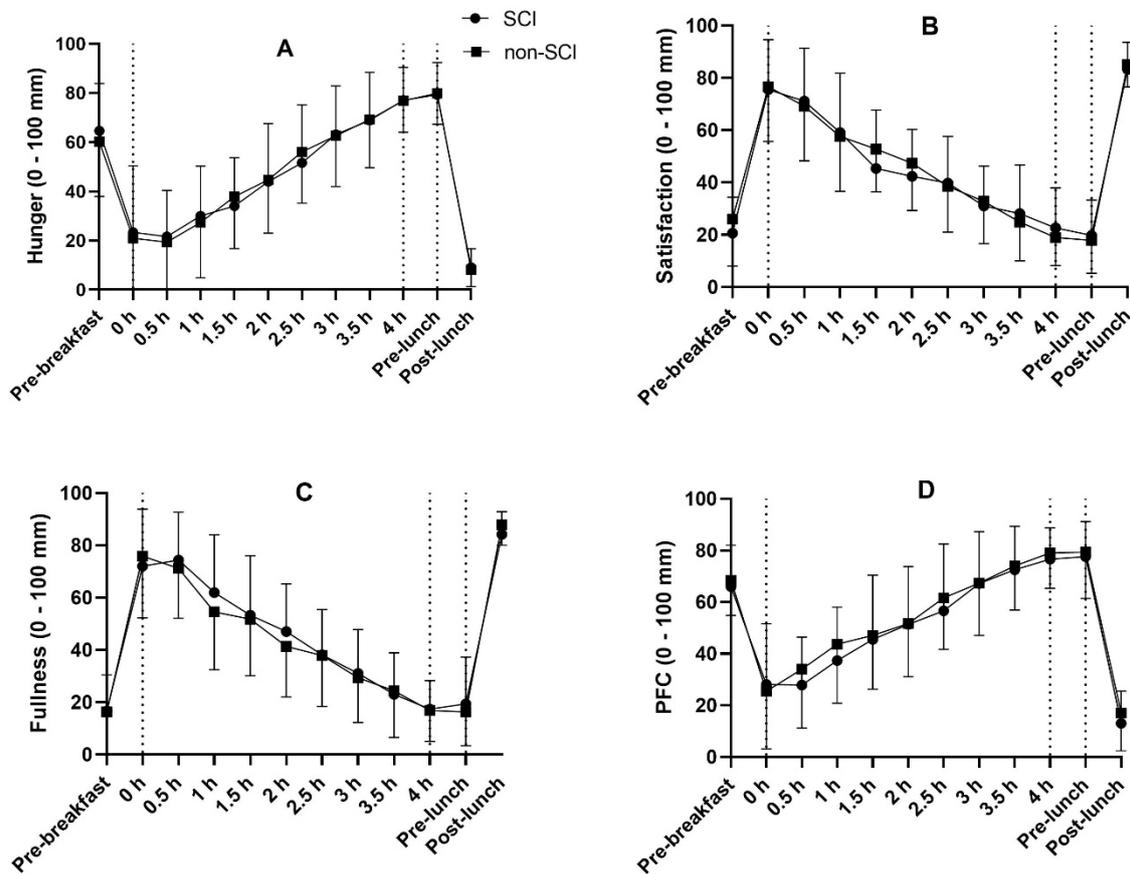
	Pre-lunch		Post-lunch		Group effect			Time effect		
	SCI	non-SCI	SCI	non-SCI	F <sub>(1,38)</sub>	p	$\eta_p^2$	F <sub>(1,38)</sub>	p	$\eta_p^2$
<b>Explicit liking</b>										
Sweet appeal bias	1 ± 18	-17 ± 22	35 ± 22	23 ± 20	14.247	0.001*	0.451	50.129	< 0.001*	0.569
Fat appeal bias	4 ± 12	4 ± 9	5 ± 11	4 ± 15	0.005	0.945	<0.001	0.026	0.872	0.001
<b>Explicit wanting</b>										
Sweet appeal bias	-3 ± 20	-20 ± 20	32 ± 24	22 ± 15	12.743	0.001*	0.251	57.674	< 0.001*	0.603
Fat appeal bias	3 ± 13	3 ± 12	5 ± 8	3 ± 12	0.14	0.710	0.004	0.208	0.651	0.005
<b>Implicit wanting</b>										
Sweet appeal bias	-14 ± 47	-32 ± 35	63 ± 26	48 ± 32	3.799	0.059	0.091	104.692	< 0.001*	0.734
Fat appeal bias	14 ± 30	6 ± 27	16 ± 17	12 ± 19	1.066	0.308	0.027	0.68	0.415	0.018
<b>Relative preference</b>										
Sweet appeal bias	-5 ± 18	-13 ± 13	24 ± 10	17 ± 11	5.507	0.024*	0.127	112.414	< 0.001*	0.747
Fat appeal bias	5 ± 11	2 ± 10	7 ± 7	5 ± 8	1.855	0.181	0.047	2.429	0.127	0.060

Values are mean ± SD.

\* Significant difference ( $p < 0.05$ ).

266 3.3. Ratings of perceived appetite and palatability

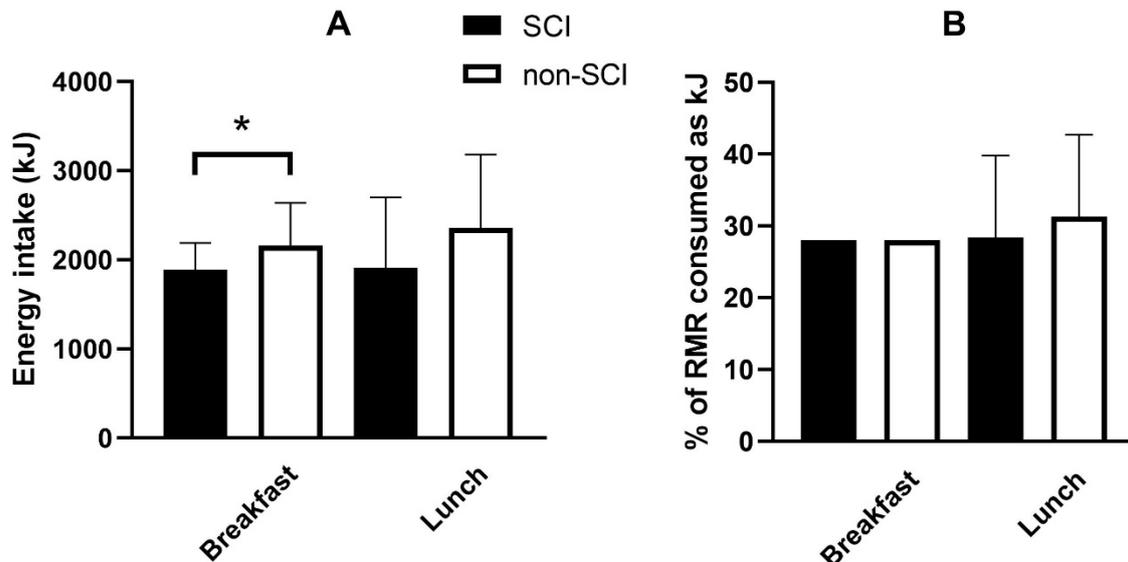
267 There were no differences between groups in pre-breakfast levels of hunger, satisfaction, fullness, or  
 268 prospective future consumption ( $p \geq 0.300$ ;  $d \leq 0.33$ ). Similarly, no group or group-by-time interaction  
 269 effects were apparent for any of the appetite perceptions from 0-4 h post breakfast consumption ( $p \geq$   
 270  $0.690$ ;  $\eta_p^2 \leq 0.018$ ) (Figure 2). In contrast, an effect of time was detected for all appetite perceptions in  
 271 the 0-4 h postprandial period (all  $p < 0.001$ ;  $\eta_p^2 \geq 0.692$ ). There was no group effect for how pleasant  
 272 participants found the breakfast ( $t(38) = -0.109$ ;  $p = 0.914$ ;  $d = 0.035$ ) or lunch ( $t(38) = -0.649$ ;  $p =$   
 273  $0.520$ ;  $d = 0.207$ ) meals. Similarly, no difference in how satisfying participants perceived breakfast to  
 274 be ( $t(38) = -1.526$ ;  $p = 0.135$ ;  $d = 0.579$ ) was identified. Conversely, the group with SCI deemed the  
 275 lunch meal to be less satisfying than participants in the non-SCI group ( $t(38) = -2.461$ ;  $p = 0.018$ ;  $d =$   
 276  $0.778$ ).



277  
 278 **Figure 2.** Perceptions of hunger (A, B) satisfaction (C, D), fullness, (E, F) and prospective future  
 279 consumption (G, H) in persons with spinal cord injury (SCI) and non-SCI individuals. Data are mean  $\pm$   
 280 SD ( $n = 20$  in each group). Vertical dotted lines were added for visual aid of the four time points.

281 3.4. Energy intake

282 Energy intake at the breakfast meal was lower in the SCI, compared to the non-SCI group ( $t(38) = -$   
 283  $2.188$ ;  $p = 0.035$ ;  $d = 0.69$ ) (Figure 3A). Trends toward a lower absolute energy intake in the group with  
 284 SCI were also seen at the *ad libitum* lunch meal ( $t(38) = -1.916$ ;  $p = 0.063$ ;  $d = 0.56$ ) (Figure 3A).  
 285 However, relative energy intake (energy intake scaled to RMR) ( $U = 166$ ;  $p = 0.358$ ;  $r = 0.15$ ) (Figure  
 286 3B) was not different between groups.



287  
 288 **Figure 3.** Breakfast and lunch energy intake (A), and energy intake as a percentage of RMR at the  
 289 breakfast and lunch meal (B). Data are mean  $\pm$  SD ( $n = 20$  in each group). \* Significant difference ( $p <$   
 290  $0.05$ ).

### 291 3.5. Trait and state eating behaviour questionnaires

#### 292 3.5.1. Adult Eating Behaviour Questionnaire

293 Satiety responsiveness was higher in the group with SCI, compared to the non-SCI group ( $t(38) = 2.266$ ;  
 294  $p = 0.029$ ;  $d = 0.72$ ; Table 3). However, no significant group differences were found for slowness in  
 295 eating ( $t(38) = 0.502$ ;  $p = 0.618$ ;  $d = 0.16$ ), emotional under eating ( $t(38) = 1.195$ ;  $p = 0.240$ ;  $d = 0.38$ ),  
 296 food fussiness ( $t(38) = -5.02$ ;  $p = 0.619$ ;  $d = 0.16$ ), food responsiveness ( $t(38) = 0.75$ ;  $p = 0.458$ ;  $d =$   
 297  $0.24$ ), emotional overeating ( $t(38) = 0.869$ ;  $p = 0.39$ ;  $d = 0.25$ ), hunger ( $t(38) = 0.234$ ;  $p = 0.816$ ;  $d =$   
 298  $0.07$ ), or enjoyment of food ( $U = 196.5$ ;  $p = 0.922$ ;  $r = 0.02$ ).

#### 299 3.5.2. Control of Eating Questionnaire

300 Craving for savoury foods was lower in the SCI group compared to the non-SCI group ( $t(38) = -2.975$ ;  
 301  $p = 0.005$ ;  $d = 0.94$ ; Table 3). However, no significant group differences for craving control ( $t(38) =$   
 302  $1.03$ ;  $p = 0.310$ ;  $d = 0.33$ ), positive mood ( $t(38) = 0.522$ ;  $p = 0.605$ ;  $d = 0.12$ ) or craving for sweet foods  
 303 ( $t(38) = -1.149$ ;  $p = 0.258$ ;  $d = 0.36$ ) were found.

304 *3.5.3. Reasons Individuals Stop Eating Questionnaire*

305 Scores of decreased food appeal ( $t(38) = 0.71$ ;  $p = 0.482$ ;  $d = 0.23$ ), physical satisfaction ( $t(38) = -0.049$ ;  
306  $p = 0.962$ ;  $d = 0.06$ ), planned amount ( $U = 197.5$ ;  $p = 0.946$ ;  $r = 0.01$ ), self-consciousness ( $U = 140$ ;  $p$   
307  $= 0.101$ ;  $r = 0.26$ ), and decreased priority of eating ( $U = 176$ ;  $p = 0.513$ ;  $r = 0.10$ ) were not significantly  
308 different between the group with SCI and the non-SCI group (Table 3).

309 *3.5.4. Three Factor Eating Questionnaire (R-18)*

310 The group with SCI reported higher levels of cognitive restraint compared to the non-SCI group ( $t(38)$   
311  $= 2.297$ ;  $p = 0.027$ ;  $d = 0.726$ ; Table 3), though no significant differences were detected when comparing  
312 uncontrolled eating ( $t(38) = -0.454$ ;  $p = 0.652$ ;  $d = 0.02$ ) or emotional eating ( $U = 155$ ;  $p = 0.216$ ;  $r =$   
313  $0.196$ ).

**Table 3**

Scores from different trait eating behaviour questionnaires in persons with and without a spinal cord injury (SCI).

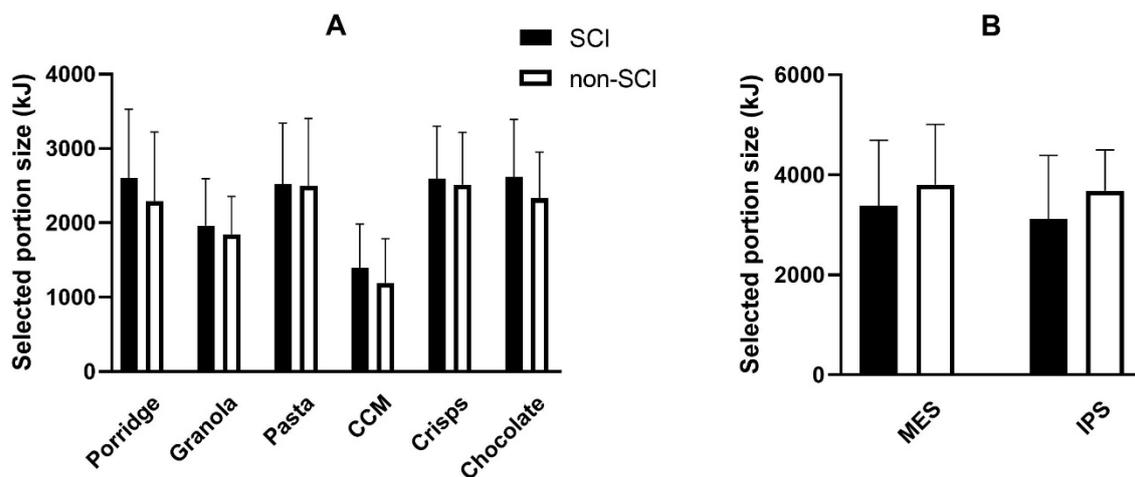
	SCI (n=20)	Non-SCI (n=20)
<b>Adult Eating Behaviour Questionnaire</b>		
<i>Food approach</i>		
Slowness in eating	9.8 ± 4.1	9.2 ± 3.4
Emotional under-eating	14.6 ± 4.8	12.9 ± 4.5
Food fussiness	6.0 ± 3.4	6.5 ± 2.8
Satiety responsiveness	10.3 ± 3.2 *	8.2 ± 2.4
<i>Food avoidance</i>		
Food responsiveness	13.1 ± 3.2	12.4 ± 2.7
Emotional overeating	11.9 ± 5.0	10.7 ± 4.6
Hunger	15.1 ± 4.3	14.8 ± 3.8
Enjoyment of food	14.0 (12.0-15.0)	14.0 (12.3-15.0)
<b>Control of Eating Questionnaire</b>		
Craving control	68.5 ± 21.5	61.7 ± 20.0
Positive mood	70.8 ± 18.8	68.6 ± 16.8
Craving for savoury	34.4 ± 14.3 *	49.1 ± 16.8
Craving for sweet	33.3 ± 20.2	40.9 ± 21.6
<b>Reasons Individuals Stop Eating Questionnaire</b>		
Decreased food appeal	2.4 ± 0.8	2.2 ± 0.8
Physical satisfaction	4.4 ± 1.0	4.4 ± 1.3
Planned amount	4.3 (4.0-5.0)	4.4 (3.7-5.2)
Self-consciousness	1.9 (1.4-3.0)	1.4 (1.0-2.3)
Decreased priority of eating	2.0 (1.8-2.8)	2.0 (1.1-2.8)
<b>Three Factor Eating Questionnaire - Revised 18-item version</b>		
Uncontrolled eating	37.8 ± 20.3	37.4 ± 12.6
Cognitive restraint	49.1 ± 16.9 *	36.8 ± 16.9
Emotional eating	33.3 (13.9-41.7)	22.2 (2.8-33.3)

Mean ± SD; Median (IQR).

\* Significant difference between groups (p < 0.05).

315 3.6. *Expected satiety and Ideal portion size*

316 For measures of hypothetical expected satiety, there was no effect of group ( $F_{(1,38)} = 1.274$ ;  $p = 0.266$ ;  
317  $\eta_p^2 = 0.032$ ), and no interaction effect ( $F_{(1,38)} = 0.346$ ;  $p = 0.884$ ;  $\eta_p^2 = 0.009$ ). However, a main effect  
318 of food was present ( $F_{(1,38)} = 26.894$ ;  $p < 0.001$ ;  $\eta_p^2 = 0.414$ ) (Figure 4A). In contrast, no main effect of  
319 group ( $F_{(1,38)} = 2.116$ ;  $p = 0.154$ ;  $\eta_p^2 = 0.054$ ), food ( $F_{(1,38)} = 1.468$ ;  $p = 0.233$ ;  $\eta_p^2 = 0.038$ ), or interaction  
320 effect ( $F_{(1,38)} = 0.193$ ;  $p = 0.663$ ;  $\eta_p^2 = 0.005$ ) was found for ideal portion size and momentary expected  
321 satiety (Figure 4B).



322  
323 **Figure 4.** Hypothetical expected satiety assessments for six test foods (A), and momentary expected  
324 satiety (MES) and ideal portion size (IPS) for cheese and tomato pasta (B) in persons with a spinal cord  
325 injury (SCI) and non-SCI individuals. Data are mean  $\pm$  SD ( $n = 20$  in each group).

326 **4. Discussion**

327 In a controlled experimental setting, this study investigated whether differences in appetite, food reward  
328 and eating behaviour traits, compared to non-SCI individuals, may contribute to the greater obesity  
329 prevalence in those with SCI. We found that meal-related perceptions of appetite were no different  
330 between groups but people with SCI exhibited stronger liking and motivation to consume sweet rather  
331 than savoury foods. We also found that people with SCI exhibit higher levels of dietary cognitive  
332 restraint compared with the non-SCI group. These findings suggest that SCI is associated with altered  
333 hedonic factors impacting appetite and eating behaviour.

334 *4.1. Food reward*

335 We found that sweet appeal bias scores for ‘relative preference’, ‘explicit wanting’, and ‘explicit liking’  
336 were greater in persons with SCI compared to those without SCI, in both hungry and fed states. Lower  
337 cravings for savoury foods in the group with SCI, as reported in the Control of Eating Questionnaire,  
338 reinforced this finding. These findings indicate that people with SCI exhibit a greater overt preference

339 and desire to consume sweet rather than savoury foods. In the context of appetite, eating behaviour and  
340 energy balance, this finding is relevant as sweet foods are typically more energy dense, contain a high  
341 proportion of simple sugars, and have a weak effect on satiety (Hogenkamp, 2016). It is possible that  
342 this pleasure-related difference in appetite may contribute to a greater propensity for overconsumption,  
343 however, the cross-sectional nature of this study does not allow causal influence to be determined.

344 This finding is consistent with previous data which found that inactive men experienced a greater liking  
345 for sweet foods compared with active men (Horner et al., 2016). Notably, whilst data from our physical  
346 activity questionnaires indicated no clear difference in activity behaviours between study groups, it  
347 should be recognised that these questionnaires lack sensitivity, and that the group with SCI almost  
348 certainly expend less energy in daily activities each day. The relevance of physical inactivity is also  
349 supported by studies measuring food reward via fMRI. Herein, individuals displaying low levels of  
350 daily physical activity experienced greater activation of reward related brain regions in response to  
351 high-energy foods (Drummen et al., 2019; Killgore et al., 2013; Luo et al., 2018). Collectively, such  
352 differences in food hedonics may contribute to the established uncoupling between energy intake and  
353 energy expenditure in people who habitually demonstrate low levels of activity-related energy  
354 expenditure (Hopkins & Blundell, 2016).

355 We speculate that the higher levels of adiposity, found in persons with a SCI may have contributed to  
356 their greater preference and motivation for sweet foods (Dressler & Smith, 2013; Lampuré et al., 2016;  
357 Nakamura et al., 2001). As expected, although we matched groups for BMI, those with SCI were found  
358 to have 20% greater skinfolds and displayed significantly greater waist circumferences than the non-  
359 SCI group. These results are in-line with other literature in SCI (Spungen et al., 2003) and studies have  
360 shown obesity to be associated with impaired signalling systems, such as insulin and leptin resistance,  
361 and oxidative and endoplasmic reticulum stress (Berthoud et al., 2011). This creates a central toxic  
362 environment which leads to neural alterations in parts involved in reward processing, such as the  
363 corticolimbic system. These changes have been shown to accentuate the progression of obesity,  
364 insinuating that elevated adiposity increases food reward circuitry activity (Berthoud et al., 2011).

#### 365 *4.2. Cognitive restraint*

366 A second notable finding was that cognitive restraint was higher in the group with SCI, indicating that  
367 a greater effort to limit day-to-day food intake is made in this group (Stunkard & Messick, 1985).  
368 Elevated levels of restraint may reflect the individuals' increased awareness of their lowered energy  
369 requirements and how, particularly in today's obesogenic environment, more psychological effort is  
370 required to prevent overconsumption and weight gain. Similarly, elevated body image concerns in  
371 persons with SCI, specifically around weight gain, may lead to changes in the quantity and type of foods  
372 the individual perceives acceptable to eat (Alysse Bailey et al., 2016).

373 It has been proposed that the elevated cognitive effort used to resist hedonic food cues for high-energy  
374 foods may result in increased wanting and craving (Adams et al., 2019; Veenstra & de Jong, 2010;  
375 Verzijl et al., 2018). This could infer that by exhibiting greater cognitive restraint against sweet foods,  
376 persons with SCI inadvertently develop an increase in food reward. However, the relationship between  
377 cognitive restraint, food reward, and energy intake, specifically of sweet or energy-dense foods, has not  
378 previously been explored in an SCI population. Further investigation into this relationship may provide  
379 more insight into how these factors influence weight gain and obesity in persons with SCI.

#### 380 4.3. Satiety responsiveness and energy intake

381 Our research group have previously noted that postprandial satiety responses were greater in persons  
382 with SCI compared to a non-SCI group following an isoenergetic preload (Fenton et al., 2021). In our  
383 previous work we proposed that the preload represented a relatively larger meal for the group with SCI  
384 compared to the non-SCI group due to their lower energy requirements. This current study is the first  
385 to extend this hypothesis across a 4 h postprandial period following a breakfast meal with an energy  
386 content normalised to participants' RMR and found that appetite perceptions, including satiety, were  
387 not different in persons with SCI. Additionally, after the 4 h period, *ad libitum* energy intake, normalised  
388 to RMR, was similar in SCI and non-SCI groups. These data suggest that meal-related satiety  
389 perceptions are not 'dysregulated' as a result of SCI. Data from our hypothetical expected satiety  
390 questionnaire also support this idea, showing that persons with SCI do not perceive there to be any  
391 differences in the satiating abilities of a variety of foods compared to the non-SCI group.

392 A trend toward a lower absolute *ad libitum* energy intake was also seen in the group with SCI. In  
393 conjunction with the aforementioned findings, this could suggest that satiety perceptions in persons  
394 with SCI are upregulated to adjust for reduced energy demands. This notion is supported by the elevated  
395 levels of satiety responsiveness and cognitive restraint seen in the group with SCI, inferring that there  
396 could be both physiological and psychological factors influencing this adjustment. Whilst scores of  
397 momentary expected satiety and ideal portion size were not significantly different between groups, the  
398 non-SCI group recorded portion sizes that were 419 kJ and 558 kJ higher than the group with SCI,  
399 respectively, which further supports this concept. Our absolute energy intake data match this, suggesting  
400 that this study may have been underpowered to detect group differences. Altogether, alterations in meal-  
401 related and trait-based satiety profiles do not appear to underpin an enhanced obesity predisposition in  
402 people with SCI.

#### 403 4.4. Strengths and limitations

404 The primary strength of this study was the recruitment of a hard-to-reach population that was matched  
405 to a control group for age, sex, and BMI. However, the heterogenous nature of the group with SCI with  
406 regards to injury level and motor function must be considered with regards to variability in study  
407 outcomes. In addition, given the exploratory nature of this study, multiple outcomes were examined

408 increasing the potential for Type I errors. Another potential limitation of this study was conducting an  
409 LFPQ after the *ad libitum* lunch meal, as the quantity of food was not standardised or normalised. Thus,  
410 the validity of this LFPQ may be questionable. Subsequent studies may benefit from conducting  
411 postprandial LFPQs following a meal normalised for energy requirements. Additionally, although  
412 skinfold thickness was (almost;  $p = 0.059$ ) greater in the group with SCI, suggesting elevated levels of  
413 adiposity, no objective measures of body composition were conducted in the current study. The  
414 influence of accentuated adiposity as a rationale for group differences in food reward or eating  
415 behaviours should therefore be interpreted with caution. The use of questionnaires to assess habitual  
416 physical activity is another limitation of the current study, given the low reliability of self-reported  
417 physical activity status (Prince et al., 2008). Although, objective measures of physical activity status,  
418 such as wrist-worn accelerometers, have been validated in SCI populations against non-SCI controls  
419 (Murphy et al., 2019), these measures would not provide an accurate reflection of the interindividual  
420 differences in energy expenditure, which is an important consideration when assessing appetite- or  
421 energy intake-related measures.

#### 422 *4.5 Conclusions*

423 In summary, food reward for sweet foods, as well as cognitive restraint, was greater in persons with  
424 SCI compared to the non-SCI group. In addition, postprandial appetite perceptions and subsequent *ad*  
425 *libitum* energy intake following a breakfast meal standardised for energy requirements, were not  
426 different between the SCI and non-SCI groups. Collectively, these findings highlight that altered satiety  
427 profiles do not appear to increase susceptibility to obesity in this population. Conversely, elevated food  
428 reward for sweet food in persons with SCI may lead to overconsumption.

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434 **Author contributions**

435 JMF, SPH, CJM, VLT and JAK conceived the study idea. JMF performed data collection, conducted  
436 the data analysis, and led the writing of the manuscript with the input of all authors. All authors reviewed  
437 and approved the final version of the manuscript.

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441 **Ethical approval**

442 All participants gave informed consent before taking part in this study. Ethical approval was given by  
443 Loughborough University Ethics Review (Human Participants) Sub-Committee (Study ID: 5574)

444 **Declaration of competing interest**

445 None.

446 **Ethical statement**

447 This research obtained ethical approval from Loughborough University's ethics review board before  
448 any study related procedures commenced (Study ID: 5574). All participants provided written informed  
449 consent to participate.

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