

1 Two inhibitory control training interventions designed to improve eating behaviour and  
2 determine mechanisms of change

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**Highlights**

- 14 • Testing the effect of Stop-Signal training on eating behaviour and self-regulation
- 15 • Training did not change eating behaviour outside the laboratory
- 16 • Improvements in resistance to depletion and inhibitory contro
- 17 • I were not maintained
- 18 • This particular training may not be intense enough to influence eating behaviour
- 19 • Improvements in self-regulation may only persist insofar as training does

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## Abstract

Inhibitory control training has been shown to influence eating behaviour in the laboratory; however, the reliability of these effects is not yet established outside the laboratory, nor are the mechanisms responsible for change in behaviour. Two online Stop-Signal Task training interventions were conducted to address these points. In Study 1, 72 participants completed baseline and follow-up measures of inhibitory control, self-regulatory depletion, fat intake and body-mass index. Participants were randomly assigned to complete one of three Stop-Signal Tasks daily for ten days: *food-specific inhibition*- inhibition in response to unhealthy food stimuli only, *general inhibition*- inhibition was not contingent on type of stimuli, and *control*- no inhibition. While fat intake did not decrease, body-mass index decreased in the food-specific condition and change in this outcome was mediated by changes in vulnerability to depletion. In Study 2, the reliability and longevity of these effects were tested by replicating the intervention with a third measurement time-point. Seventy participants completed baseline, post-intervention and follow-up measures. While inhibitory control and vulnerability to depletion improved in both training conditions post-intervention, eating behaviour and body-mass index did not. Further, improvements in self-regulatory outcomes were not maintained at follow-up. It appears that while the training paradigm employed in the current studies may improve self-regulatory outcomes, it may not necessarily improve health outcomes. It is suggested that this may be due to the task parameters, and that a training paradigm that utilises a higher proportion of stop-signals may be necessary to change behaviour. In addition, improvements in self-regulation do not appear to persist over time. These findings further current conceptualisations of the nature of self-regulation and have implications for the efficacy of online interventions designed to improve eating behaviour.

44 Two inhibitory control training interventions designed to improve eating behaviour and  
45 determine mechanisms of change

46 The prevalence of overweight and obesity is increasing (Colagiuri et al., 2010; Flegal,  
47 Carroll, Ogden, & Curtin, 2010). Although the current food-rich environment, in which  
48 unhealthy choices are readily available, may make achieving and maintaining the goal of  
49 healthy eating difficult (Stroebe, 2008; Wansink, 2004), some individuals are able to resist  
50 high calorie foods and maintain a healthy diet and weight. Research suggests that inhibitory  
51 control may be one important factor implicated in the regulation of eating behaviour  
52 (Hofmann, Friese, & Roefs, 2009; Houben & Wiers, 2009).

53 Inhibitory control refers to the ability to overrule impulsive reactions in order to  
54 regulate behaviour in line with long-term goals (Miyake et al., 2000). In the case of eating  
55 behaviour, this may involve resisting the impulse to eat high-calorie food in order to meet the  
56 goal of adhering to a healthy diet. Individual differences in measures said to assess inhibitory  
57 control such as the Go/No-Go Task (GNG; Miller, Schäffer, & Hackley, 1991) and the Stop-  
58 Signal Task (SST; Logan, Schachar, & Tannock, 1997) consistently predict eating behaviours  
59 (Allom & Mullan, 2014; Hall, 2012; Hofmann et al., 2009), as well as weight gain  
60 (Nederkoorn, Houben, Hofmann, Roefs, & Jansen, 2010), among non-clinical participants.  
61 Further, inhibitory control can be undermined leading to greater consumption of high calorie  
62 foods (Hofmann, Rauch, & Gawronski, 2007; Vohs & Heatherton, 2000). This effect, termed  
63 depletion, derives from the strength model of self-regulation (Baumeister, Vohs, & Tice,  
64 2007), in which self-regulation is assumed to rely on a limited resource. Goal directed  
65 behaviours are rarely performed in isolation, or without the influence of external stressors-  
66 two factors which lead to depletion and compromise the capacity to enact goal directed  
67 behaviour (Hagger, Wood, Stiff, & Chatzisarantis, 2009). Therefore, in order to achieve the  
68 goal of healthy eating, both inhibitory control and resistance to depletion are necessary.

69 Current research suggests that inhibitory control training can influence eating  
70 behaviour using both GNG and SST paradigms (Lawrence, Verbruggen, Morrison, Adams, &  
71 Chambers, 2015; Veling, van Koningsbruggen, Aarts, & Stroebe, 2014). In GNG training  
72 paradigms, participants are required to respond as rapidly as possible to a neutral set of  
73 stimuli while always withholding responses to a set of stimuli representing the target  
74 behaviour (Veling, Aarts, & Papies, 2011; Veling, Aarts, & Stroebe, 2013). Consistent  
75 pairings of the no-go response with target stimuli facilitates the retrieval of no-go-target  
76 stimuli associations and results in improved inhibition of responses to target stimuli (Spierer,  
77 Chavan, & Manuel, 2013). SST training paradigms differ from GNG as participants are  
78 instructed to respond as rapidly as possible to both target stimuli and neutral stimuli and only  
79 inhibit responses to target stimuli on a proportion of trials (Jones & Field, 2013; Lawrence et  
80 al., 2015). Improvement in behaviour is typically assessed using a between-participants  
81 design wherein participants who are randomly assigned to receive inhibitory control training  
82 consume or select less unhealthy foods in an immediately administered laboratory-based task,  
83 compared to those assigned to an inert or alternative form of training (Houben, 2011; Veling  
84 et al., 2011).

85 To date, only one study has assessed *change* in ecologically valid health outcomes as  
86 a result of inhibitory control training (Veling et al., 2014). This study demonstrated that four  
87 sessions of GNG training resulted in decreased BMI. However, underlying mechanisms  
88 responsible for change in health outcomes were not directly tested. As described above, the  
89 two training paradigms differ in that in the GNG, the go response is consistently inhibited for  
90 all members of a certain category, while in the SST the ‘go’ response does not need to be  
91 inhibited for all members of a certain category, only for a certain proportion. Therefore, it is  
92 suggested that the effectiveness of these paradigms may differ, and the mechanisms by which  
93 they influence health behaviour may also differ. Preliminary evidence suggests that GNG

94 training results in the devaluation of unhealthy food stimuli and that this is responsible for  
 95 differences in eating behaviour (van Koningsbruggen, Veling, Stroebe, & Aarts, 2013). While  
 96 no direct evidence exists as to what mechanism of change underlies SST training, Jones and  
 97 Field (2013) demonstrated that alcohol-specific SST training led to a reduction in inhibition  
 98 errors to alcohol stimuli across training blocks, which may suggest that SST training  
 99 improves health behaviour by increasing inhibitory control. Nevertheless, this assumption  
 100 was not directly tested as no additional measure of inhibitory control was included, thus this  
 101 result may have been due to a practice effect. Therefore, not only is there is a need to  
 102 examine whether SST training produces changes in ecologically valid eating behaviour  
 103 outcomes, but to also examine the mechanisms that underlie the effect of training.

104         It is proposed that SST training may not only influence eating behaviour by  
 105 improving inhibitory control, but also by decreasing vulnerability to depletion. Vulnerability  
 106 to depletion has been shown to decrease after behaviour regulation training (Muraven, 2010),  
 107 which involves regulating an element of behaviour that is unrelated to the target behaviour,  
 108 such as speech, posture, or mood, for a period time in order to improve self-regulation and  
 109 consequently health behaviour (Muraven, Baumeister, & Tice, 1999; Oaten & Cheng,  
 110 2006b). For example, Oaten and Cheng (2006a) demonstrated that reductions in depletion  
 111 effects after training resulted in improvement in a variety of self-reported health behaviours,  
 112 including improvements in healthy eating. Therefore, it may be worthwhile to examine  
 113 whether inhibitory control training not only improves inhibitory control capacity but also  
 114 decreases vulnerability to depletion, and to examine whether changes in these elements of  
 115 self-regulation account for changes in eating behaviour.

116                                   **Present research**

117         Therefore, the aim of the present research was to improve self-reported eating  
 118 behaviour through online SST training and to test two potential mechanisms by which this

119 particular version of SST training may improve health behaviour, by examining the extent to  
120 which training effects can be attributed to improvements in inhibitory control and/or a  
121 decreased vulnerability to depletion. In order to achieve these aims a SST with 25% stop-  
122 signal trials was employed, and three conditions, each with a different version of the SST,  
123 were included: (1) *food-specific inhibition* condition in which the stop-signals were paired  
124 only with unhealthy food stimuli, (2) *general inhibition* condition in which the same stimuli  
125 and proportion of stop-signals were used; however, the stop-signals were not contingent on a  
126 particular category of stimuli, and (3) *control* condition that included the same stimuli as  
127 other conditions but without stop-signals. This final condition was included in order to  
128 determine whether general inhibition training was sufficient enough to change behaviour. The  
129 stop-signal density was kept at 25% of trials in order to ensure that the training was  
130 influencing inhibitory control, or the ability to cancel a response, rather than devaluating the  
131 stimuli associated with the stop response, as is proposed to be the case with GNG training in  
132 which a stop response is always paired with the target stimuli (Schachar et al., 2007).

133         It was hypothesised that inhibitory control and vulnerability to depletion would  
134 improve in both training conditions compared to the control; however, greater improvement  
135 in eating behaviour was expected in the food-specific inhibition condition as inhibition  
136 training was targeted to this behaviour. Finally, it was expected that changes in inhibitory  
137 control and changes in vulnerability to depletion would mediate the effect of food-specific  
138 inhibition training on changes in eating behaviour. Study 1 reports a preliminary investigation  
139 into the effect of training on health and self-regulatory outcomes, while Study 2 reports a  
140 replication of the training intervention with an additional measurement point in order to test  
141 the reliability and longevity of any training effects observed in Study 1.

142 **Study 1**

143 **Method**

144 **Participants**

145 Eighty-two undergraduate students from a variety of disciplines (age = 20.43 years,  $SD =$   
146 4.86; BMI = 22.62,  $SD = 2.64$ ; 66 females) were recruited to participate in a study in  
147 exchange for course credit. The number of participants recruited was based on an a-prior  
148 power analysis using G-Power 3 software (Faul, Erdfelder, Lang, & Buchner, 2007), which  
149 indicated that a sample size of 69 would be sufficient to detect a small to medium (0.15)  
150 interaction effect between three conditions at two time points with a power of .80 and an  
151 alpha of .05.

152 Inclusion criteria included having the intention to change dietary behaviour, not  
153 colour blind, fluent in English, and having access to the internet. Additionally, participants  
154 were excluded if they indicated that they had a current or prior eating disorder diagnosis.  
155 Participants were randomly allocated to one of three conditions: *food-specific inhibition* ( $n =$   
156 29), *general inhibition* ( $n = 25$ ), and *control* ( $n = 28$ ) by clicking a URL, which randomly  
157 directed them to one of three pages. The university's human research ethics committee  
158 approved the study and participants provided informed consent prior to participation.

159 **Materials and measures**

160 **BMI & saturated fat intake.** BMI was calculated from participants' self-reported  
161 height and weight. Saturated fat intake in grams was calculated from responses on the Block  
162 food screener (Block, Gillespie, Rosenbaum, & Jenson, 2000), which has been validated  
163 against a 100-item food frequency questionnaire (Block et al., 2000). Participants indicated  
164 how often they ate 17 meat and snack items (e.g. bacon, full-fat ice-cream, fried potatoes) on  
165 a 5 point scale ranging from: never (0), to 5 or more times per week (4).



166           **Stroop interference task.** Change in inhibitory control capacity was assessed using  
167 the computerised version of the Stroop, in which participants were required to name the  
168 colour in which a written colour word is printed while inhibiting the tendency to read the  
169 word itself. For example, when the word ‘red’ is printed in blue, the tendency to respond  
170 ‘red’ must be inhibited in order to provide the correct response of ‘blue’. The task consisted  
171 of three types of trials presented in three experimental blocks of 60 trials each and one  
172 practice block of 20 trials. *Congruent trials* consisted of colour words that were printed in the  
173 corresponding colour. In *incongruent trials*, the colour of the colour word was different to the  
174 word itself. *Control trials* consisted of strings of letters matched in length to the colour  
175 words. Stimuli were displayed until the participant responded, and the response-stimulus  
176 interval was 500ms. The Stroop interference score was calculated as the difference between  
177 mean response time of correct responses on incongruent trials and control trials (MacLeod,  
178 2005), where a larger score indicated poorer inhibitory control. Response times that fell three  
179 standard deviations above or below a participant’s mean reaction time per block were deemed  
180 to be outliers and were deleted (MacLeod, 2005).

181           **Depletion task.** Participants were asked to write about what they had done over the  
182 weekend for five minutes with the instructions not to use two common letters, namely, a or n.  
183 This task has been used in previous research to induce depletion (Lewandowski, Ciarocco,  
184 Pettenato, & Stephan, 2012; Schmeichel, 2007). Participants also completed a four item  
185 questionnaire measuring their perceptions regarding the depletion task (Muraven &  
186 Slessareva, 2003), including how difficult and unpleasant (1 = extremely easy/pleasant – 7 =  
187 extremely difficult/unpleasant), and frustrating (1 = not at all frustrating – 5 = extremely  
188 frustrating), the depletion task had been for them. In addition, participants indicated how  
189 much effort the task required: “How much were you fighting against an urge while working  
190 on the task?” (1 = not at all – 5 = extremely), and written responses were reviewed to ensure

191 that participants had completed the task correctly. Depletion was calculated as the difference  
192 between Stroop interference before and after the depletion task, where a larger score  
193 indicated greater vulnerability to depletion.

194       **Stop-signal task.** The current study utilised three versions of the SST with cues,  
195 which included three experimental blocks of 64 trials and a practice block of 32 trials. In all  
196 versions, each trial began with a fixation cross (+) presented in the centre of the screen for  
197 500ms, followed by a picture of either an unhealthy food or a healthy food. All conditions  
198 were exposed to the same number of unhealthy and healthy food stimuli (50% unhealthy,  
199 50% healthy). Participants in all conditions were required to categorise the content of the  
200 picture by pressing the “D” key for an unhealthy food picture or the “K” key for a healthy  
201 food picture, which was counterbalanced across participants. For the two training conditions,  
202 on 25% of trials an auditory tone occurred after a delay which signified that participants  
203 should inhibit their response on that trial and wait for the next trial. The stop-signal delay  
204 (SSD) was initially set at 250ms and was adjusted dynamically according to participants’  
205 responses using a staircase tracking procedure: When inhibition was successful, SSD  
206 increased by 50ms; when inhibition was unsuccessful, SSD decreased by 50ms. On stop-  
207 signal trials, responses within the 1500ms timeout period were classed as inhibition errors  
208 (Verbruggen, Logan, & Stevens, 2008).

209       For the *food-specific inhibition* condition, the stop-signal was only presented after  
210 unhealthy food images. Therefore, each block consisted of 16 unhealthy food-stop trials, 16  
211 unhealthy food go-trials, 0 healthy food-stop trials and 32 healthy food-go trials. For the  
212 *general inhibition* condition, the stop-signal was randomly presented either after a healthy or  
213 an unhealthy food image. Therefore, each block consisted of 8 unhealthy food-stop trials, 24  
214 unhealthy food-go trials, 8 healthy food-stop trials and 24 healthy food-go trials. For the  
215 *control* condition, participants performed the same task as the other conditions; however, no

216 stop-signals were presented. If participants in either training condition inhibited their  
217 responses less than 50% of the time on inhibition trials this was an indication that they were  
218 not responding to the stop-signal correctly and thus that session was not included as a training  
219 session. Similarly, if participants inhibited their responses more than 50% of the time, this  
220 was not counted as a training session and was excluded (Verbruggen et al., 2008).

221 Stimuli consisted of eight colour pictures of both sweet and savoury unhealthy foods  
222 (e.g., potato chips, chocolate) and eight colour pictures of fruit and vegetables (e.g., apple,  
223 carrot) displayed on a white background and were approximately 450 by 400 pixels in size.  
224 The stimuli were comparable to those used in previous research on eating behaviour and  
225 impulsive responses (Veling et al., 2013), and those represented in the Block food screener.

## 226 **Procedure**

227 The study was conducted entirely online over 12 days. Once participants had signed  
228 up to the study, and provided informed consent, they completed the pre-intervention  
229 measures in the following order: Stroop task, depletion task, Stroop task, the Block food  
230 screener, and reported their height and weight. Finally, participants completed demographic  
231 measures and the questionnaire measuring their perceptions of the depletion task. On Days 2  
232 – 11, participants completed one of three SST, depending upon the condition to which they  
233 had been randomly assigned. Finally, on Day 12 participants completed the same measures as  
234 Day 1, with the exception of height, and demographic measures.

## 235 **Data analyses**

236 In order to confirm that randomisation was successful the three experimental  
237 conditions were compared with respect to scores on age, BMI, Stroop interference,  
238 vulnerability to depletion, and saturated fat intake using a one-way analysis of variance  
239 (ANOVA), while a chi-squared analysis was utilised to assess sex differences between  
240 conditions. Similarly, one-way ANOVAs were used to determine differences on all variables,

241 including condition, between those who completed the study and those who dropped out,  
242 with the exception of sex where a Fisher's Exact Test was used. To ensure that the depletion  
243 task influenced participants' self-regulatory resources, pre-intervention Stroop interference  
244 scores were compared pre- to post- depletion across all conditions using a paired samples *t*-  
245 test. To assess the effect of training on Stroop performance and vulnerability to depletion two  
246 2(time: pre-intervention; Day 1, post-intervention; Day 12) by 3(condition: food-specific  
247 inhibition, general inhibition, control) mixed ANOVAs were conducted. If a significant time  
248 by condition interaction was detected, planned contrasts examining whether change in self-  
249 regulatory outcomes experienced by the training conditions differed from that experienced by  
250 the control, as well as whether the two training conditions differed from each other.  
251 Similarly, to assess the effect of training on saturated fat intake, a 2 x 3 mixed ANOVA was  
252 conducted; with planned contrasts examining whether change experienced by the food-  
253 specific condition differed to that experienced by the general inhibition and control  
254 conditions, as well as whether the two training conditions differed from each other. Finally,  
255 bootstrapping techniques for simple mediation (Hayes, 2012), were utilised to test whether  
256 changes in either inhibitory control or vulnerability to depletion mediated the effect of food-  
257 specific training related changes in saturated fat intake.

## 258 **Results**

### 259 **Randomisation check**

260 There were no significant differences in any tested variables between conditions, all *p*  
261 > .05. Additionally, the number of SSTs performed did not differ between conditions, *p* > .05.

### 262 **Attrition**

263 Ten participants did not complete post-intervention measures (food-specific  
264 inhibition: *n* = 3, general inhibition: *n* = 4, control: *n* = 3). Three participants dropped out of

265 the study and seven did not sufficiently engage with all tasks. There were no differences  
266 between those who completed the study and those who did not on any tested variables all,  $p >$   
267  $.05$ .

## 268 **Depletion**

269 Participants' performance on the Stroop task was significantly poorer following the  
270 depletion task,  $MD = -107.870$ ,  $SE = 8.531$ ;  $t(81) = -12.644$ ,  $p < .001$ . Additionally, on  
271 average participants reported the task as difficult,  $M = 6.27$ ,  $SD = 0.92$ , unpleasant,  $M = 5.12$ ,  
272  $SD = 1.29$ , frustrating,  $M = 3.61$ ,  $SD = 1.24$ , and effortful,  $M = 3.35$ ,  $SD = 1.07$ .

## 273 **Training effects**

274 **Inhibitory control.** There was a significant main effect of time indicating that all  
275 conditions improved on Stroop performance post-intervention,  $F(1, 69) = 4.635$ ,  $p = .035$ ,  
276 partial  $\eta^2 = .063$ . There was no main effect of condition, nor was the time by condition  
277 interaction effect significant, all  $p > .05$ . See Table 1 for pre- and post- intervention means  
278 and standard deviation of all test variables.

279 **INSERT TABLE 1 NEAR HERE**

280 **Vulnerability to depletion.** A comparison of pre- and post- intervention depletion  
281 scores revealed a significant main effect of time such that all conditions were less vulnerable  
282 to depletion post-intervention,  $F(1, 69) = 15.097$ ,  $p < .001$ , partial  $\eta^2 = .180$ , which was  
283 qualified by a significant time by condition interaction effect,  $F(2, 69) = 3.781$ ,  $p = .028$ ,  
284 partial  $\eta^2 = .099$ ; see Figure 1. A planned contrast examining the significant interaction  
285 revealed that both training conditions experienced improvement in vulnerability to depletion,  
286 compared to the control condition,  $\psi = 55.146$ ,  $F(1,69) = 6.377$ ,  $p = .014$ . Further,  
287 improvement in the food-specific inhibition condition did not differ significantly from the

288 general inhibition condition,  $\psi = 23.953$ ,  $F(1,69) = .8599$ ,  $p = .357$ . There was no main effect  
289 of condition on depletion,  $p > .05$ .

290 INSERT FIGURE 1 NEAR HERE

291 **Saturated fat intake.** There was no main effect of condition, time, nor was the time  
292 by condition interaction effect significant, all  $p > .05$ .

293 **BMI.** There was a significant main effect of time on BMI such that all conditions  
294 decreased in BMI post-depletion,  $F(1, 69) = 10.048$ ,  $p = .002$ , partial  $\eta^2 = .127$ , which was  
295 qualified by a significant time by condition interaction effect,  $F(2, 69) = 5.086$ ,  $p = .009$ ,  
296 partial  $\eta^2 = .128$ , see Figure 2. A planned contrast examining the significant interaction  
297 revealed that BMI decreased in the food-specific inhibition condition post-intervention, while  
298 BMI did not change in the general inhibition condition and the control,  $\psi = .354$ ,  $F(1,69) =$   
299  $10.171$ ,  $p = .002$ . Additionally, a contrast comparing change in BMI in the food-specific  
300 inhibition condition to the general inhibition condition revealed that BMI decreased more in  
301 the food-specific inhibition condition compared to the general inhibition condition,  $\psi = .365$ ,  
302  $F(1,69) = 7.53$ ,  $p = .008$ . There was no main effect of condition,  $p > .05$ .

303 INSERT FIGURE 2 NEAR HERE

304 **Mediation analysis.** As there were no changes in saturated fat intake the original  
305 mediation analysis was not conducted. However, the indirect effect of food-specific  
306 inhibition training on BMI through vulnerability to depletion was tested. In order to conduct  
307 this analysis, the general inhibition condition was grouped with the control condition and  
308 compared to the food-specific inhibition condition. Change in vulnerability to depletion and  
309 change in BMI variables were created by subtracting post-intervention scores from pre-  
310 intervention scores. The significance of the indirect effect was assessed using 95%  
311 confidence intervals, calculated using 5000 bootstrap re-samples (Hayes, 2012). The indirect  
312 effect from food-specific training, through change in vulnerability to depletion, to change in

313 BMI was significant,  $\beta = 0.071$ , 95% [CI: 0.01, 0.20]. The  $R^2$  mediation effect size was  
314 .0527;  $SE = .0386$ , indicating that 5.27% of the variance in change in BMI was explained by  
315 the mediating effect of change in vulnerability to depletion on the type of training effect, see  
316 Figure 3 for standardised coefficients between all variables.

317 INSERT FIGURE 3 NEAR HERE

## 318 Discussion

319 As expected, both training conditions demonstrated a decrease in vulnerability to  
320 depletion, and within the food-specific training condition; changes in vulnerability to  
321 depletion mediated changes in BMI. However, food-specific training did not result in changes  
322 in saturated fat intake, nor did type of training influence inhibitory control.

323 It is possible that training did not differentially influence inhibitory control capacity as  
324 Stroop interference is not reflecting the same specific inhibitory control mechanism that SST  
325 training is influencing. However, given that previous research has shown there to be overlap  
326 between the two tasks (Allom & Mullan, 2014; Miyake et al., 2000; Verbruggen, Liefoghe,  
327 & Vandierendonck, 2004), it is unlikely that these measures are wholly independent. While  
328 the Stroop procedure used in the current study has been frequently used in previous research  
329 (Cassiday, McNally, & Zeitlin, 1992; Formea & Burns, 1996; McNally, Riemann, & Kim,  
330 1990), it may be that not enough practice trials were used. A sufficient number of practice  
331 trials is essential in order to acclimatise participants to the display and response  
332 characteristics of the task so that response times are based on interference rather than the  
333 novelty of the task (MacLeod, 2005).

334 Despite this, the present results indicated a significant change in vulnerability to  
335 depletion in the training conditions. These results are similar to Muraven et al. (1999), who  
336 found that behavioural regulation training results in reduced depletion. Similarly, Oaten and  
337 Cheng (2007) found that after four months of engaging in financial monitoring participants

338 were not only less vulnerable to depletion but also reported engaging in more health  
339 enhancing behaviours. In contrast, within the current study this improvement only transferred  
340 to change in BMI in the food-specific condition, suggesting that behavioural specificity of the  
341 task, coupled with decrease in vulnerability to depletion may be necessary to change  
342 behaviour. Alternatively, it may be that more intense training is required for improvements to  
343 translate across behavioural domains. Further research is required to determine the optimal  
344 intensity and length of training required to achieve such transfer effects.

345 SST training did not appear to alter self-reported eating behaviour. Previous research  
346 using the SST to influence eating behaviour has demonstrated differences between training  
347 and control conditions in the amount consumed in a taste test (Lawrence et al., 2015). Future  
348 research should compare both laboratory-based measures of eating behaviour and other  
349 measures to ascertain the external validity of SST training. Despite the null result for  
350 saturated fat intake, SST training did result in a small but significant decrease in BMI  
351 amongst the participants in the food-specific condition. This reflects recent findings that  
352 GNG task training improves weight loss (Veling et al., 2014) and may indicate that the  
353 current training did alter eating behaviour, but the measure used to assess this outcome was  
354 not sensitive enough to detect such changes. While food frequency questionnaires in general  
355 have been shown to be effective at assessing change in eating behaviour in intervention  
356 studies (Kristal, Beresford, & Lazovich, 1994), it is possible that this particular questionnaire  
357 was not appropriate. However, it must be noted that the training paradigm used in the current  
358 study differed from that used by Houben (2011) and Veling et al. (2014), which may account  
359 for the dissimilar results rather than an issue with the instrument used to measure eating  
360 behaviour.



## 361 **Limitations**

362           Insufficient practice trials in the Stroop task may have precluded the observation of  
363 changes in inhibitory control. Secondly, using a food frequency questionnaire that does not  
364 take into account portion size may not have been sufficient to capture subtle changes in  
365 eating behaviour. Finally, these results need to be replicated with objectively measured height  
366 and weight, as it may be the case that the change observed in BMI was an artefact of self-  
367 report.

## 368 **Study 2**

369           Study 2 was designed to address these limitations and establish the reliability of the  
370 previously observed effects. Namely, by using an objective measure of BMI, increasing the  
371 number of practice trials used in the Stroop, and using an alternative measure of eating  
372 behaviour. The National Cancer Institute (NCI) percentage energy from fat screener  
373 (Thompson et al., 2007) has been validated in intervention studies (Thompson et al., 2008;  
374 Williams et al., 2008), finding that the instrument was consistent at two time points with the  
375 gold-standard method of assessing dietary behaviour: the 24-hour food recall (Carter,  
376 Sharbaugh, & Stapell, 1981). An additional objective was to include follow-up assessments  
377 in order to determine whether training gains persist over time.

## 378 **Method**

### 379 **Participants**

380           Seventy-eight students and staff from a variety of disciplines at an Australian  
381 university (age = 22.97 years,  $SD = 5.81$ ; BMI = 23.11,  $SD = 2.56$ ; 61 females) were  
382 recruited to participate in a study in exchange for course credit or \$20. The number of  
383 participants recruited was based on an a-prior power analysis conducted using G-Power  
384 software (Faul et al., 2007), which indicated that a sample size of 57 would be sufficient to  
385 detect a small to medium (0.15) interaction effect between three conditions at three time

386 points with a power of .80 and an alpha of .05. Inclusion criteria and randomisation did not  
387 differ from Study 1. Participants were randomly allocated to the following conditions: *food-*  
388 *specific inhibition* ( $n = 27$ ), *general inhibition* ( $n = 26$ ), and *control* ( $n = 25$ ).

### 389 **Materials and measures**

390 **BMI & fat intake.** Participants' height was recorded at Time 1 and weight was  
391 measured at each time point on the same set of digital weight scales. Eating behaviour was  
392 operationalised as percentage daily fat intake as measured using the 17-item NCI percentage  
393 energy from fat screener (Thompson et al., 2007). Participants indicated how often they ate  
394 15 food items (e.g., fruit, sausage or bacon, full fat cheese) on a 6-point scale ranging from 0  
395 to 5: never (0), to 2 or more times per day (5). Additionally, participants were asked to  
396 indicate how often they used a reduced-fat butter or margarine when they prepared foods with  
397 butter or margarine, on a 6-point scale ranging from 0 to 5: Didn't use butter or margarine  
398 (0), to almost always or always (5). Finally, participants were asked to indicate whether they  
399 considered their diet to be low, medium, or high in fat. Percentage energy from fat was  
400 calculated using scoring algorithms that assign sex- and age- specific median portion sizes in  
401 grams to each item and then uses a regression model to estimate the expected intake given the  
402 screener responses.

403 **Stroop interference.** Inhibitory control capacity was assessed using the same  
404 computerised version of the Stroop task as Study 1; however, the number of practice trials  
405 was increased from 20 to 50.

406 **Depletion task and Stop-signal task.** The depletion task and the three versions of the  
407 SST did not differ from Study 1.

## 408 **Procedure**

409           This was identical to Study 1 with two exceptions. Measurements of all outcomes  
410 were conducted in the laboratory and a third measurement time point was included one week  
411 after training was completed.

## 412 **Data analyses**

413           Randomisation checks, drop-out analyses and depletion checks were performed as per  
414 Study 1. To assess the effect of training on Stroop performance and vulnerability to depletion  
415 two 3(time: pre-intervention, post-intervention, follow-up) by 3(condition: food-specific  
416 inhibition, general inhibition, control) mixed ANOVAs were conducted. Overall effects were  
417 examined; however, focus was placed on time by condition interactions between two sets of  
418 levels of the within-participants factor (pre-intervention versus post-intervention, and pre-  
419 intervention versus follow-up). If a significant time by condition interaction was detected for  
420 either comparison, planned contrasts examining differences between the two training  
421 conditions and the control, and between the two training conditions themselves, were  
422 conducted. Similarly, to assess the effect of training on percentage energy from fat and BMI,  
423 two 3 x 3 mixed ANOVAs were conducted; with planned contrasts examining pre- to post-  
424 intervention, and pre-intervention to follow-up differences between the food-specific  
425 inhibition condition and other conditions, and between the training conditions themselves.

## 426 **Results**

### 427 **Randomisation check**

428           There were no significant differences on measured variables between conditions pre-  
429 intervention, all  $p > .05$ . Additionally, the number of SSTs performed across the training  
430 period did not differ between conditions,  $p > .05$ .

### 431 Attrition

432 Eight participants did not complete post-intervention and follow-up data (food-  
433 specific inhibition:  $n = 3$ , general inhibition:  $n = 3$ , control:  $n = 2$ ). Five participants dropped  
434 out of the study and three did not sufficiently engage with all tasks. All drop-out occurred at  
435 the second time point (post-intervention). There were no differences on measures, all  $p > .05$ ,  
436 between those who completed the study and those who did not.

### 437 Depletion

438 Participants' performance on the Stroop task was significantly poorer following the  
439 depletion task,  $MD = -109.527$ ,  $SE = 15.323$ ;  $t(77) = -7.148$ ,  $p < .001$ . Additionally, on  
440 average participants reported the task as difficult,  $M = 6.28$ ,  $SD = 0.79$ , unpleasant,  $M = 5.23$ ,  
441  $SD = 1.01$ , frustrating,  $M = 3.23$ ,  $SD = 0.82$ , and effortful,  $M = 3.58$ ,  $SD = 0.85$ .

### 442 Training effects

443 Means and standard deviation of all test variables at pre-intervention, post-  
444 intervention, and follow-up are displayed in Table 2.

445 INSERT TABLE 2 NEAR HERE

446 **Inhibitory control.** There was a significant main effect of time indicating that  
447 averaged across all conditions, there were differences in Stroop performance according to the  
448 three time points,  $F(2, 134) = 22.687$ ,  $p < .001$ , partial  $\eta^2 = .253$ . Additionally, there was a  
449 significant time by condition interaction, indicating that the differences in Stroop  
450 performance according to time were not the same for each condition,  $F(4, 134) = 4.489$ ,  $p =$   
451  $.002$ , partial  $\eta^2 = .118$ . There was no main effect of condition,  $p > .05$ .

452 A planned contrast examining the significant interaction effect revealed that both  
453 training conditions performed better on the Stroop post-intervention compared to the control  
454 condition,  $\psi = 92.492$ ,  $F(1, 67) = 11.973$ ,  $p = .001$ . However, this improvement was not

455 maintained at follow-up as a planned contrast between pre-intervention and follow-up  
456 performance did not indicate significant differences between training conditions and the  
457 control,  $\psi = 9.105$ ,  $F(1,67) = .163$ ,  $p = .688$ . Additionally, improvement in performance  
458 demonstrated by the food-specific condition from pre- to post- intervention did not differ to  
459 that demonstrated by the general training condition,  $\psi = 4.358$ ,  $F(1,67) = .020$ ,  $p = .887$ ,  
460 indicating that both forms of SST training improved inhibitory control as measured by the  
461 Stroop. The performance of all conditions across all time points is displayed in Figure 4.

462 INSERT FIGURE 4 NEAR HERE

463 **Vulnerability to depletion.** There was a significant main effect of time indicating  
464 that averaged across all conditions, there were differences in vulnerability to depletion  
465 according to the three time points,  $F(2, 134) = 7.765$ ,  $p = .001$ , partial  $\eta^2 = .104$ .  
466 Additionally, there was a significant time by condition interaction, indicating that the  
467 differences in vulnerability to depletion according to time were not the same for each  
468 condition,  $F(4, 134) = 2.661$ ,  $p = .035$ , partial  $\eta^2 = .074$ . There was no main effect of  
469 condition,  $p > .05$ .

470 A planned contrast examining the significant interaction revealed that both training  
471 conditions decreased in vulnerability to depletion post-intervention compared to the control  
472 condition,  $\psi = 76.995$ ,  $F(1, 67) = 8.347$ ,  $p = .001$ . However, this improvement was not  
473 maintained at follow-up as a planned contrast between pre-intervention and follow-up  
474 performance did not indicate significant differences between training conditions and the  
475 control,  $\psi = 12.181$ ,  $F(1,67) = .195$ ,  $p = .661$ . Additionally, the decrease in vulnerability to  
476 depletion demonstrated by the food-specific condition from pre- to post- intervention did not  
477 differ to that demonstrated by the general training condition,  $\psi = .837$ ,  $F(1,67) = .001$ ,  $p =$   
478  $.975$ , indicating that both forms of SST training resulted in decreased vulnerability to  
479 depletion. The performance of all conditions across all time points is displayed in Figure 5.

480 INSERT FIGURE 5 NEAR HERE

481 **Percentage energy from fat.** There were no effects of time, condition, nor were any  
482 time by condition interactions effects significant, all  $p > .05$ .

483 **BMI.** There were no effects of time, condition, nor were any time by condition  
484 interactions effects significant, all  $p > .05$ .

#### 485 **Discussion**

486 The aim of this study was to replicate and address the limitations of Study 1. The  
487 results suggested that both forms of training led to improvement in inhibitory control and  
488 vulnerability to depletion; however, this improvement did not lead to changes in eating  
489 behaviour or BMI. Therefore, the effect of training on vulnerability to depletion was  
490 replicated; however, the effect of food-specific training on BMI was not. The results also  
491 suggested that these improvements in inhibitory control and vulnerability to depletion did not  
492 persist after the training period had ended, suggesting that inhibitory control training may  
493 only improve self-regulatory outcomes in the short-term.

494 The results indicated that both inhibitory control capacity, and vulnerability to  
495 depletion improved after both forms of training. This suggests that repeatedly performing a  
496 task that requires inhibitory control results in improvements in this capacity and in the ability  
497 to exert this capacity after performing another task that requires self-regulation. This is in line  
498 with the strength model of self-regulation, which suggests that self-regulation relies on a  
499 limited pool of resources that can become depleted in the short-term, but strengthened over  
500 time with repeated acts of self-regulation (Baumeister et al., 2007). Additionally, these results  
501 reflect previous research that has used self-regulation training to improve self-regulatory  
502 outcomes. Specifically, Muraven (2010) demonstrated that participants who were instructed  
503 to avoid unhealthy foods for a two week period, or perform a handgrip task daily for two  
504 weeks, showed improved performance on an SST compared to control conditions that did not

505 receive training. However, it appears that while modifying eating behaviour leads to  
506 improvement in inhibitory control, as measured by the SST, practicing the SST does not lead  
507 to changes in eating behaviour. It may be the case that exerting self-regulation in real-life  
508 situations requires more control and results in larger effects that are easily detectable on a  
509 reaction time measure, whereas practicing an abstract task may be a less intense form of  
510 training that does not translate to improvements in everyday behaviour.

511         The finding that SST training, as employed in the current study, did not result in  
512 changes in eating behaviour is unexpected given that research employing other inhibitory  
513 control training paradigms has demonstrated an influence on eating behaviour (Houben,  
514 2011; Houben & Jansen, 2011; Veling et al., 2011; Veling et al., 2013). However, the training  
515 paradigm adopted in the current studies differs substantially from previous research and  
516 therefore may account for the differing results. Firstly, the majority of previous research has  
517 utilised a GNG paradigm in which unhealthy food stimuli are always paired with no-go  
518 responses, rather than only a proportion of them. Thus, it may be the case that target stimuli  
519 have to be consistently paired with a stop response in order to induce change in behaviour.  
520 Additionally, Veling et al. (2014) demonstrated weight loss after four 30 minutes sessions of  
521 GNG spread across four weeks, using greater variety of stimuli. Thus, training may not have  
522 been effective not only due to the low proportion of stop-signals used in the current  
523 paradigm, but also the timing of training sessions and lack of variety in the stimuli that were  
524 used. It is recommended that future research aiming to replicate these training effects employ  
525 a more intense and varied paradigm. Finally, given that the results of Study 2 did not replicate  
526 the change in BMI finding of Study 1, we suggest that this finding may have been due to the  
527 self-report measurement of BMI.

528         The observed changes in inhibitory control and vulnerability to depletion in the two  
529 training conditions were not maintained at follow-up. Although different training paradigms

530 and behavioural outcomes were measured, these results are similar to that of Verbruggen et  
531 al. (2013), who did not find that inhibitory control training produced long-lasting effects.  
532 These results appear to indicate that inhibitory control training may only improve self-  
533 regulation outcomes in the short-term. While Baumeister and colleagues did not directly  
534 hypothesise about the maintenance of improvements in self-regulation (Baumeister et al.,  
535 2007; Hagger, Wood, Stiff, & Chatzisarantis, 2010), the muscle metaphor commonly used to  
536 conceptualise self-regulation can be extended to account for these effects. Specifically, while  
537 exercise can strengthen a muscle, if exercise is not maintained- strength will slowly decline.  
538 Similarly, it appears that if training is not continued, self-regulatory capacity may return to  
539 initial levels. Future research should attempt to replicate these effects in order to further  
540 knowledge regarding the nature of self-regulation.

#### 541 **General Discussion**

542 These studies represent some of the first to assess the efficacy of an SST training  
543 paradigm in the improvement of self-reported health behaviour, in order to determine  
544 whether training translates into change in everyday behaviour and to directly test potential  
545 mechanisms of change. However, there are limitations to these studies that must be  
546 acknowledged. Firstly, it may be the case that presenting stop-signals on only 25% of trials  
547 with the target stimuli was not intense enough to induce a change in eating behaviour.  
548 Research in the field of alcohol consumption demonstrated a change in laboratory based  
549 drinking behaviour after SST training with a 50% stop-signal density (Jones & Field, 2013).  
550 Further, GNG training, in which all trials that display the target stimuli are ‘no-go’ (i.e. stop)  
551 trials, has more consistently resulted in behaviour change (Bowley et al., 2013; Veling et al.,  
552 2014). Therefore, a higher density of stop responses associated with the target behaviour may  
553 be necessary to induce behaviour change and future research should systematically vary the  
554 density of stop-signal trials in order to determine whether this influences the transfer of



555 training to health behaviour. Further, comparing the efficacy of SST training to GNG  
556 training, and whether these paradigms influence behaviour via different mechanisms (i.e.  
557 inhibitory control versus automatic evaluations) is warranted.

558         Additionally, previous research has shown that individual difference variables such as  
559 dietary restraint (Houben & Jansen, 2011; Veling et al., 2011), and homeostatic variables  
560 such as previous food intake and hunger (Loeber, Grosshans, Herpertz, Kiefer, & Herpertz,  
561 2013), influence food cue processing. Future research may benefit from including and  
562 controlling for these variables. Additionally, while the stimulus set used in both interventions  
563 reflected that used in other inhibitory control training and eating behaviour interventions  
564 (Veling et al., 2013), it was not validated for the respective samples. Future research should  
565 assess participants' perceptions of the palatability of food items in order to ensure that the  
566 selected stimuli are considered palatable by the target sample. Finally, because there was not  
567 a control condition in which participants did not receive a depletion task, it is difficult to  
568 ascertain whether the vulnerability to depletion measure accurately assessed this construct.  
569 However, all participants performed poorer on the Stroop that followed the depletion task,  
570 suggesting that this task did in fact induce a depletion effect. Nevertheless, future research  
571 attempting to determine whether SST training can improve vulnerability to depletion should  
572 include a depletion control condition in order to test this assumption.

### 573 **Implications**

574         Despite these limitations, the current results have several implications for  
575 interventions designed to improve self-regulatory outcomes and eating behaviour. Namely, it  
576 appears that this particular inhibitory control training paradigm does not result in changes in  
577 everyday eating behaviour. Comparing the current paradigm to that used in previous research,  
578 it appears that training needs to be of a certain intensity in order to induce change in health  
579 behaviour, such that the proportion of unhealthy food – stop-signal pairings used in the

580 current studies was not intense enough. . Additionally, these results contribute to theoretical  
581 explanations regarding the nature of self-regulation. While it has been established that  
582 elements of self-regulation can be improved through training (Muraven, 2010), the current  
583 results suggest that the benefits of training are only maintained insofar as training is  
584 maintained.

## 585 **Conclusions**

586         The results of two inhibitory control training studies in which the aim was to improve  
587 eating behaviour and demonstrate the mechanism by which this improvement occurs were  
588 reported. The results of Study 2 did not replicate those of Study 1, such that inhibitory control  
589 training in this intervention did not appear to influence health outcomes. However, the results  
590 indicated that inhibitory control training does appear to improve inhibitory control, as  
591 measured by a related task, and the construct of vulnerability to depletion, but these effects  
592 do not appear to persist after training has ceased.

593

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Table 1

*Means and Standard Deviations of All Outcome Variables for Each Condition Pre- and Post- Intervention*

	Pre-intervention						Post-intervention					
	Food-specific		General		Control		Food-specific		General		Control	
	<i>n</i> = 29		<i>n</i> = 25		<i>n</i> = 28		<i>n</i> = 26		<i>n</i> = 21		<i>n</i> = 25	
	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>
Inhibitory control	159.06	114.26	151.79	104.05	132.63	63.56	130.82	81.81	118.74	78.48	107.96	84.72
Depletion	124.90	74.93	100.62	84.58	96.71	72.36	57.47	59.88	47.35	59.85	95.53	83.33
Saturated fat intake	23.16	7.49	24.34	7.04	23.06	6.74	22.01	7.14	23.03	6.28	22.02	6.71
BMI	22.21	2.04	22.78	2.43	22.90	3.31	21.96	2.08	22.65	2.51	22.84	2.94

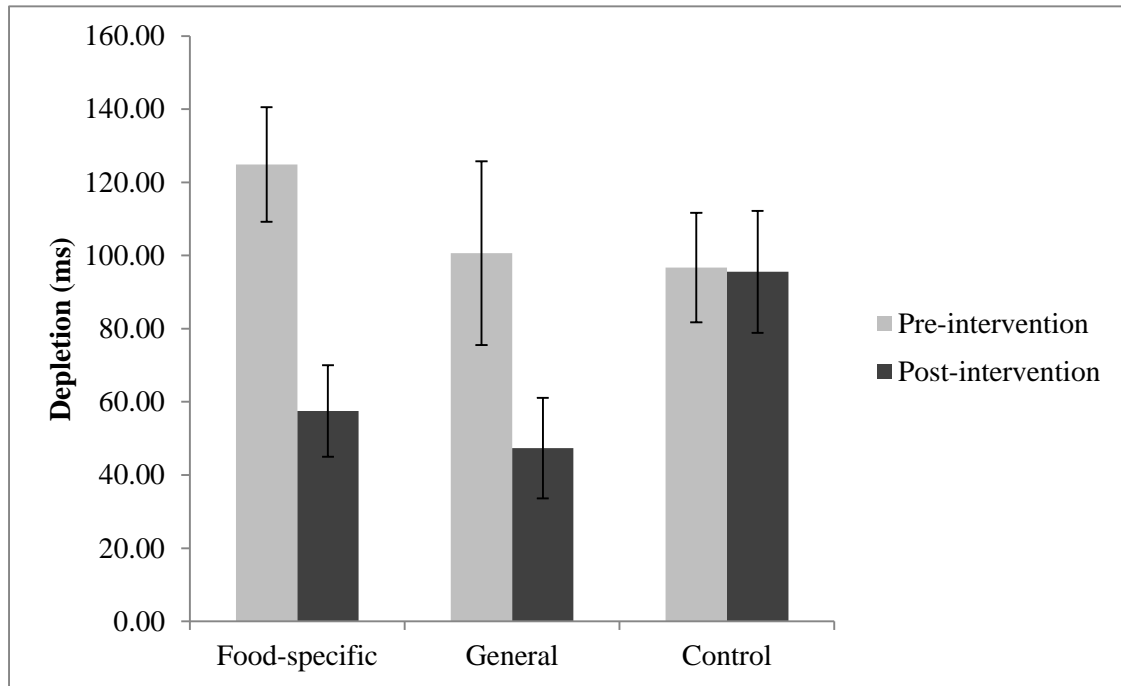
*Note.* Inhibitory control = Stroop interference score (ms); Depletion = difference in Stroop interference scores pre- to post- depletion task (ms), Saturated fat intake = g/day calculated from dietary fat items of the Block food screener, BMI = body mass index.

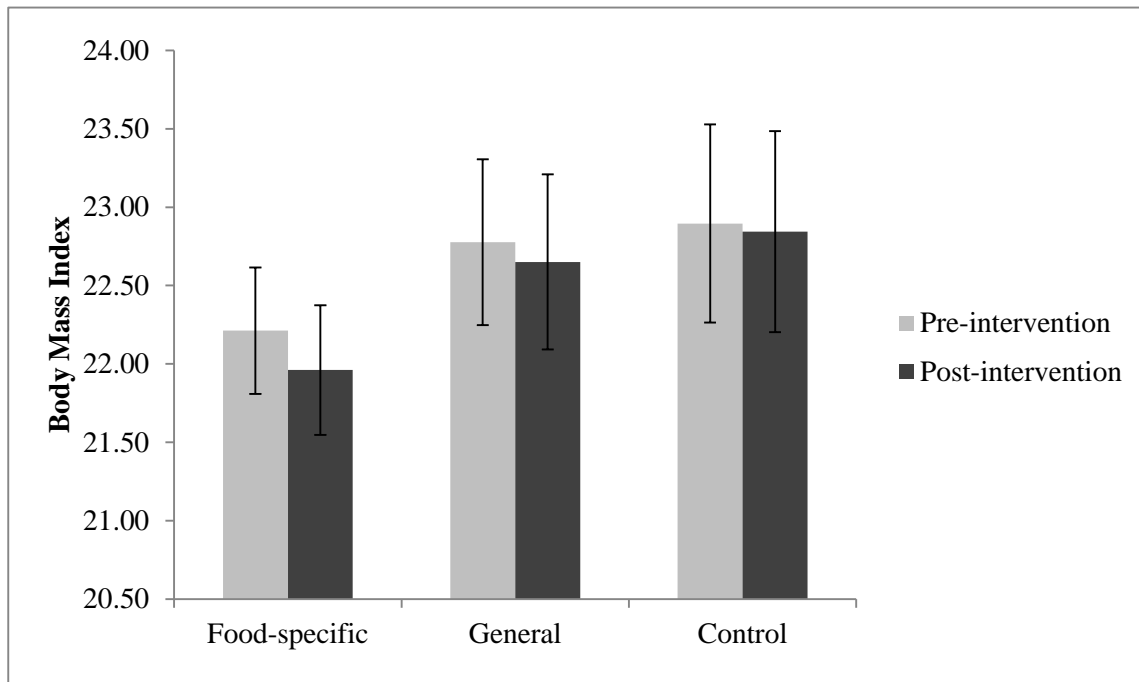
Table 2

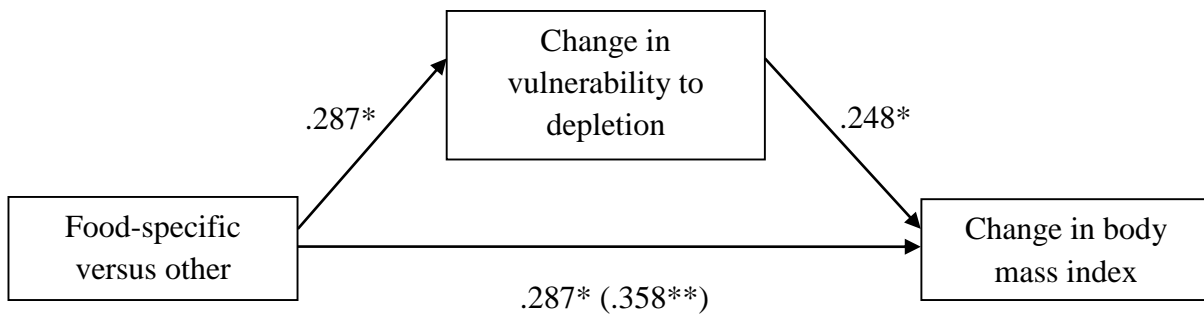
*Means and Standard Deviations of All Outcome Variables for Each Condition at Pre-Intervention, Post-Intervention, and Follow-Up*

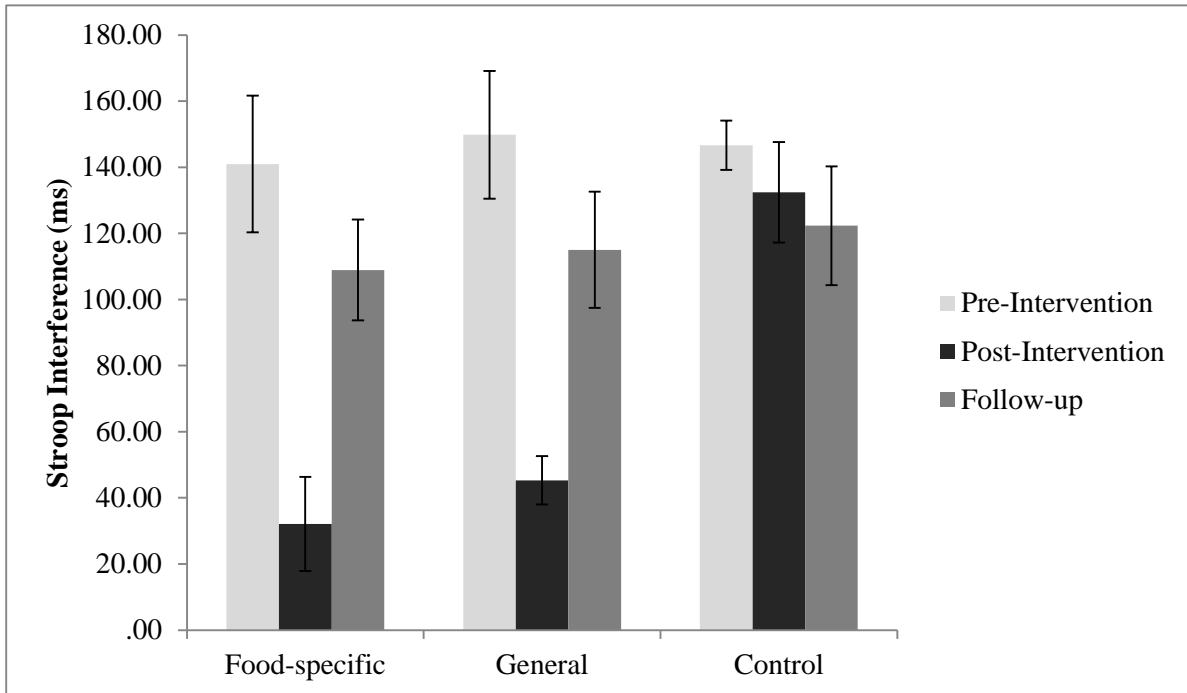
	Pre-intervention						Post-intervention						Follow-up					
	Food-specific		General		Control		Food-specific		General		Control		Food-specific		General		Control	
	<i>n</i> = 27		<i>n</i> = 26		<i>n</i> = 25		<i>n</i> = 24		<i>n</i> = 23		<i>n</i> = 23		<i>n</i> = 24		<i>n</i> = 23		<i>n</i> = 23	
	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>
Inhibitory control	138.86	99.62	145.49	89.47	141.62	38.84	32.10	69.64	45.33	35.21	132.45	72.86	108.92	74.55	115.03	84.25	122.33	86.05
Depletion	114.59	165.03	110.57	120.15	120.91	98.87	54.24	70.62	48.68	75.54	129.88	87.45	119.96	111.29	110.04	101.87	128.61	89.33
% energy from fat	34.63	14.36	34.49	14.24	35.95	12.05	34.02	14.83	34.16	14.41	34.65	13.77	34.95	12.67	35.68	14.21	35.09	17.32
BMI	23.11	2.50	23.01	2.73	23.21	2.54	23.18	2.53	23.01	2.89	23.20	2.72	23.14	2.45	22.97	2.93	23.13	2.60

*Note.* Inhibitory control = Stroop interference score (ms); Depletion = difference in Stroop interference scores pre- to post- depletion task (ms); % energy from fat = fat intake calculated from NCI Percentage Energy from Fat Screener, BMI = body mass index.

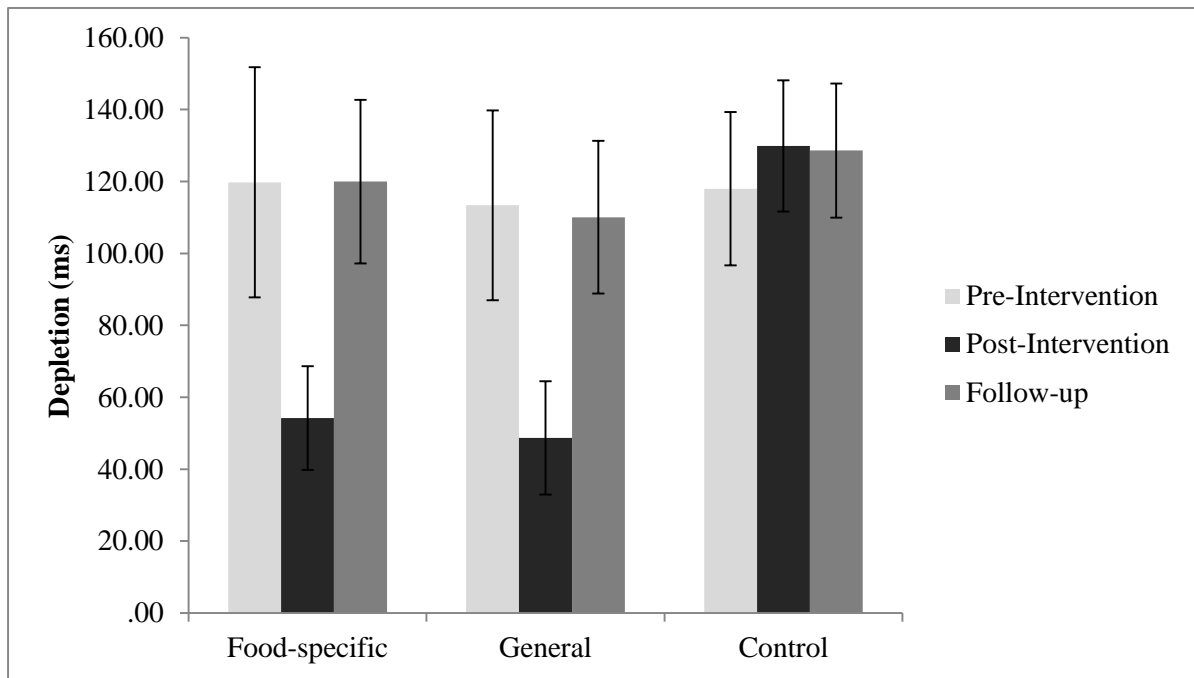
**Figures**











### Figure Captions

*Figure 1.* Amount of depletion (difference in Stroop interference scores pre- to post-depletion task in ms) experienced pre- and post- intervention for each condition. Error bars display standard error.

*Figure 2.* Body mass index pre- and post- intervention for each condition. Error bars display standard error.

*Figure 3.* Simple mediation model depicting the indirect effect of type of training on change in body mass index through change in vulnerability to depletion. Standardised beta coefficients are noted in the diagram, \* $p < .05$ , \*\* $p < .01$ .

*Figure 4.* Inhibitory control performance (Stroop interference scores in ms) pre-intervention, post-intervention and at follow-up for each condition. Error bars display standard error.

*Figure 5.* Amount of depletion (difference in Stroop interference scores pre- to post-depletion task in ms) experienced pre-intervention, post-intervention and at follow-up for each condition. Error bars display standard error.