Running Head: INHIBITORY CONTROL TRAINING

1	Two inhibitory control training interventions designed to improve eating behaviour and
2	determine mechanisms of change
3	
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13		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	•	l 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

21

Abstract

Inhibitory control training has been shown to influence eating behaviour in the laboratory;

22 however, the reliability of these effects is not yet established outside the laboratory, nor are 23 the mechanisms responsible for change in behaviour. Two online Stop-Signal Task training 24 interventions were conducted to address these points. In Study 1, 72 participants completed 25 baseline and follow-up measures of inhibitory control, self-regulatory depletion, fat intake 26 and body-mass index. Participants were randomly assigned to complete one of three Stop-27 Signal Tasks daily for ten days: *food-specific inhibition-* inhibition in response to unhealthy 28 food stimuli only, general inhibition- inhibition was not contingent on type of stimuli, and 29 control- no inhibition. While fat intake did not decrease, body-mass index decreased in the 30 food-specific condition and change in this outcome was mediated by changes in vulnerability 31 to depletion. In Study 2, the reliability and longevity of these effects were tested by 32 replicating the intervention with a third measurement time-point. Seventy participants 33 completed baseline, post-intervention and follow-up measures. While inhibitory control and 34 vulnerability to depletion improved in both training conditions post-intervention, eating 35 behaviour and body-mass index did not. Further, improvements in self-regulatory outcomes 36 were not maintained at follow-up. It appears that while the training paradigm employed in the 37 current studies may improve self-regulatory outcomes, it may not necessarily improve health 38 outcomes. It is suggested that this may be due to the task parameters, and that a training 39 paradigm that utilises a higher proportion of stop-signals may be necessary to change 40 behaviour. In addition, improvements in self-regulation do not appear to persist over time. 41 These findings further current conceptualisations of the nature of self-regulation and have 42 implications for the efficacy of online interventions designed to improve eating behaviour.

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Two inhibitory control training interventions designed to improve eating behaviour and 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 pairings of the no-go response with target stimuli facilitates the retrieval of no-go-target 75 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 a result of inhibitory control training (Veling et al., 2014). This study demonstrated that four 86 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 no direct evidence exists as to what mechanism of change underlies SST training, Jones and 96 97 Field (2013) demonstrated that alcohol-specific SST training led to a reduction in inhibition errors to alcohol stimuli across training blocks, which may suggest that SST training 98 99 improves health behaviour by increasing inhibitory control. Nevertheless, this assumption was not directly tested as no additional measure of inhibitory control was included, thus this 100 101 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

Therefore, the aim of the present research was to improve self-reported eating
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 and proportion of stop-signals were used; however, the stop-signals were not contingent on a 125 126 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 = 4.86; BMI = 22.62, SD = 2.64; 66 females) were recruited to participate in a study in 146 147 exchange for course credit. The number of participants recruited was based on an a-prior power analysis using G-Power 3 software (Faul, Erdfelder, Lang, & Buchner, 2007), which 148 149 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 directed them to one of three pages. The university's human research ethics committee 157 158 approved the study and participants provided informed consent prior to participation. Materials and measures 159

BMI & saturated fat intake. BMI was calculated from participants' self-reported
height and weight. Saturated fat intake in grams was calculated from responses on the Block
food screener (Block, Gillespie, Rosenbaum, & Jenson, 2000), which has been validated
against a 100-item food frequency questionnaire (Block et al., 2000). Participants indicated
how often they ate 17 meat and snack items (e.g. bacon, full-fat ice-cream, fried potatoes) on
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 'red' must be inhibited in order to provide the correct response of 'blue'. The task consisted 170 171 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 =extremely difficult/unpleasant), and frustrating (1 = not at all frustrating - 5 = extremely187 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 on the task?" (1 = not at all - 5 = extremely), and written responses were reviewed to ensure 190

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

194 Stop-signal task. The current study utilised three versions of the SST with cues, which included three experimental blocks of 64 trials and a practice block of 32 trials. In all 195 196 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).

For the *food-specific inhibition* condition, the stop-signal was only presented after unhealthy food images. Therefore, each block consisted of 16 unhealthy food-stop trials, 16 unhealthy food go-trials, 0 healthy food-stop trials and 32 healthy food-go trials. For the *general inhibition* condition, the stop-signal was randomly presented either after a healthy or an unhealthy food image. Therefore, each block consisted of 8 unhealthy food-stop trials, 24 unhealthy food-go trials, 8 healthy food-stop trials and 24 healthy food-go trials. For the *control* condition, participants performed the same task as the other conditions; however, no stop-signals were presented. If participants in either training condition inhibited their
responses less than 50% of the time on inhibition trials this was an indication that they were
not responding to the stop-signal correctly and thus that session was not included as a training
session. Similarly, if participants inhibited their responses more than 50% of the time, this
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 up to the study, and provided informed consent, they completed the pre-intervention 228 229 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

In order to confirm that randomisation was successful the three experimental
conditions were compared with respect to scores on age, BMI, Stroop interference,
vulnerability to depletion, and saturated fat intake using a one-way analysis of variance
(ANOVA), while a chi-squared analysis was utilised to assess sex differences between
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 inhibition, general inhibition, control) mixed ANOVAs were conducted. If a significant time 247 248 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 p261 > .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 >

.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**

Inhibitory control. There was a significant main effect of time indicating that all conditions improved on Stroop performance post-intervention, F(1, 69) = 4.635, p = .035, partial eta² = .063. There was no main effect of condition, nor was the time by condition interaction effect significant, all p > .05. See Table 1 for pre- and post- intervention means 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

general inhibition condition, $\psi = 23.953$, F(1,69) = .8599, p = .357. There was no main effect 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 decreased in BMI post-depletion, F(1, 69) = 10.048, p = .002, partial eta² = .127, which was 294 qualified by a significant time by condition interaction effect, F(2, 69) = 5.086, p = .009, 295 296 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 BMI did not change in the general inhibition condition and the control, $\psi = .354$, F(1,69) =298 299 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 effect from food-specific training, through change in vulnerability to depletion, to change in 312

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, Liefooghe,
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

were not only less vulnerable to depletion but also reported engaging in more health
enhancing behaviours. In contrast, within the current study this improvement only transferred
to change in BMI in the food-specific condition, suggesting that behavioural specificity of the
task, coupled with decrease in vulnerability to depletion may be necessary to change
behaviour. Alternatively, it may be that more intense training is required for improvements to
translate across behavioural domains. Further research is required to determine the optimal
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 for the dissimilar results rather than an issue with the instrument used to measure eating 359 360 behaviour.

361 Limitations

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

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

Seventy-eight students and staff from a variety of disciplines at an Australian university (age = 22.97 years, SD = 5.81; BMI = 23.11, SD = 2.56; 61 females) were recruited to participate in a study in exchange for course credit or \$20. The number of participants recruited was based on an a-prior power analysis conducted using G-Power software (Faul et al., 2007), which indicated that a sample size of 57 would be sufficient to detect a small to medium (0.15) interaction effect between three conditions at three time points with a power of .80 and an alpha of .05. Inclusion criteria and randomisation did not differ from Study 1. Participants were randomly allocated to the following conditions: *foodspecific 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

Randomisation checks, drop-out analyses and depletion checks were performed as per 413 414 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 inhibition condition and other conditions, and between the training conditions themselves. 425

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.

19

431 Attrition

Eight participants did not complete post-intervention and follow-up data (foodspecific inhibition: n = 3, general inhibition: n = 3, control: n = 2). Five participants dropped out of the study and three did not sufficiently engage with all tasks. All drop-out occurred at the second time point (post-intervention). There were no differences on measures, all p > .05, 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 three time points, F(2, 134) = 22.687, p < .001, partial eta² = .253. Additionally, there was a 448 449 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 =.002, partial $eta^2 = .118$. There was no main effect of condition, p > .05. 451 452 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

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

INSERT FIGURE 4 NEAR HERE

Vulnerability to depletion. There was a significant main effect of time indicating that averaged across all conditions, there were differences in vulnerability to depletion according to the three time points, F(2, 134) = 7.765, p = .001, partial eta² = .104. Additionally, there was a significant time by condition interaction, indicating that the differences in vulnerability to depletion according to time were not the same for each condition, F(4, 134) = 2.661, p = .035, partial eta² = .074. There was no main effect of 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 =.975, indicating that both forms of SST training resulted in decreased vulnerability to 478 479 depletion. The performance of all conditions across all time points is displayed in Figure 5.

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 reflect previous research that has used self-regulation training to improve self-regulatory 501 502 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

receive training. However, it appears that while modifying eating behaviour leads to improvement in inhibitory control, as measured by the SST, practicing the SST does not lead to changes in eating behaviour. It may be the case that exerting self-regulation in real-life situations requires more control and results in larger effects that are easily detectable on a reaction time measure, whereas practicing an abstract task may be a less intense form of 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 the change in BMI finding of Study 1, we suggest that this finding may have been due to the 526 527 self-report measurement of BMI.

528 The observed changes in inhibitory control and vulnerability to depletion in the two529 training conditions were not maintained at follow-up. Although different training paradigms

23

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 hypothesise about the maintenance of improvements in self-regulation (Baumeister et al., 534 535 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 be necessary to induce behaviour change and future research should systematically vary the 553 554 density of stop-signal trials in order to determine whether this influences the transfer of

- 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 dietary restraint (Houben & Jansen, 2011; Veling et al., 2011), and homeostatic variables 559 560 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 current studies was not intense enough. Additionally, these results contribute to theoretical explanations regarding the nature of self-regulation. While it has been established that elements of self-regulation can be improved through training (Muraven, 2010), the current results suggest that the benefits of training are only maintained insofar as training is maintained.

585 Conclusions

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

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Table 1

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

			Pre-inte	rvention			Post-intervention								
	Food-s	pecific	General		Control		Food-s	pecific	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				
	М	SD	М	SD	М	SD	М	SD	М	SD	М	SD			
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.

		on		Post-intervention							Follow-up							
	Food- specific		Food- General specific		Control		Food- specific		General		Control		Food- specific		General		Control	
	<i>n</i> = 27		= 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	
	М	SD	М	SD	М	SD	М	SD	М	SD	М	SD	М	SD	М	SD	М	SD
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

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

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 postdepletion task in ms) experienced pre-intervention, post-intervention and at follow-up for each condition. Error bars display standard error.