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1        **No Effects of Ingesting or Rinsing Sucrose on Depleted Self-Control Performance**

2  
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9  
10  
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12 **Short title:** Self-control, Ego-depletion, & Carbohydrates

## Abstract

15  
16 Self-control tasks appear to deplete a limited resource resulting in reduced subsequent self-  
17 control performance; a state of ego depletion. Evidence of reduced peripheral glucose by  
18 exertion of self-control, and attenuation of ego depletion by carbohydrate metabolism  
19 underpins the proposition that this macronutrient provides the energetic source of self-  
20 control. However, the demonstration of positive, non-metabolic effects on ego depletion  
21 when merely sensing carbohydrates orally contradicts this hypothesis. Recent studies have  
22 also failed to support both metabolic and non-metabolic accounts. The effects of ingesting or  
23 rinsing a carbohydrate (sucrose) and an artificially sweetened (sucralose) solution on  
24 capillary blood and interstitial glucose, and depleted self-control performance were examined  
25 in older adults. Forty, healthy, adults (50–65 years) ingested and rinsed sucrose and sucralose  
26 solutions in a 2(method) x 2(source), fully counterbalanced, repeated measures, crossover  
27 design. Capillary blood and interstitial glucose responses were assayed. Depleted self-control  
28 performance (induced by the Bakan visual processing task) on an attention switch task was  
29 assessed under each study condition. Ego depletion had no consistent effects on peripheral  
30 glucose levels and no significant effects of ingesting or rinsing sucrose on self-control were  
31 observed. The act of rinsing the solutions, independent of energetic content, resulted in a  
32 small, non-significant enhancement of performance on the attention switch task relative to  
33 ingesting the same solutions (RT:  $p = .05$ ; accuracy:  $p = .09$ ). In conclusion, a metabolic  
34 account of self-control was not supported. Whilst a positive effect of rinsing on depleted self-  
35 control performance was demonstrated, this was independent of energetic content. Findings  
36 suggest glucose is an unlikely physiological analogue for self-control resources.

37

38

39 **The Effect of Ingesting and Rinsing Sucrose and Sucralose on Depleted Self-Control**  
40 **Performance**

41  
42 Acts of self-control require the effortful inhibition of predominant responses,  
43 emotions, thoughts, and impulses, permitting behaviour to vary adaptively moment to  
44 moment <sup>(1, 2)</sup>. Exertion of self-control is considered to be a key process in the human  
45 personality structure as flexibility in behavioural response permits the attainment of goals,  
46 and facilitates adherence to rules, laws, and social norms and standards <sup>(3)</sup>. Indeed, self-  
47 control capacity has been positively associated with an impressive array of behaviours of  
48 personal and societal significance (e.g., reduced aggression <sup>(4)</sup>; scholastic achievement <sup>(5)</sup>;  
49 interpersonal success <sup>(6)</sup>; criminality <sup>(7)</sup>).

50  
51 The capacity to exert self-control appears to be limited <sup>(8)</sup>. The resource strength  
52 model maintains that acts of self-control consume and temporarily deplete a common, and  
53 crucially limited, resource; ultimately resulting in ‘ego depletion’ <sup>(9, 10)</sup>. Self-control  
54 performance is therefore determined by the current strength or level of depletion of this  
55 common resource. Indeed, initial expenditure of self-control has been repeatedly  
56 demonstrated to result in reduced subsequent performance on self-control tasks independent  
57 of differing task modalities <sup>(11)</sup>. This would fit with the claim that the resource involved in  
58 inhibition and self-control reflects a form of executive function <sup>(12, 13)</sup>.

59  
60 Gailliot and Baumeister <sup>(14)</sup> extended the strength model from the metaphorical to the  
61 physical by proposing that glucose is the central energy source of self-control. This  
62 proposition was founded on evidence of (i) reduced blood glucose levels after initial exertion  
63 of self-control; (ii) an association between subsequent, post-depletion, self-control  
64 performance and blood glucose decline, and (iii) attenuation of the detrimental ego depletion  
65 effect on self-control performance after ingestion of glucose, but not artificial sweetener  
66 (sucralose) <sup>(14)</sup>. Subsequent attempts to replicate the moderation of peripheral blood glucose  
67 by exertion of self-control have not supported this original finding <sup>(15)</sup>. The amount of glucose  
68 required for acts of self-control is likely to be negligible in absolute brain energy cost terms.  
69 Furthermore, reduced peripheral glucose is unlikely considering the efficiency of homeostatic  
70 systems in maintaining brain energy levels <sup>(16)</sup>.

71

72           The capacity of glucose ingestion to counteract the impairing effect of ego depletion  
73 has however been replicated <sup>(17, 18, 19, 20)</sup>. Nevertheless, the precise role of glucose in self-  
74 control performance remains indistinct. Firstly, a number of studies have demonstrated that  
75 glucose can influence performance on self-control tasks in a non-energetic manner. Merely  
76 sensing carbohydrates in the oral cavity can confer a restorative benefit on cognitive self-  
77 control performance under conditions of ego depletion <sup>(21, 15, 22)</sup>. The positive effect of  
78 carbohydrate oral rinsing has also been demonstrated in physical endurance performance <sup>(23,</sup>  
79 <sup>24)</sup>, conferring greater performance benefits than ingestion <sup>(25)</sup>. Such findings suggest a  
80 potential motivational rather than metabolic effect of carbohydrates on performance,  
81 underpinned by activation of motivational neural reward pathways <sup>(24, 26, 15)</sup>. Ego depletion  
82 effects can also be moderated by manipulation of subjective states such as motivation <sup>(27)</sup>,  
83 expectation of self-control capacity <sup>(28)</sup>, and self-affirmation <sup>(29)</sup>. Evidence of moderation of  
84 self-control performance by subjective state casts further doubts on the specific role of  
85 glucose.

86

87           The existing evidence for metabolic accounts of self-control (e.g., <sup>(14, 11)</sup>) has also  
88 received criticism on the statistical grounds of potentially inflated effect sizes and  
89 methodological shortcomings <sup>(16, 30, 31, 32)</sup>. Further, failed attempts to replicate the effects of  
90 carbohydrate ingestion and oral sensing suggest the effect of carbohydrate on self-control  
91 performance may not be as robust as the literature suggests <sup>(30, 31)</sup>. In sum, whilst the ingestion  
92 and rinsing of carbohydrate-containing solutions have previously been shown to attenuate the  
93 ego depletion effect, critiques of the existing evidence, and recent failures to replicate the  
94 effect, necessitate further independent examination of the relationship between carbohydrate  
95 and self-control.

96

97           The present study examined the effect of ingesting and rinsing a sucrose and sucralose  
98 solution on a self-control task under conditions of ego depletion. The potential depletion of  
99 glucose by the exertion of self-control, and moderation of self-control by the metabolism of  
100 glucose, were rigorously assessed using formal laboratory standard capillary blood glucose  
101 analysis techniques and continuous interstitial glucose monitoring. Previous studies  
102 examining the effects of glucose ingesting and rinsing on self-control have recruited young,  
103 predominantly student samples. The facilitative effects of carbohydrate intake on cognitive  
104 performance may be more potent in individuals with disrupted metabolic or cognitive  
105 functioning <sup>(33, 34, 35, 36, 37)</sup> rather than young, high functioning students. Indeed, glucose

106 administration has been shown to selectively enhance cognitive performance in elderly, but  
107 not the young<sup>(33, 34)</sup> and was additionally mediated by glucoregulatory control in older but not  
108 younger adults<sup>(33)</sup>. Ageing is associated with cognitive decline and disturbed regulation of  
109 primary hormones and neurotransmitters mediating glucose regulation and cognitive function  
110 (e.g., acetylcholine, adrenaline<sup>(37)</sup>). Therefore, older adults were considered a highly relevant  
111 population in which to study the effects of ingesting and rinsing carbohydrate solutions on  
112 self-control performance, due to commonly observed age-related decline in cognitive control  
113<sup>(38)</sup> and an age-associated deterioration in glucose regulation<sup>(37)</sup>.

## 114 **Methods**

### 115 **Design**

116 A 2 (method: rinse or ingest) X 2 (source: sucrose or sucralose) fully counterbalanced  
117 (William's Latin Square), within subjects, crossover design was employed to expose  
118 participants to each experimental condition. Participants were required to ingest or orally  
119 rinse sucrose and sucralose solutions (sucrose/ingest; sucrose/rinse; sucralose/ingest;  
120 sucralose/rinse) over four study visits separated by one week. At each visit, self-control was  
121 depleted prior to solution intake and performance on a different task requiring self-control  
122 was assessed post-intake. This study was conducted according to the guidelines laid down in  
123 the Declaration of Helsinki and all procedures involving human participants were approved  
124 by the University of Leeds' School of Psychology Research Ethics Committee. The study  
125 was registered on Clinicaltrials.gov on February 26, 2014, registration identification number:  
126 NCT02075333 (available at <https://www.clinicaltrials.gov/ct2/show/NCT02075333>) Written  
127 informed consent was obtained from all participants. An honorarium of £140 was paid upon  
128 completion of the study.

### 130 **Participants**

131 Forty (16 males; 24 females), non-smoking, non-obese ( $\bar{x} = 25.20$ ,  $SD = 2.53$  kg/m<sup>2</sup>;  
132 BMI < 30 kg/m<sup>2</sup>; WHO, 2013), older adults ( $\bar{x} = 57.75$ ,  $SD = 5.79$  years) were recruited from  
133 the University campus and local community. Participants self-reported to be free from  
134 symptoms of dementia, depression, Type II diabetes, phenylketonuria or other conditions that  
135 precluded the ingestion of sucralose, and were not taking medication likely to influence  
136 glucose metabolism or cognitive function. Volunteers with impaired glucose tolerance (2 hr  
137 post-prandial capillary blood glucose  $\geq 7.8$  mmol/L) were excluded at screening. Fifteen  
138 female participants reported themselves as post-menopausal. After eligibility screening,  
139 participants were randomly assigned to a counterbalanced experimental condition order.

140

## 141 **Glucose Measures**

142 Capillary blood glucose was obtained via finger prick measures at - 40, - 24, - 2, + 5,  
143 + 19, + 33, + 37, and + 44 minutes relative to solution intake across study visits. Blood  
144 glucose was analysed using a YSI 2700 Glucose/Lactate Analyzer (Yellow Springs  
145 Instruments, Yellow Springs, OH). Interstitial glucose was also measured in a subgroup of  
146 the total sample (5 male; 5 female) using a subcutaneous continuous glucose monitoring  
147 system (CGMS iPRO, Medtronic MiniMed, CA, USA). The CGMS was fitted the day prior  
148 to each study visit and continuously measured interstitial glucose every 5 minutes until the  
149 end of the test day. The time of solution ingestion/rinse was recorded and used to centre the  
150 response profile (0 minutes). Consequently, measures taken - 50 minutes pre- and + 40  
151 minutes post-solution intake are reported. A finger prick measure collected upon waking on  
152 the study visit morning and measures collected by the experimenter at - 40, + 5, and + 33  
153 minutes at each visit were used to calibrate the CGMS. Capillary blood and interstitial  
154 glucose levels are reported in millimols per litre (mmol/L).

155

## 156 **Self-Control Tasks**

157 All cognitive tests were presented using E prime software (Psychology Software  
158 Tools, Inc, PA, USA) on a Dell Optiplex 760 desktop computer with a 17" monitor (screen  
159 resolution 1280 x 800 pixels). Responses for both tasks were made on a keyboard spacebar.

160

161 **Self-control depletor task.** The Bakan task <sup>(39)</sup> was employed to deplete self-control.  
162 The Bakan is a rapid visual information processing task that requires respondents to attend to  
163 and monitor the rapid continuous presentation of single digits whilst maintaining sufficient  
164 self-control to be able to detect and respond to infrequent target stimuli (the presentation of  
165 three consecutive odd or even digits). The BAKAN can be classified within Baumeister et  
166 al.s self-control spheres of attentional and impulse control <sup>(11)</sup>. Attentional control was  
167 required to focus attention (monitoring stimuli for targets) and disregard distractions (not  
168 responding to the more frequent non-target stimuli). Impulse control was required to override  
169 the prepotent non-response to the more frequently presented non-target stimuli when  
170 infrequent target stimuli appeared. Stimuli were presented at a rate of 500 milliseconds per  
171 item, with a 100 ms inter-stimulus delay. One hundred stimuli were presented per one minute  
172 block. Participants were required to identify ten randomly presented targets (three  
173 consecutive digits) in each one minute block. Nine, one minute blocks were presented (900

174 hundred total trials). Stimuli were presented in the centre of a light grey display in black  
175 (bold, Courier New, 18 point) font. Target accuracy and target reaction time (RT) were  
176 recorded to monitor compliance with the task to ensure comparable engagement with the  
177 depletor task across each experimental condition.

178

179 **Attention switching task.** An attention switch task, originally devised by Wylie et al.  
180 <sup>(40)</sup> that combines a task-switch paradigm with a Go/noGo task was employed as a measure of  
181 self-control performance. Attention switch tasks typically require respondents to repeatedly  
182 perform a task on some trials then switch to another task when cued to do so, thus requiring  
183 the effortful suppression of a dominant response (the repeated trials). Performance on  
184 repeated trials is typically superior to performance on “switch” trial due to the time and effort  
185 needed to switch between the two tasks – the switch cost <sup>(40)</sup>. Letter-number pairs were  
186 presented on a horizontal plane in the centre of the screen for 1800 ms (120 ms inter-stimulus  
187 delay). Each character was positioned 1° to the left or right of the central fixation point  
188 (randomly determined). Letters were taken from a set containing 4 vowels (A, E, I, and U)  
189 and four consonants (G, K, M, and R). The numbers were taken from a set containing 4 even  
190 (2, 4, 6, and 8) and 4 odd numbers (3, 5, 7, and 9). The letter-number pairs were presented in  
191 two alternating colours every three trials. Respondents were required to make a Go/noGo  
192 choice based upon the colour of the letter-number pairs. For example, when letter-number  
193 pairs were red, respondents were required to respond when the letter was a vowel (Go), but  
194 not when the letter was a consonant (noGo). Alternatively, when the letter-number pairs  
195 switched to blue, respondents were required to respond when the number was even (Go), but  
196 not when the number was odd (noGo). The three trials in each task-run are split into switch,  
197 nested, and pre-switch trials. Switch trials are the first letter-number pairs presented after the  
198 task-switch (i.e., the Go/noGo colour switch). Nested and pre-switch are the subsequent  
199 repeat trials within the same task-run (see Figure 1 for stimulus configuration).

200 <FIGURE 1>

201 In total 144 trials were presented with target trials randomly presented with a  
202 probability of 50%. Eight parallel versions of the task, differing only with respect to colours  
203 used to cue the task-switch, were employed, and administered in a counterbalanced manner.  
204 Accuracy (number of correctly identified targets) and RT (target response latency) across  
205 each trial type were assessed.

206

## 207 **Subjective Measures**

208 The Profile of Mood States (POMS; <sup>(41)</sup>) was administered to measure transient  
209 subjective mood (tension-anxiety, depression-dejection, anger-hostility, vigour-activity,  
210 fatigue-inertia, and confusion-bewilderment) across each study visit.

211

212 Visual analogue scales (VAS) assessing the sensory properties of the experimental  
213 solutions were employed. How “sweet”, “pleasant”, and “satisfying” participants found the  
214 solutions were rated using 100 mm scales anchored by “*Not at all*” and “*Very*”.

215

## 216 **Experimental Conditions**

217 All study solutions were isoweight (381.4 g) water diluted, commercially available  
218 blackcurrant cordial based solutions. (Sainsbury’s® High Juice Blackcurrant Squash (sucrose  
219 solution) and No Added Sugar Double Strength Blackcurrant Squash (artificially sweetened  
220 equivalent control solution). The solution compositions and nutritional values are shown in  
221 Table 1. The sucrose solution contained 50 g of sugars, providing 25 g of glucose. The  
222 placebo solution was artificially sweetened with sucralose (0.92 g of sugars).

223

224 <TABLE 1>

225

## 226 **Procedure**

227 All study visits were undertaken between 0700 – 1030 hr after an overnight fast  
228 (approximately 10–12 hr fasted state). The timing of all procedures is shown in Figure 2.  
229 Eight capillary blood samples were taken on each test visit. To control for baseline  
230 differences in self-control resources, participants completed a parallel version of the attention  
231 switch task upon arrival at the laboratory. Baseline POMS and the initial self-control  
232 depletory task (BAKAN) were completed before rinsing or ingesting the solutions.  
233 Participants were then required to ingest or rinse a sucrose or sucralose blackcurrant cordial  
234 solution. In the ingest condition, participants drank the entire volume of the solution. In the  
235 rinse condition, participants were instructed to take a mouthful of the solution, rinse it around  
236 the mouth for 5 seconds, then spit it into an empty cup, repeating these steps until they had  
237 rinsed with the entire volume of the solution. In both conditions participants were given 5  
238 minutes to ingest or rinse the solutions. A palatability VAS was completed immediately post-  
239 solution intake. The self-control attention switch task was administered +21 minutes post-

240 solution intake to allow sufficient time for glucose metabolism. Post-solution POMS ratings  
241 were collected after solution intake (+ 7 minutes and + 39 minutes).

242

243 <FIGURE 2>

244

## 245 **Statistical Analyses**

246 Statistical analyses were performed using SAS (Statistical Analysis System, Version  
247 9.2; SAS Institute, Inc., Cary, NC). The SAS PROC PLAN procedure was employed to  
248 randomly generate the William Latin Square counterbalancing schedule. All data were  
249 screened and residual outliers were removed ( $> 2.58$  SD) and residual plots inspected for  
250 deviations from normality.

251

252 A repeated measures analysis (using PROC MIXED) was employed using a mixed  
253 model with subjects as the random factor and a compound symmetry covariance structure of  
254 the repeated measures. For all glucose measurements, method (ingest and rinse), source  
255 (sucrose and sucralose), and time (glucose measurement time points) were the fixed effects in  
256 the model. For the BAKAN (depletor task), method (ingest and rinse), source (sucrose and  
257 sucralose) and trial block (1-9 mins trial blocks) were the fixed effects in the model. For the  
258 attention switch task, method (ingest and rinse), source (sucrose and sucralose) and trial  
259 (switch, nested, preswitch) were the fixed effects in the model. The potential effect of self-  
260 control depletion on peripheral glucose levels was explored by examining interstitial and  
261 capillary blood glucose over the period that participants completed the initial self-control  
262 tasks (i.e. the ego depletion period prior to solution intake). For capillary blood glucose, 3  
263 samples at - 40, - 24, and - 2 minutes (relative to solution intake), corresponding with 10  
264 minutes before the first attention switch task, and 2 minutes prior to, and 2 minutes  
265 immediately after, the BAKAN task, were analysed. Continuous interstitial glucose levels  
266 were analysed between - 40 and - 5 minutes (relative to solution intake) corresponding to  
267 approximately - 10 minutes before the first attention switch task and subsequently at 5  
268 minute intervals until 5 minutes post BAKAN task. Time (blood glucose: - 40, - 24, - 2;  
269 interstitial glucose: - 40, - 35, - 30, - 25, - 20, - 15, - 10, - 5 minutes relative to solution  
270 intake) was the fixed effect in the model.

271

272 Age, sex, and BMI were entered as covariates in all models. Baseline blood glucose  
273 and subjective solution sweetness, pleasantness and satisfaction ratings were entered as

274 covariates in models for the attention switch task. Visit order was entered into all models as a  
275 fixed effect to rule out order effects. All models included the first and second order  
276 interactions of these stated fixed effects. Non-significant covariates and interactions were  
277 removed if they did not contribute to the model and are not reported.

278

279 Participants completed the attention switch task upon arrival at the laboratory to  
280 standardise self-control capacity prior to ego-depletion. It was considered that performance  
281 on this initial task would be too variable, compared to the standardised self-control capacity  
282 post ego-depletion, to serve as a reliable baseline comparison measure for post-depletion  
283 attention switch performance. Moreover, analyses controlling for baseline attention switch  
284 task outcomes revealed inclusion of baseline performance did not significantly alter the  
285 observed effects and were removed from statistical models.

286

287 Blood glucose area under the curve with respect to increase (AUC<sub>i</sub>) was calculated  
288 using the trapezoid method <sup>(42)</sup>. For all analyses, the significance level was set at  $\alpha = 5\%$ . The  
289 nominal  $\alpha$  level was adjusted for multiple post-hoc least squares mean comparisons using the  
290 Tukey-Kramer correction <sup>(43)</sup>. All results (including figures and tables) are presented as mean  
291 and standard error of the mean (SEM).

292

## Results

### 293 Participant Characteristics

294 The continuous interstitial glucose monitor subsample (IPRO) did not significantly  
295 differ from the remaining sample for age,  $t(38) = -0.40$ ,  $p = .69$ , BMI,  $t(38) = 0.42$ ,  $p = .68$ ,  
296 or baseline fasted blood glucose,  $t(38) = -1.25$ ,  $p = .22$ .

297

### 298 Capillary Blood Glucose

299 A significant source\*method\*time interaction was revealed for capillary blood  
300 glucose response,  $F(7, 240) = 42.02$ ,  $p < .001$ . This interaction reflects a significant increase  
301 in capillary blood glucose level in the sucrose/ingest condition. Following intake of the  
302 sucrose solution, blood glucose was significantly higher at + 19, + 33, + 37, and + 44 minutes  
303 compared to - 40, - 24, - 2, and + 5 minutes relative to solution intake (all significant at  $p <$   
304  $.001$ ; Figure 3). Capillary blood glucose was significantly higher in the sucrose/ingest  
305 condition at + 19, + 33, + 37, and + 44 minutes compared to the corresponding time points in  
306 all other experimental conditions ( $p < .001$ ).

307

308 <FIGURE 3>

309

310 Capillary blood glucose AUCi further corroborates the glucose response profile data.  
311 The blood glucose AUCi response was significantly higher in the sucrose/ingest condition  
312 ( $\bar{x} = 102.39 \pm 7.14$  mmol/L),  $F(3,87) = 30.36$ ,  $p < .001$ , compared to all other experimental  
313 conditions (sucrose/rinse:  $\bar{x} = 5.85 \pm 9.33$  mmol/L; sucralose/ingest:  $\bar{x} = - 9.12 \pm 8.90$   
314 mmol/L; sucralose/rinse:  $\bar{x} = - 8.41 \pm 8.29$  mmol/L;  $p < .001$ ). No significant differences  
315 were demonstrated between the remaining experimental conditions.

316

### 317 **Interstitial Blood Glucose**

318 Consistent with the capillary blood glucose response, an interstitial glucose response  
319 was evident in the sucrose/ingest condition only (significant source\*method\*time interaction,  
320  $F(18, 160) = 2.62$ ,  $p < .001$ ; Figure 4). Interstitial glucose was significantly elevated from +  
321 10 minutes onwards after sucrose solution ingestion (compared to prior measurements  
322 between - 50 to 0 minutes ( $p < .001$ ). Interstitial glucose was significantly higher in the  
323 sucrose/ingest condition from + 10 to + 40 minutes compared to the corresponding time  
324 points in all other experimental conditions ( $p < .001$ ).

325

326 <FIGURE 4>

327

### 328 **Effect of Ego Depletion on Peripheral Glucose**

329 **Capillary blood glucose.** A significant main effect of time  $F(2, 78) = 3.75$ ,  $p = .03$ ,  
330 was revealed for capillary blood glucose levels over the self-control depletion period  
331 (corresponding to - 40, -24, - 2 minutes relative to solution intake shown in Figure 3). Post  
332 hoc comparisons revealed a marginal increase in blood glucose between - 40 ( $\bar{x} = 4.64 \pm$   
333  $0.06$  mmol/L) and - 24 ( $\bar{x} = 4.77 \pm 0.06$  mmol/L;  $p = .06$ ) minutes, indicating an increase in  
334 blood glucose during completion of the first attention switch task. Conversely, a significant  
335 decrease was revealed between - 24 and - 2 ( $\bar{x} = 4.62 \pm 0.05$  mmol/L ;  $p = .04$ ) minutes,  
336 indicating a decrease in blood glucose during completion of the BAKAN.

337

338 **Interstitial glucose.** A non-significant effect of time  $F(6, 54) = 0.40$ ,  $p = .10$ , was  
339 revealed for interstitial glucose levels over the self-control depletion period (corresponding to

340 measures between – 40 and – 5 minutes relative to solution intake shown in Figure 4)  
341 indicating no significant effect of self-control depletion on peripheral glucose levels.

342

### 343 **Self-Control Tasks**

344 **Depletor task.** A significant main effect of trial block was revealed for BAKAN  
345 target accuracy,  $F(8, 312) = 20.95$ ,  $p = .001$ , and RT,  $F(8, 312) = 4.25$ ,  $p = .001$ . Accuracy  
346 was significantly higher during the first two trial blocks ( $p < .03$ ), and RT faster during the  
347 first trial block ( $p < .05$ ), of the task compared to performance across the remaining trial  
348 blocks. This effect was independent of experimental condition and reflects a comparable  
349 temporal decline in self-control performance over task exposure at each study visit. Mean  
350 target accuracy and RT across the trial blocks are shown in Figure 5.

351

352 <FIGURE 5>

353

### 354 **Attention Switch Performance**

355 **Reaction time.** A significant main effect of method,  $F(1, 39) = 4.11$ ,  $p < .05$ , trial,  
356  $F(2, 78) = 294.77$ ,  $p < .001$ , and visit,  $F(3, 108) = 21.45$ ,  $p < .001$ , were revealed for attention  
357 switch RT. The method of solution intake (rinse or ingest) irrespective of source (sucrose or  
358 sucralose) significantly affected overall RT performance (i.e., independent of trial type).  
359 Post-hoc tests revealed that RT collapsed across all trials was marginally faster during rinse  
360 compared to ingest conditions ( $p = .05$ ; Table 2).

361

362 Closer inspection of the RTs across trial suggest enhanced performance on switch  
363 trials in the sucrose rinse condition, whilst the enhancing effects of sucralose rinse are only  
364 evident in the repeat nested and preswitch trials. An additional analysis examining the  
365 differences between conditions for each specific trial type (switch, nested, preswitch),  
366 revealed a significant source\*method interaction on switch trials,  $F(1,37) = 4.56$ ,  $p = .04$ .  
367 However, post hoc comparisons revealed no significant RT performance difference on switch  
368 trials between conditions (faster RT after rinsing with sucrose vs. ingesting sucrose and  
369 rinsing with sucralose  $p = .10$ ).

370

371 <TABLE 2>

372 Participants overall RT decreased over the study visits indicating a practice effect.  
373 However, no interaction with source or method suggests counterbalancing ensured this effect  
374 was equally balanced across the experimental conditions

375

376 **Accuracy.** A significant trial\*visit interaction  $F(6, 195) = 2.71, p = .02$ , main effects  
377 of trial,  $F(2, 78) = 23.72, p < .001$ , visit,  $F(3, 104) = 21.65, p < .001$ , and a marginal main  
378 effect of method,  $F(1, 39) = 3.85, p = .06$ , were revealed for attention switch accuracy (Table  
379 2).

380

381 The trial\*visit interaction reflects significantly lower accuracy on switch trials at visit  
382 1 compared to visits 2 and 3, and on preswitch trials at visit 1 compared to visits 2, 3, and 4;  
383 this is consistent with a practice effect. However, no interaction with source or method  
384 suggests counterbalancing ensured this effect was equally balanced across the experimental  
385 conditions.

386

387 Post hoc tests revealed the marginal main effect of method to be reflective of non-  
388 significantly higher accuracy during rinse compared to ingest conditions ( $p = .09$ ). Additional  
389 analyses revealed no significant differences between solution conditions for each specific  
390 trial type. It should be noted that many participants' accuracy was close to ceiling.

391

392 The main effect of trial for both attention switch RT and accuracy reflects  
393 significantly lower performance on switch vs nested and preswitch trials ( $p < .001$ ) and  
394 preswitch vs nested trials (RT:  $p < .05$ ; accuracy:  $p = .01$ ), indicating the cognitive cost of  
395 switch between task contingencies. However, this was independent of experimental  
396 conditions so does not reflect an effect of the intervention.

397

398 To permit a more direct comparison with previous studies reporting the specific  
399 facilitative effect of ingesting carbohydrates vs artificially sweetened drinks (e.g., <sup>(18)</sup>), and  
400 rinsing carbohydrates vs artificially sweetened drinks (e.g., <sup>(15)</sup>), two separate analyses  
401 comparing the ingestion of sucrose vs. sucralose and comparing the rinsing of sucrose vs.  
402 sucralose solutions were conducted.

403 **Ingest.** No significant effects of ingesting sucrose or sucralose were revealed for  
404 switch task accuracy or RT. Main effects of trial (accuracy:  $F(2,78) = 11.72, p < .001$ ; RT:  
405  $F(2,78) = 151.45, p < .001$ ) and visit (accuracy:  $F(3,34) = 10.75, p < .001$ ; RT:  $F(3,34) =$

406 12.79,  $p < .001$ ) reflected greater accuracy and RT on nested and preswitch vs switch trials,  
407 and improved performance over study visits ( $p < .05$ ).

408

409 **Rinse.** No significant effects of rinsing sucrose or sucralose were revealed for switch  
410 task accuracy or RT. Main effects of trial (accuracy:  $F(2,77) = 14.38$ ,  $p < .001$ ; RT:  $F(2,76) =$   
411  $145.63$ ,  $p < .001$ ) and visit (accuracy:  $F(3,34) = 12.82$ ,  $p < .001$ ; RT:  $F(3,34) = 5.72$ ,  $p =$   
412  $.003$ ) reflected greater accuracy and RT on nested and preswitch vs switch trials, and  
413 improved performance over study visits ( $p < .05$ ). A non-significant source (sucrose vs  
414 sucralose)\*trial interaction was revealed for RT,  $F(2,79) = 2.45$ ,  $p = .09$ . Post hoc comparison  
415 revealed no significant differences in RT performance between the rinse conditions.

416

### 417 **Subjective Response**

418 **Subjective mood (POMS).** Significant baseline\*source\*method,  $F(1,221) = 11.97$ ,  $p$   
419  $< .001$ , source\*method,  $F(1,28) = 7.43$ ,  $p = .01$ , baseline\*method,  $F(1,217) = 5.22$ ,  $p = .02$ ,  
420 interactions and an effect of baseline,  $F(1,217) = 67.98$ ,  $p < .001$ , were revealed for  
421 subjective fatigue-inertia rating. A positive relationship between baseline and post-solution  
422 overall fatigue rating was demonstrated. This positive association was strongest in the  
423 sucralose/rinse and weakest in the sucrose/ingest condition specifically at higher fatigue  
424 loads. This suggests ingestion of glucose reduced fatigue more than rinsing with sucralose in  
425 those reporting high baseline fatigue. Overall, higher subjective fatigue ratings were shown in  
426 rinse ( $\bar{x} = 4.22 \pm 0.35$ ) compared to ingest conditions ( $\bar{x} = 3.39 \pm 0.29$ ;  $p = .01$ ).  
427 Specifically, significantly higher fatigue was reported following rinsing of sucralose ( $\bar{x} =$   
428  $4.65 \pm 0.5$ ) compared to ingesting sucralose ( $\bar{x} = 3.28 \pm 0.43$ ;  $p = .03$ ). Marginally higher  
429 fatigue ratings were also reported after rinsing with sucrose ( $\bar{x} = 3.78 \pm 0.47$ ) compared to  
430 ingesting sucralose ( $p = .06$ ).

431

432 Examination of the remaining POM dimensions revealed no further significant effects of  
433 exposure to the experimental conditions on subjective mood.

434

### 435 **Solution Palatability**

436 Participants rated the sucrose solutions ( $\bar{x} = 72.55 \pm 2.05$ ) to be significantly sweeter  
437 than the sucralose ( $\bar{x} = 66.75 \pm 2.27$ ) solutions ( $p = .01$ ; main effect of source,  $F(1,39) =$   
438  $7.44$ ,  $p = .001$ ). However, solution rinsing ( $\bar{x} = 71.77 \pm 1.94$ ) was rated to be significantly

439 sweeter than ingesting ( $\bar{x} = 67.53 \pm 2.38$ ) the solutions ( $p = .05$ ; main effect of method,  
440  $F(1,39) = 3.95, p = .05$ ). This likely reflects the longer retention of solutions in the oral cavity  
441 in the rinse conditions. The sucrose solutions ( $\bar{x} = 59.66 \pm 2.91$ ) were also perceived to be  
442 more pleasant than the sucralose solutions ( $\bar{x} = 51.53 \pm 3.06; p = .01$ ; main effect of source,  
443  $F(1,39) = 9.60, p = .001$ ). Specifically, ingesting the sucrose solution ( $\bar{x} = 64.95 \pm 3.67$ ) was  
444 rated as significantly more pleasant than all other conditions ( $p < .02$ ; source\*method  
445 interaction,  $F(1,39) = 7.73, p = .01$ ). Participants also rated the ingestion ( $\bar{x} = 57.12 \pm 2.68$ )  
446 of solutions to be significantly more satisfying than rinsing ( $\bar{x} = 44.99 \pm 2.91; p = .001$ ; main  
447 effect of method,  $F(1,39) = 17.03, p = .01$ ).

448

449

### Discussion

450

451 Carbohydrates have been associated with a positive effect on depleted self-control  
452 task performance. This effect has been the subject of both metabolic and motivational  
453 explanations. The findings from the present study contribute to the current understanding of  
454 the relationship between carbohydrates and self-control in a number of ways. Firstly, the  
455 findings contradict metabolic accounts of the relationship between self-control and  
456 carbohydrate metabolism. Despite the use of high precision laboratory standard measurement  
457 of capillary blood and interstitial glucose, no consistent effects of self-control exertion on  
458 glucose metabolism were observed. A small, but significant drop in capillary blood glucose  
459 was demonstrated over the period participants completed the BAKAN depletion task.  
460 However, this constituted an average decrease of 0.15 mmol/L. This is lower than the  
461 peripheral blood glucose decrease under depletion conditions previously reported (0.33  
462 mmol/L<sup>[14]</sup>). A comparable increase (0.13 mmol/L) was also demonstrated during exposure  
463 to the initial attention switch task; therefore, this finding likely reflects variation in  
464 endogenous blood glucose levels and/or precision variability of the glucose analyser.  
465 Furthermore, self-control exertion had no effect on interstitial glucose levels which were  
466 continuously measured throughout the ego-depletion period, and can thus be considered a  
467 more robust measure of this effect. This supports previous evidence that the moderation of  
468 peripheral glucose by self-control exertion is unlikely<sup>(16, 30, 15)</sup> and contradicts evidence  
469 directly linking glucose metabolism to self-control<sup>(3, 14)</sup>. Inconsistency in the literature may  
470 stem from the use of less reliable and precise commercially available capillary blood glucose  
monitors which are not sufficiently sensitive to measure fluctuations in the euglycaemic

471 range <sup>(44, 45)</sup>. The use of a subcutaneous continuous glucose monitor also confirmed, for the  
472 first time, the lack of this effect in interstitial fluid.

473

474 The contention that glucose administration has “a large homogenous effect” on ego  
475 depletion (p. 514; <sup>(11)</sup>) was not supported. No discernible difference in self-control  
476 performance under conditions of ego depletion was found despite large concomitant increases  
477 in both blood and interstitial glucose levels. This lack of effect in older adults replicates  
478 findings demonstrated in a young adult sample <sup>(31)</sup> and contributes to growing evidence that  
479 the role of glucose as the physical mediator of self-control has been overstated <sup>(16)</sup>. The non-  
480 energetic capacity of carbohydrate sensing in the mouth to moderate self-control performance  
481 has been at the forefront of criticism of the metabolic-based resource strength model. The  
482 present study also failed to demonstrate any specific, significant effect of sensing  
483 carbohydrate in the mouth on depleted self-control performance. This contradicts previous  
484 evidence of the capacity of carbohydrate rinsing to protect performance <sup>(21, 15, 22)</sup>. However,  
485 the positive effect of carbohydrate rinsing has not been consistently reported <sup>(30)</sup>.

486

487 The act of rinsing both a caloric and an artificially sweetened solution was associated  
488 with superior performance on an attention switch task compared to ingestion of the same  
489 solutions. It is important to note that improved attention switch performance after rinsing the  
490 solutions was marginal ( $p = .05$ ) and very small (RT saving 17 ms; Cohen’s  $d = .1$ ). This  
491 effect is certainly not as robust as previous demonstrations of the facilitative effects of rinsing  
492 (e.g.,  $d = .63, .73$  <sup>(15)</sup>). The effect of rinsing on accuracy was marginal and likely influenced  
493 by ceiling effects. However, a positive effect of rinsing in general on cognitive performance  
494 is difficult to assimilate with existing evidence of the effect of rinsing carbohydrate and  
495 artificially sweetened solutions. The potential effect of ingesting the solutions having a  
496 greater fatiguing effect than rinsing, due to the digestive processes required, is contradicted  
497 by higher subjective levels of fatigue in rinse conditions, and evidence of a specific  
498 restorative effect of sucrose in those reporting higher fatigue at baseline. Previous evidence of  
499 the positive effects of rinsing has been attributed to the specific sensing of energy in the oral  
500 cavity. Oral energy sensing of carbohydrate may stimulate activity in neural regions  
501 associated with exertion of self-control (namely the anterior cingulate cortex [ACC]) thus  
502 countering reduced activity when depleted. Oral sensing may also activate neural reward  
503 pathways shifting the motivational state towards persistence <sup>(24, 15)</sup>. This proposed mechanism  
504 cannot easily explain the observation of general beneficial effects of rinsing with both

505 sucrose and sucralose that were observed in the current study. fMRI studies have located  
506 divergent neural activation for caloric and artificially sweetened oral solutions. For example,  
507 stronger activation of the primary taste areas (anterior insula and frontal operculum),  
508 prefrontal cortex, and reward circuit regions (striatum) is evident after tasting sucrose  
509 compared to an artificially sweetened solution <sup>(24, 47)</sup>. Divergent activation by caloric and  
510 artificially sweetened oral solutions has also been reported in the amygdala, median  
511 cingulate, thalamus and pre-central gyrus <sup>(48, 49)</sup>.

512

513 However, the proposed mechanisms of oral sensing remain speculative and  
514 unverified, and neural areas activated by oral carbohydrate sensing are rather heterogeneous.  
515 Furthermore, a divergent response across energy content has not been consistently shown.  
516 van Rijn et al. <sup>(50)</sup> reported no overall difference in neural activation for caloric (maltodextrin  
517 and maltodextrin + sucralose) and sucralose solutions. Activation of reward circuit areas  
518 (striatum) after tasting caloric and non/low-caloric solutions (including water) were also  
519 indistinguishable. Divergent neural activation was only demonstrated when hunger state was  
520 taken into account. Greater activation of the median cingulate, thalamus, anterior insula and  
521 inferior frontal gyrus, was revealed after tasting a caloric solution in a satiated compared to  
522 hungry state. Thus energy sensing may be a hunger state dependent process. This emphasises  
523 that the mechanisms assumed to underpin the effects of oral energy sensing on self-control  
524 are yet to be fully elucidated.

525

526 Lange and Eggert <sup>(30)</sup> attempted to further clarify how the proposed rewarding effect  
527 of oral sensing acts to moderate self-control by manipulating the temporal relationship  
528 between ingestion and rinsing of carbohydrate and self-control tasks. Oral sensing of  
529 carbohydrate may operate via (i) retrospectively rewarding initial self-control exertion if  
530 administered after the initial depletory task, or (ii) prospectively indicating the availability of  
531 future resources signalling ‘affordable’ continued exertion of self-control if administered  
532 prior to the secondary ego depleted task <sup>(30)</sup>. No effect of rinsing, or ingesting, carbohydrates  
533 on self-control were observed. Such findings are incompatible with the assumed rewarding  
534 effect of oral sensing. Psychological manipulations (e.g., thinking about a cherished value  
535 <sup>(29)</sup>) have also been shown to attenuate depleted self-control performance. Mere perception of  
536 feeling less (versus more) depleted, irrespective of actual level of depletion, also results in  
537 better self-control performance <sup>(51)</sup>. Therefore, self-control capacity appears sensitive to  
538 multiple factors independent of metabolism or the rewarding properties of energy sensing.

539

540 Findings demonstrate that RT in general, i.e., independent of trial type, was faster  
541 after rinsing, independent of carbohydrate content. The exertion of self-control on this task  
542 specifically relates to the effortful suppression of the dominant response to repeated trials  
543 when responding to switch trials. It is therefore debatable if a facilitation of RT, independent  
544 of trial, is indicative of a clear effect on self-control performance. A facilitative effect on self-  
545 control performance would be more clearly indexed by faster RT or greater accuracy on  
546 switch trials compared to repeat trials. However, across all conditions, performance on  
547 preswitch trial was lower than nested trials. This performance deficit on preswitch trials is  
548 likely reflective of a preparatory response or inhibition of a premature response to the  
549 impending task switch. Therefore, whilst performance on the switch trials relative to repeated  
550 trials is the primary indicator of self-control performance, performance across the task in  
551 general can still be considered a measure of self-control. Comparisons of performance on  
552 each trial type between conditions did reveal a trend for faster RT on switch trials after  
553 rinsing sucrose ( $p = .10$ ); enhancement after sucralose rinsing was confined to repeat trials.  
554 This provides only very limited support for the non-metabolic enhancing effect of oral sensing  
555 carbohydrates on depleted self-control performance.

556

557 The strengths of the reported study lie in the robust methodology adopted. Highly  
558 precise methods of assessing the metabolic effects of ingesting and rinsing carbohydrates and  
559 a rigorous standardised, repeated measures, counterbalanced crossover design were applied.  
560 The cognitive tasks employed to deplete and measure self-control performance were also  
561 well-established and validated measures of performance. This contrasts with previous studies  
562 that have adopted less standardised methods of depleting self-control (e.g., writing about  
563 one's death <sup>(3)</sup>; a high score on a questionnaire assessing attitudes towards homosexuality  
564 <sup>(18)</sup>). To detect an effect of carbohydrate ingestion as large as that reported by Hagger et al.  
565 <sup>(11)</sup> ( $d = .75$ ) or Gailliot et al. <sup>(11)</sup> ( $d = .55$ ), sample sizes of  $n = 16$  and  $n = 28$  would be  
566 required respectively using the employed design. Similarly, a sample size of  $n = 22$  would be  
567 required to detect an effect of rinsing carbohydrate of the magnitude reported by Molden et  
568 al. <sup>(15)</sup> ( $d = .63$ ; all  $\alpha = .05$ ,  $1 - \beta = .8$ ). Therefore, the lack of convincing effects in this sample  
569 of 40 older adults is unlikely to be due to an insufficient level of power. However, a number  
570 of additional factors are worthy of discussion. Different self-control tasks were employed to  
571 deplete and assess ego depleted performance in the present study. The use of differing tasks  
572 should not be expected to unduly influence the findings considering the assumption that self-

573 control capacity is mediated by a common, modality independent, resource <sup>(11)</sup>. Furthermore,  
574 the positive effects of carbohydrates on self-control performance, and failed attempts to  
575 replicate the effects, have been demonstrated using the same <sup>(20, 31)</sup> and dissimilar <sup>(21, 30)</sup> tasks.

576

577 In this study, the placebo solutions were sweetened with sucralose which, whilst  
578 sufficiently low in energy to be considered non-caloric (< 4 kcal in study solutions), contains  
579 a small amount of maltodextrin filler (< 1 g in study solutions). The majority of studies  
580 examining the effects of rinsing carbohydrates have employed artificial sweeteners  
581 completely free from carbohydrates (e.g., aspartame <sup>(15, 21)</sup>). Therefore, it could be argued that  
582 the facilitative effects of rinsing with both sucrose and sucralose may be indicative of an  
583 enhancing effect of carbohydrate even at very low levels. However, specific enhancing  
584 effects of rinsing glucose but not sucralose have been reported <sup>(22-46)</sup> which contradicts this  
585 proposition.

586

587 Evidence that greater initial exertion on multiple tasks of self-control does not  
588 inevitably result in more severe subsequent performance decrement – as may be assumed by  
589 the self-control resource model – has been reported <sup>(46)</sup>. Indeed, the inclusion of multiple  
590 initial depletory self-control tasks has been shown to increase performance <sup>(52)</sup>. Therefore, the  
591 initial pre-solution intake exposure to the attention switch task to standardise baseline self-  
592 control resources may have influenced subsequent self-control outcomes. However, this  
593 would not explain why improved performance was only demonstrated in the rinse conditions  
594 since all visits were strictly standardised and counterbalanced.

595

596 The recruitment of an older adult sample was informed by evidence of greater  
597 potential efficacy of glucose administration in populations with increased likelihood of  
598 disrupted metabolic or cognitive activity (e.g., <sup>(52, 33, 35)</sup>). It is however acknowledged that ego  
599 depletion deficits may not be as potent in older versus younger samples (more evident in < 25  
600 vs 40 – 65 year olds <sup>(53)</sup>).

601

602 Finally, participants reported a discernible taste perception difference between the  
603 experimental solutions which may be argued to have unduly influenced the results. However,  
604 the inclusion of perceived sweetness, satisfaction and pleasantness rating in statistical models  
605 did not significantly affect the outcome variables and modulation of performance was  
606 observed independently of source.

607

608

609 **Conclusions**

610 Findings contribute to recent evidence casting doubts upon a specific role of  
611 carbohydrate in self-control performance. Exertion of self-control did not moderate blood or  
612 interstitial glucose, nor did metabolism or oral sensing of sucrose significantly affect self-  
613 control performance in the ego depleted state. In sum, findings do not support the proposition  
614 that carbohydrate is vital for self-control in older adults and suggests that the role of glucose  
615 as the physical mediator of self-control has been overstated.

616

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620

621 **Conflict of Interest**

622 On behalf of all authors, the corresponding author states that there is no conflict of interest.  
623 The authors have full control of all primary data and will permit access to review if  
624 requested. The funding agent provided editorial assistance in the completion of the draft for  
625 publication (KN). The original concept, review of literature, and study design was conceived  
626 by LD and CL. The collection (with assistance of FC and KS), analysis and interpretation of  
627 the data were completed by the corresponding author (NB). The corresponding author  
628 completed the draft for publication with editorial assistance from LD, CL, and RA.

629

630

631 **TABLES**

632

633 Table 1 Solution compositions and nutritional content

<b>Solution</b>	<b>Serving</b>	<b>Solution Composition</b>	<b>Sugar Content</b>	<b>Kcal</b>
Sucrose	381.4g	125.9g cordial + 255.5g water	50g	200
Sucralose	381.4g	63.5g cordial + 317.5g water	0.92g	3.7

634

635 Table 2 Descriptive statistics of attention switch task performance outcomes according to experimental condition\*trial and total  
 636 performance\*method of solution intake (ingest vs. rinse). Mean  $\pm$  SEM are shown

Experimental Condition	Reaction Time (ms)						Accuracy					
	Trial						Trial					
	Switch		Nested		Preswitch		Switch		Nested		Preswitch	
	<i>X</i>	<i>SEM</i>	<i>X</i>	<i>SEM</i>	<i>X</i>	<i>SEM</i>	<i>X</i>	<i>SEM</i>	<i>X</i>	<i>SEM</i>	<i>X</i>	<i>SEM</i>
Sucrose/ingest	1154.45	19.49	931.30	20.90	950.71	20.81	21.33	0.37	22.56	0.26	22.21	0.33
Sucrose/rinse	1110.18	18.72	910.97	21.78	947.27	24.34	21.67	0.37	23.05	0.18	22.42	0.29
Sucralose/ingest	1148.35	25.34	931.16	23.09	944.94	22.55	21.50	0.47	22.15	0.43	21.49	0.50
Sucralose/rinse	1157.16	20.93	915.80	22.87	933.54	20.09	<b>Total</b>					
							<i>X</i>	<i>SEM</i>			<i>X</i>	<i>SEM</i>
Ingest							1010.49	10.91			21.87	0.16
Rinse							993.49	10.83**			22.12	0.15*

637 \*\*  $p = .05$  \*  $p = .09$

638

639

640 **Figure 1** Attention switch task stimulus configuration showing seven consecutive trials.

641 Participants were required to make a Go/noGo response if the letter shown was a vowel or  
642 consonant (red coloured stimuli), and if the number shown was odd or even (green coloured  
643 stimuli). The task run switched between the two categorisation Go/noGo tasks every third  
644 trial. Figure adapted with permission from Wylie, G. R., Javitt, D. C., & Foxe, J. J. (2003).  
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646

647 **Figure 2** Procedural timeline

648

649 **Figure 3** Mean capillary blood glucose response (mmol/L) according to experimental  
650 condition

651

652 **Figure 4** Mean interstitial glucose response (mmol/L) according to experimental condition.

653 Exposure to the experimental solutions is centred at 0 minutes which represents time of  
654 solution intake

655

656 **Figure 5** Mean Bakan target accuracy (A) and RT (B) across each one minute trial block  
657 according to experimental condition

658

659

660

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