

1 16/3/16

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3 **Chemosensory abilities in consumers of a Western-style diet**

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Abstract

People vary in their habitual diet and also in their chemosensory abilities. In this study we examined whether consumption of a Western-style diet, rich in saturated fat and added sugar, is associated with either poorer or different patterns of chemosensory perception, relative to people who consume a healthier diet. Participants were selected based on a food frequency questionnaire, which established whether they were likely to consume a diet either higher or lower in saturated fat and added sugar. Eighty-seven participants were tested for olfactory ability (threshold, discrimination, identification), gustatory ability (PROP sensitivity, taste intensity, quality and hedonics), and flavour processing (using dairy fat-sugar-odour mixtures). A Western-style diet was associated with poorer odour identification ability, greater PROP sensitivity, poorer fat discrimination, different patterns of sweetness taste enhancement, and hedonic differences in taste and flavour perception. No differences were evident for odour discrimination or threshold, in perception of taste intensity/quality (excluding PROP) or the ability of fats to affect flavour perception. The significant relationships were of small to moderate effect size, and would be expected to work against consuming a healthier diet. The discussion focuses on whether these diet-related differences precede adoption of a Western-style diet and/or are a consequence of it.

Keywords: Olfaction, Gustation, Flavour, Diet, Food choice

Introduction

48
49 Although people change what they eat, the idea that they can maintain broadly stable
50 patterns of food intake over the long term (e.g., years to decades) has been confirmed in several
51 studies (e.g., Pachuki, 2012; Newby, 2006). Routinely eating diets rich in saturated fat, and
52 added sugar - a Western-style diet - in contrast to a plant-based diet, is associated with poorer
53 health outcomes (e.g., Appel et al., 1996; Mente et al., 2009). This makes it important to study
54 the drivers of dietary choice. While many factors impact dietary choice (e.g., education,
55 poverty, impulsivity), one potentially important factor is chemosensory ability. This is
56 relatively unexplored, with most focus to date on alcohol consumption (e.g., Bachmanov,
57 2003) and on genetically based sensitivity to bitter tasting propylthiouracil (PROP; Duffy et
58 al., 2009; Feeney, 2011; Hayes et al., 2010; Hayes et al., 2013). Far less attention has been
59 paid to how variation in olfactory abilities, taste perception beyond PROP sensitivity, and the
60 integration of taste, olfaction and somatosensation into flavour, may relate to diet. Here, we
61 examine whether and how variation in chemosensory ability, for olfaction, taste (including
62 PROP) and flavour, is linked to consumption of a Western-style diet.

63 Dietary preferences may be influenced by pre-existing perceptual differences. In the
64 context of a Western-style diet, we hypothesise that one such difference may be a poorer sense
65 of smell. If a person's olfactory ability is poor, this may be compensated for by choosing
66 foods that involve greater stimulation of the taste and oral somatosensory systems (i.e., irritant
67 and fat perception). While there is no direct data as yet regarding habitual diet and olfactory
68 ability in general populations, there is circumstantial evidence favouring a link. One line of
69 evidence comes from data we have collected. Some participants who had completed the
70 Sniffin sticks (Hummel et al., 1997) odour discrimination and identification tests as part of one
71 study in our laboratory, also provided dietary data as part of another. There were significant
72 associations between their olfactory ability and reported diet, with a Western-style diet
73 associated with poorer discrimination ($n = 86$; $r = -0.25$) and identification ($n = 86$; $r = -0.22$).

74 There are two further reasons to think that poorer olfactory ability may be associated
75 with an inferior diet. First, obese individuals show preferences for sweeter (e.g., Drewnowski
76 et al., 1985) and especially fattier (Cox et al., 2016) stimuli, and some studies find olfactory
77 impairments among the obese (e.g., Richardson et al., 2004; but see Stafford & Whittle, 2015).
78 Olfactory impairments could of course be caused by obesity, rather than predating it, but these
79 findings are at least consistent with the view that poorer olfaction might result in weight gain.
80 Second, people with an impaired sense of smell, including the elderly (Duffy & Hayes, 2014),
81 often report dietary alterations. These may include adding more sugar, increasing spice use,
82 and reducing plant-based foods (Merkonidis et al., 2015; Miwa et al., 2001; Van Toller, 1999)
83 - although the impact of impaired olfaction on body weight is variable (e.g., Ferris & Duffy,
84 1989). Here, we tested whether performance on standardised tests of olfaction (Sniffin Sticks;
85 Hummel et al., 1997) would be poorer in consumers of a Western-style diet.

86 We also examined whether a Western-style diet would be related to variation in taste
87 perception. Two hypotheses can be advanced here. First, as a Western-style diet involves
88 more processed food, and less fruit and vegetables, this would suggest the possibility of greater
89 sensitivity to bitter tasting PROP. This is premised upon the association between sensitivity to
90 genetically determined bitter taste perception ability and dietary preferences for cruciferous
91 vegetables (Feeney, 2011; Hayes et al., 2013). Second, as processed foods contain higher
92 levels of sugar, salt and fat than non-processed food, exposure to them might increase
93 preference for more concentrated forms of these tastants (with fat mentioned here simply as it
94 seems to be perceived via multiple sensory channels; Frost & Janhoj, 2007). In contrast,
95 reduced intake of fruits and vegetables, might be associated with reduced exposure to sour and
96 bitter tastes, and hence (e.g., via mere exposure) reduced preference for such tastants. That
97 dietary exposure can selectively affect taste preference has been shown in several studies (e.g.,
98 Bertino et al 1982; Bertino et al., 1986) and in naturalistic studies of populations who consume
99 different diets (e.g., bitterness/sourness; Moskowitz et al., 1975). While preference changes

100 seem well supported taste intensity perception appears to be stable and largely independent of
101 dietary exposure (e.g., Cicerale et al., 2012; Mattes, 1985; Moskowitz et al., 1975), thus no
102 relationship with habitual diet would be expected here.

103 A third set of predictions relate to flavour. Flavour perception, in the context of this
104 study, has two main attributes. The first concerns sensory interactions in the mouth. These
105 come in two forms: (1) the ability of certain tastants and odourants to affect odour and taste
106 quality perception such that sweet tastants can enhance the intensity of previously co-
107 experienced odours (e.g., Von Sydow et al., 1974), and that previously co-experienced odours
108 can enhance the intensity of sweet tastants (e.g., Frank & Byram, 1988); and (2) the ability of
109 fats to suppress the intensity of odourants. Both of these effects have a well-established
110 psychological basis (Bult, de Wijk & Hummel, 2007; Sakai et al., 2001) and both could either
111 be influenced by or influence habitual diet (e.g., poorer odour-taste integration resulting in less
112 sweetness taste enhancement could be associated with a preference for more added sugar).
113 Several possible effects are plausible, but the absence of any prior data precludes direction-
114 specific predictions. The second main flavour attribute is hedonics (Stevenson, 2009). Not
115 only may liking for different flavour combinations differ depending upon prior dietary
116 exposure but capacity to integrate the sensory dimensions of flavour could also affect liking.

117 To examine whether participants who differ in adherence to a Western-style diet also
118 differ in their olfactory, taste and flavour perception abilities, we recruited people differing on
119 this dietary dimension. Western-style dietary intake was established using a reliable and
120 validated brief food frequency scale (Dietary Fat and Sugar questionnaire; DFS; Francis &
121 Stevenson, 2013). While we used standardised tests for olfactory ability and PROP taste
122 intensity – as these are well established commercially (for Sniffin sticks) and in the literature
123 (for both; e.g., Hummel et al., 1997; Kirkmeyer & Tepper, 2005; Prescott, Ripandelli &
124 Wakeling, 2001), we employed bespoke tests to examine taste and flavour perception. For
125 taste, participants evaluated sweet and salty tastes at two superthreshold concentrations, along

126 with the PROP sample and a sour taste. For flavour, we loosely modelled our design on the
127 Drewnowski et al., (1985) approach, providing participants with dairy samples that varied in
128 fat, sugar and fruit odourant concentration. For both the tastes and flavour samples,
129 participants evaluated their intensity, qualities and hedonics using labelled magnitude scales
130 (Green et al., 1996; Lim, Wood & Green, 2009). Finally, we collected basic demographic,
131 medical chemosensory information and body mass index (BMI), to check that these variables
132 were not responsible for any observed dietary associates of chemosensory ability.

133

134 Method

135 Participants

136 Participants were recruited in two ways. The first involved asking people who had
137 completed an earlier study looking at the relationship between diet and memory if they wished
138 to take part in a further (i.e., this) study. Originally, all participants in the earlier study had
139 completed the Dietary Fat and Sugar questionnaire (DFS; Francis & Stevenson, 2013) as part
140 of a screening program to identify individuals who differed maximally in saturated fat and
141 added sugar intake, so as to ensure a wide spread of DFS scores. Participants who had a DFS
142 score above 70 or below 55 (scores on the DFS can range from 26 to 130), who reported a BMI
143 between 17 and 26 (broader because this was a self-estimate and included people of both
144 Caucasian and Asian descent), were aged between 17 and 35, who had consented to be
145 approached, and passed the medical screening (described below), formed the pool from which
146 the earlier diet and memory study were recruited – and from which 60 people here were
147 recruited.

148 The second recruitment method drew upon the university community. Using
149 advertisements posted around campus, interested parties were invited to phone the study team
150 about participation. On phoning they were asked about their frequency of consumption for the
151 seven items from the DFS that have the highest item-total correlations (Soft drinks; Cakes &

152 Cookies; Pizza; Fried chicken, or chicken burgers; Doughnuts, pastries, croissants; Corn chips,
153 potato chips, popcorn with butter; French fries, fried potatoes). Participants aged 17-35, with a
154 BMI between 17-26, and who scored below 16 or above 21 on this short-form of the DFS were
155 potentially eligible to take part. Potential participants also received a medical screening
156 interview, which was also successfully completed by participants in our earlier diet and
157 memory study. This assessed current (physical or mental illness; chronic conditions; recent
158 hospitalisations; any history of eating disorders; any head injuries; food allergies), and past
159 health issues. Participants who reported anything other than minor health complaints were
160 excluded. Using this advertisement route, 28 participants were recruited.

161 Eligible participants were instructed to breakfast or lunch as per normal, but to refrain
162 from eating (and smoking if they were a smoker) in the 2 hours before testing. Participants
163 were also told that they could drink water in this period but not caloric beverages and not to
164 exercise beyond their normal pattern.

165 In total 88 participants completed the study. Data from one participant was excluded as
166 they were unable to breathe properly due to nasal congestion during testing. The same pattern
167 of significant findings obtains when this participant's available data are included.

168

169 Materials

170 Olfactory testing: This was undertaken using the Sniffin-sticks test battery (Hummel et
171 al., 1997), which involves a 16-item odour identification test, a 16-item odour triad
172 discrimination test, and a butanol odour threshold test (Burghart Medizintechnik, Germany).

173 Gustatory testing; Seven test solutions were prepared: 0.17M ('strong salt') and 0.03M
174 ('weak salt') saline solutions, plus a 0.1M saline standard; 0.36M ('strong sugar') and 0.03M
175 ('weak sugar') sucrose solutions; a 0.04M citric acid (Sigma-Aldrich, Sydney, Australia)
176 solution; and a 0.32mM PROP (Sigma-Aldrich, Sydney, Australia) solution - this
177 concentration and that of the preceding saline standard were based upon prior studies (Prescott,

178 Ripandelli & Wakeling, 2001; Kirkmeyer & Tepper, 2005). 10ml of each tastant were
179 presented in disposable 30ml sample cups.

180 Flavour testing: Factorial combinations of three components were used to make the 18
181 samples used for flavour testing. These components were full cream milk (3.4% fat, 3.4%
182 protein, 4.9% carbohydrates) and skimmed milk (0.1% fat, 3.4% protein, 4.9% carbohydrates),
183 three levels of added sucrose, and three levels of added cherry odourant (Givaudan). The final
184 percentages of fat, added sucrose (noting that both milks contain the same base level of
185 naturally occurring sugars) and cherry flavourant for the samples were: (1) all 9 skimmed milk
186 samples contained 0.08% fat, and all 9 full fat milk samples contained 2.8% fat; (2) the 6
187 samples with lowest level of added sugar contained 0.5% w/v (0.015M), the 6 samples with an
188 intermediate level of added sugar contained 3.8% w/v (0.11M), and the 6 samples with the
189 highest level of added sugar contained 11.7% w/v (0.34M); and (3) for the cherry odourant, 6
190 samples contained no cherry odourant, 6 contained 1.3×10^{-4} % w/v, and 6 contained 4.2×10^{-4} %
191 w/v. All samples were of 10ml presented in disposable 30ml sample cups.

192 DFS (diet) questionnaire: This 26 item food frequency questionnaire was developed
193 specifically to detect variation in saturated fat and added sugar intake. The questionnaire has
194 established reliability and validity (see Francis & Stevenson, 2013, for details). To summarise,
195 dietary intakes of saturated fat and added sugar obtained from an extensive Australian (CSIRO)
196 food frequency question, and from a 4-day Medical Research Council diet diary, both
197 significantly correlated with DFS (diet) scores (r 's from 0.36 to 0.71), indicating that higher
198 scores on the DFS equate to higher intakes of saturated fat and added sugar. More recently, we
199 have shown that a skin spectrophotometry estimate of subcutaneous carotenoid levels, which
200 are primarily derived from fruit and vegetable intake, is significantly and negatively associated
201 with DFS (diet) score ($r = -0.21$; Attuquayefio et al., Submitted), indicating that dietary fruit
202 and vegetable intake is lower in individuals scoring higher on the DFS. The DFS is also
203 reliable, even over fairly extended intervals (22 weeks; $r = 0.84$).

204 Procedure

205 The study protocol was approved by Macquarie University Human Research Ethics
206 Committee and informed consent was provided by all participants. After consenting,
207 participants completed three questionnaires: (1) a biographical questionnaire to obtain age,
208 gender, along with general health questions and ones pertaining to adherence to the pre-
209 experimental instructions; (2) a chemosensory health-screening questionnaire, developed in our
210 lab to identify potential olfactory impairments (history relating to allergies, sinusitis, facial
211 surgery, facial injury, current or recent respiratory infections, current or recent nasal
212 congestion, any past or current problems with taste or smell, current and past smoking history,
213 any history of head injury, and any previous periods of unconsciousness/concussion); (3) a
214 current-state questionnaire, with rating of hunger, thirst, fullness, happiness, sadness,
215 relaxedness and alertness - in that order - on 120mm line scales (anchors Not at all and Very).

216 Participants then completed, in this order, the following study tasks: (1) Threshold
217 testing with the Sniffin Sticks; (2) Discrimination testing with the Sniffin Sticks; (3) Odour
218 identification testing with the Sniffin Sticks; (4) Gustatory testing; (5) Flavour testing; and (6)
219 Final study measurements. Each is described in more detail below.

220 Sniffin-sticks test battery: Threshold testing was conducted as per the manual, in all but
221 one regard, as only five sets of reversals were employed. The 16-odour triad discrimination
222 task (using a forced choice oddity [triangle] test) and the 16-item odour identification task
223 (using a four response option forced choice procedure) were both conducted as per the manual.

224 Gustatory testing: On this test participants sampled and evaluated seven tastants. Five
225 of these were presented in random order (strong and weak sucrose and saline solutions, and
226 citric acid solution), with the salt standard always being the penultimate sample, and the PROP
227 sample being the last to be evaluated. Following instructions on scale usage, participants were
228 asked to pour the whole of the first sample into their mouth and then expectorate, and
229 immediately complete the six evaluations. Evaluations were made on 12cm labelled

230 magnitude scales (based upon Green et al., 1996, and using the same ratios for the anchors) for
231 intensity, sweetness, bitterness, saltiness, sourness and hedonics (in this last case using a
232 bipolar labelled magnitude scale, based upon Lim, Wood & Green, 2009). Participants then
233 rinsed with water, waited 5s, and then commenced the next trial, repeating this procedure for
234 all of the remaining tastants.

235 Flavour testing: On this test participants sampled and evaluated 18 solutions presented
236 in randomised order. The same pour-into-the-mouth, sample, spit, rate and rinse procedure
237 was used here as for the gustatory testing. However, participants made a slightly different set
238 of evaluations here, rating intensity, sweetness, fattiness, fruitiness and hedonics, again using
239 labelled magnitude scales.

240 Final study measures: All participants completed the DFS (diet) scale so as to obtain
241 the most recent estimate of their use of a Western-style diet, with this estimate being used in
242 the analysis. Height and weight were measured to calculate body mass index (BMI).

243

244 Analysis

245 The DFS (diet) score obtained at the end of testing formed the key dietary measure. Its
246 26 item scores were summed for each participant. This score was then used as a continuous
247 variable in all of the analyses, rather than as a grouping variable (i.e., high vs. low). Treating
248 this score as a continuous variable is more powerful than using it to form groups, as it utilises
249 all of the information in the dietary measure (Preacher et al., 2005).

250 Threshold scores were the mean of the final four reversals (higher numbers indicate
251 greater sensitivity), and discrimination and identification scores were the number of correct
252 responses out of 16.

253 For measures using the labelled magnitude scale scores, we found these to more closely
254 approximate to a normal distribution than when a log-transformation was applied and so these
255 data were analysed without further transformation. Three types of score were assembled for

256 the Gustatory tests: Intensity – from the intensity rating in mm along the labelled magnitude
257 scale; Hedonics, with scores in mm on the hedonic scale being positively signed for liked
258 responses and negatively signed for disliked responses; and Quality – the nominal taste quality
259 (e.g., sweetness for sucrose) minus the mean taste quality scores for the remaining qualities (in
260 the case of sweetness - sourness, saltiness and bitterness). This latter approach to analysing the
261 taste quality ratings is both sensitive to impairments in taste processing (e.g., Stevenson, Miller
262 & McGrillen, 2013) and reduces the number of comparisons necessary to analyse these types
263 of data.

264 For the Flavour test, all of the ratings were analysed separately, because of interest in
265 certain rating scale specific effects (i.e., odour/taste enhancement and fat suppression). A
266 chemosensory problem score was also calculated, based upon responses to the 12-item
267 chemosensory health-screening questionnaire (scores could vary between 0 and 12, with 1
268 being given for each response indicative of possible impairment and 0.5 for unsure responses).

269 Two main analysis approaches were used. The first involved descriptive statistics and
270 then zero order correlations between DFS (diet) score and particular variables of interest. This
271 approach was then followed up in cases where significant relationships with DFS (diet) score
272 emerged, by a further correlation in which the effects of age, gender, BMI and chemosensory
273 problem score (and certain other variables as identified in the text) were partialled out. The
274 second approach was to use ANCOVA (dependent variables being intensity, quality and
275 hedonic ratings, and independent variables being stimulus concentration and/or type), notably
276 on the flavour and taste data. In these analyses DFS (diet) score (transformed into a Z-score)
277 was included as the covariate, which can be thought of as being akin to a continuous between-
278 participant independent variable. This allowed us to test for any heterogeneity in relationships
279 between the covariate and the factors included in each analysis, as well as establishing any
280 relationship between the covariate and the grand mean. This approach was also used in
281 conjunction with the first, to explore particular effects identified a priori as being of interest

282 (i.e., taste and odour enhancement, fat odour suppression effects). For succinctness, the
283 ANCOVAs are reported in summary form, except where DFS-related effects emerged.
284 Finally, alpha was set at 0.05, with 1-tailed tests for a priori directional hypotheses and with
285 Bonferroni adjusted alpha's for multiple comparisons as described in the Results section.

286

287 Results

288 There were no correlations between age, gender, BMI and DFS (diet) score – see Table
289 1 for details. While the chemosensory problem score (see Table 1) correlation was non-
290 significant ($p = .095$), it would appear that there is a slightly heightened rate of factors
291 associated with impaired chemosensory function in more frequent consumers of a Western-
292 style diet (i.e., a higher DFS score). There were no significant correlations between hunger,
293 thirst, fullness or mood ratings and DFS (diet) score, suggesting similarity in state at testing. A
294 summary of the key findings, including variance accounted for, is presented in Table 2.

295

296 Odour testing

297 Threshold: Mean butanol threshold was 7.2 (SD = 2.9), with no observed relationship
298 with DFS (diet) score.

299 Discrimination: Mean odour discrimination score was 11.5 (SD = 2.0), with no
300 observed correlation with DFS (diet) score.

301 Identification: Mean identification score was 11.5 (SD = 2.5), and there was a
302 significant correlation between this variable and DFS (diet) score ($r = -0.20$, $p < 0.05$, 1-tailed).
303 This relationship was not attenuated when age, gender, BMI, chemosensory problem score,
304 threshold score and discrimination score, were partialled out ($r = -0.24$, $p < 0.05$, 1-tailed).
305 This suggests that greater reported consumption of a Western-style diet is associated with
306 poorer odour identification ability, confirming earlier preliminary findings for identification.

307

308 Gustatory testing

309 PROP: There was a significant correlation between PROP intensity ratings ($M = 57.1$;
310 $SD = 37.5$) and DFS (diet) score ($r = 0.20$, $p < 0.05$, 1-tailed). This relationship was not
311 attenuated when partialling out the effects of age, gender, BMI and chemosensory problem
312 score ($r = 0.20$, $p < 0.05$, 1-tailed). This suggests that greater consumption of a Western-style
313 diet is associated with greater sensitivity to bitter tasting PROP.

314 There was also a significant negative correlation between hedonic ratings for PROP (M
315 $= -25.9$; $SD = 21.0$) and DFS (diet) score ($r = -0.21$, $p < 0.05$, 1-tailed). This relationship was
316 not attenuated when partialling out the effects of age, gender, BMI and chemosensory problem
317 score ($r = -0.19$, $p < 0.05$, 1-tailed). Complimentary to the findings above, greater
318 consumption of a Western-style diet is associated with greater dislike for PROP.

319 Finally, there was no association between taste quality score (i.e., bitter rating minus
320 the mean of the other taste quality ratings) for PROP ($M = 53.0$; $SD = 38.8$) and DFS (diet)
321 score, indicating that all participants were readily able to discern its principal taste quality.

322 Citric acid: There were no significant correlations between DFS (diet) score and citric
323 acid intensity, quality or hedonic ratings.

324 Sucrose and saline – Intensity: ANCOVA revealed no effects including DFS (diet)
325 score.

326 Sucrose and saline – Quality: ANCOVA revealed no effects including DFS (diet)
327 score.

328 Sucrose and saline – Hedonics: A two-way repeated measures ANCOVA, with Tastant
329 (sucrose vs. saline) and Concentration (strong vs. weak), and DFS (diet) score entered as a
330 covariate, revealed main effects of Concentration and Tastant, and a Tastant by Concentration
331 interaction. There was also an interaction between DFS (diet) score and Tastant ($F(1,85) =$
332 6.09 , $p < 0.02$, partial eta-squared = 0.07), indicating that DFS (diet) score correlated with one
333 of the tastant variables significantly more so than with another. DFS (diet) score was

334 significantly correlated with hedonic ratings for salt solutions (collapsed across Concentration;
335 $r = -.26$, $p < 0.02$; accounting for 6.8% of the variance in the dietary measure), but not with
336 sucrose solutions (collapsed across Concentrations; $r = 0.02$). The relationship between diet
337 and salt solution hedonic ratings was maintained even when partialling out age, gender, BMI
338 and chemosensory problem score, indicating that relatively neutral ratings were provided by
339 participants with a low DFS (diet) score, while those who consumed diets richer in saturated
340 fat and added sugar were more negative in their evaluation.

341

342 Flavour testing

343 Intensity: ANCOVA revealed no effects involving DFS (diet) score.

344 Sweetness, and sweetness enhancement effects: ANCOVA revealed no effects
345 involving DFS (diet) score.

346 We then calculated mean linear and quadratic slope coefficients across the three levels
347 of factor Odour, collapsing across Sugar and Fat levels, to determine if the pattern of sweetness
348 enhancement was related to DFS (diet) score. While there was no association with the linear
349 coefficient, the relationship with the quadratic coefficient was significant ($r = -0.21$, $p < 0.05$).
350 Participants with a healthier diet tended to have positively signed quadratic functions, with
351 degree of sweetness enhancement increasing most between the low and high levels of factor
352 Odour. In contrast, participants with a more Western-style diet tended to have negatively
353 signed quadratic functions, with maximal taste enhancement for the lower odour level and
354 minimal enhancement for the higher level. This correlation was attenuated when partialling
355 out the effects of age, gender, BMI and chemosensory problem score ($r = -0.21$, $p = 0.059$).

356 Fattiness: The fattiness data were analysed with a three-way repeated measures
357 ANCOVA, with Fat level (skimmed vs. full fat milk), Sugar level (low vs. medium vs. high)
358 and Odourant level (zero vs. low vs. high concentration) as within factors and DFS (diet) score
359 as the covariate. There were main effects of Sugar level and Fat level, with the Fat level effect

360 being moderated by DFS (diet) score ($F(1,85) = 4.58, p < 0.05, \text{partial eta-squared} = 0.05$). To
361 examine this diet-related effect, we subtracted the mean fattiness rating of all of the skimmed
362 milk samples ($M = 35.2$) from the mean fattiness rating of all of the full fat milk samples ($M =$
363 40.9). There was a significant correlation between DFS (diet) score and this fattiness
364 difference score ($r = -.29, p < 0.01$). Poorer discrimination of the two fat levels in terms of a
365 smaller difference score, was reported by participants who habitually ate a diet rich in saturated
366 fat and added sugar. This relationship was not attenuated by partialling out the effects of age,
367 gender, BMI and chemosensory problem score ($r = -0.28, p < 0.01$).

368 Fruity odour ratings, fat suppression and flavour enhancement: ANCOVA revealed no
369 effects involving DFS (diet) score. We also tested if the degree of fat suppression of the cherry
370 odourant was associated with DFS (diet) score, but no diet-related effects emerged.

371 Hedonics: The hedonic data were analysed using the same ANCOVA design as above.
372 There were main effects of Sugar and Fat level and an interaction between Sugar level and Fat
373 level, and by that between DFS (diet) score, Sugar level, Fat level and Odourant level
374 ($F(4,340) = 3.19, p < 0.02, \text{partial eta-squared} = 0.04$). To examine the source of this four-way
375 effect we conducted eight further ANCOVA's – Fat and Odourant at each Sugar level (3
376 analyses), Fat and Sugar at each Odourant level (3 analyses) and Sugar and Odourant at each
377 Fat level (2 analyses). In each case we examined for the interaction between the covariate and
378 the two within participant variables present in each analysis (respectively; Fat by Odourant, Fat
379 by Sugar, Sugar by Odourant), setting alpha at 0.00625 (Bonferonni correction). One
380 interaction effect was detected in this way between Sugar, Odourant and DFS (diet) score in
381 the low fat skimmed milk samples ($p < 0.003$). We then examined this further by looking at
382 the difference in liking ratings between the unodourised and the highly odourised samples (to
383 maximise any differences), at each level of sweetness. These three mean difference scores
384 were then analysed using a one-way repeated measures ANCOVA, with Sugar level as the
385 within factor and DFS (diet) score as the covariate. There was one effect, a significant

386 interaction between the covariate and Sugar level ($F(2,170) = 5.46, p < 0.005$, partial eta-
387 squared = 0.06). We then examined for the source of this effect by comparing the correlation
388 between the DFS (diet) score and each of these three difference scores using the Williams test,
389 with alpha set at 0.017 (Bonferroni correction). The difference emerged between the medium
390 and high sugar level, and DFS (diet) score, with a resultant correlation of 0.32 ($p < 0.005$). In
391 skimmed milk, the addition of the odourant enhanced pleasantness most in the medium sugar
392 level for participants who consumed a more healthful diet, while the odourant enhanced
393 pleasantness most in the high sugar level for participants who consumed a Western-style diet.
394 This correlation was not attenuated, when partialling out the effects of age, gender, BMI and
395 chemosensory problem score ($r = 0.31, p < 0.005$).

396

397

Discussion

398 The aim of this study was to examine the chemosensory correlates of a Western-style
399 diet. Several findings emerged (see Table 2 for summary). On the olfactory tests of threshold,
400 discrimination and identification, only an association between DFS (diet) score and
401 identification was observed, supporting our previous unpublished findings for identification,
402 but not for discrimination. We also expected to observe differences relating to PROP
403 perception and these too were noted. Participants who consumed a Western-style diet judged
404 bitter tasting PROP to be more intense and liked it less than those who reported consuming a
405 diet lower in saturated fat and added sugar. The study also explored whether certain aspects of
406 flavour perception might be related to diet. Only one effect was observed relating to sweetness
407 enhancement (albeit weakened when the control variables were partialled out), with no effects
408 for odour enhancement or fat induced odour suppression. The Flavour and Taste tests both
409 revealed some additional diet-related effects. Consumers of a Western-style diet were poorer
410 at discriminating the fat levels used in the experiment and their hedonic responses to some of
411 the flavour and taste stimuli also differed. The Flavour and Taste tests yielded no diet-related

412 differences in intensity perception this being consistent with many previous reports (e.g.,
413 Bertino et al., 1986; Mattes, 1985; Moskowitz et al., 1975).

414 Before discussing these findings it is important to consider their limitations. One
415 potential limitation relates to the small to moderate effect sizes observed here. First, these
416 effects could be an artefact of multiple comparisons, but this suggestion would seem less likely
417 in that several findings were anticipated (e.g., PROP, fat discrimination, odour identification).
418 Second, it could be concluded that even if the observed effects are genuine and replicable, their
419 impact on food choice and ultimately human health would be correspondingly small.
420 However, this might not be the case, as effect size does not directly relate to an effect's
421 importance (e.g., McCartney & Rosenthal, 2000). Small effects can exert large impacts
422 especially at the population level, and more so if multiple small effects independently influence
423 behaviour. Third, it has to be born in mind that measurement error is a significant issue in this
424 field. Dietary intake measures are noisy, laboratory based measures of perceptual ability (e.g.,
425 watery taste solutions; non-food odour for threshold) may not fully relate to the way these
426 abilities manifest outside of the laboratory, and there may be disagreements between studies
427 due to differences in the measures used (see Cox et al., 2016; Mattes, 1985). While we note
428 that the techniques used here were not exceptional relative to other studies in the area, we did
429 attempt to use the standardised procedures when available (i.e., Sniffin sticks; PROP protocol).

430 A further consideration is whether some other variable(s) might be mediating the
431 observed relationships between diet and chemosensory performance. In epidemiological
432 studies, dietary associates of psychological variables (notably relating to cognition) are often
433 mediated by socioeconomic status (SES) and especially by education (e.g., Akbaraly et al.,
434 2009). It would seem unlikely that SES would be a major factor in moderating the
435 chemosensory variables tested here, but even if it were, our sample were all receiving a
436 university education in a catchment that draws from a wealthy area of Sydney, making it fairly
437 homogenous with regard to SES. Gender is clearly another variable that may affect diet (e.g.,

438 Rozin, Bauer & Catanese, 2003) and chemosensory ability (e.g., Dempsey & Stevenson,
439 2002). While we found no relationship between DFS (diet) score and gender in the sample
440 used here, this variable was nonetheless included in all of the partial correlations. The same
441 also applies to age and BMI, which are factors also linked to both diet and chemosensory
442 ability. A further control factor used in the partial correlation analyses was the chemosensory
443 problem score, which tended ($p = 0.095$) to be somewhat higher in participants who consumed
444 a Western-style diet. Finally, while we did not explore cultural background as a factor, this
445 could be important in future studies. Early experience with flavours and smells within a
446 particular culture, may affect later processing of these stimuli as an adult (e.g., Poncelet et al.,
447 2010), providing a further factor that might affect diet-perception relationships.

448 In the Introduction we suggested that people with poorer olfactory abilities might
449 gravitate to less healthy food choices. The basic rationale for this assertion is that a major
450 component of flavour perception comes from olfaction (Stevenson, 2014), and if this input is
451 weakened, participants might compensate by choosing diets that offer greater taste and
452 somatosensory impact. Some support for this idea came from the finding in anosmics that
453 dietary shifts are made towards more energy dense foods. However, anosmia appears to have
454 divergent effects on food intake and BMI, with some anosmic participants reporting both BMI
455 and food intake reductions and others the reverse (e.g., Ferris & Duffy, 1989; Merkonidis et al.,
456 2015). The main problem with the idea that dietary choices may be shifted towards less
457 healthy alternatives if olfactory ability is poorer, is that we only found evidence of poorer
458 odour identification ability. Differences in threshold and discrimination would have provided
459 far more robust support to this idea, because they would have suggested that frequent
460 consumers of a Western-style diet could not properly detect and distinguish odours. Perhaps
461 then poorer odour identification ability is just a consequence of reduced exposure to food-
462 based odours – noting their predominance in the Sniffin sticks test battery (e.g., eating fewer
463 fruits and vegetables, less buying, cooking and preparing food, etc).

464 A further perspective is available on the odour identification data. Animal studies
465 indicate that the hippocampus become rapidly and adversely affected by a Western-style diet
466 (e.g., Beilharz, Maniam & Morris, 2014). The hippocampus may be especially sensitive to
467 environmental insults (such as from diet) because it exhibits high synaptic plasticity and
468 neurogenesis (Murray & Holmes, 2011; Walsh & Emerich, 1988). The olfactory system might
469 be similarly vulnerable, as it too demonstrates high synaptic plasticity and neurogenesis
470 (Lledo, Alonso & Grubb, 2006). Indeed, recent animal work has shown olfactory impairments
471 following a high fat diet (Thiebaud et al., 2014). As human olfactory identification is
472 supported at least in part by the hippocampus (e.g., Kjelvik et al., 2012) it is plausible that
473 consuming a Western-style diet could also cause impaired identification.

474 For the PROP-related findings, the direction of the causal arrow would seem far more
475 assured. There are many studies (see Feeney, 2011; Hayes et al., 2013, for reviews) that show
476 a weak to moderate relationship ($r \approx 0.2$) between intake of cruciferous vegetables and PROP
477 sensitivity, as measured in one of several different ways. Presumably this relationship occurs
478 because the bitterness of these vegetables is unpleasantly intense to individuals who have a
479 genetic predisposition to strongly experience PROP bitterness. The effects found here are of
480 broadly similar magnitude to the vegetable-PROP literature, but are interesting for two
481 additional reasons. First, they are novel because the dietary variable here is generic, unlike
482 much of this literature, which has focussed on specific foods likely to be impacted by PROP
483 sensitivity. Second, while greater PROP sensitivity might reduce fruit and vegetable intake, it
484 has also been argued that it increases sensitivity to sweet and fatty tastes due to the greater
485 number of fungiform papillae (e.g., Bartoshuk et al., 2006; Hayes & Duffy, 2007). This may
486 result in a PROP-sensitive person preferring lower concentrations of sweet and fatty tastes,
487 which might moderate some of the effects on fruit and vegetable intake (Bartoshuk et al.,
488 2006). However, few studies have examined actual fat intake and PROP perception, and the
489 only finding to emerge has been of greater fat intake in PROP sensitive people (Yackinous &

490 Guinard, 2002). Nonetheless, other studies suggest that the relationship between PROP
491 sensitivity and sweet and fatty taste perception may be more nuanced, with other factors
492 affecting these relationships including quinine sensitivity (Hayes & Duffy, 2008). In sum, it
493 seems likely that the positive association revealed in this study between PROP sensitivity and
494 DFS (diet) score arises because the main causal relationship is one in which dislike of bitter
495 tastes produces preferences for diets low in plant based foods.

496 It is clear from laboratory studies that there is considerable individual variation in fat
497 perception (e.g., Tucker & Mattes, 2013), but it is far less clear how this impacts on dietary
498 choice and fat intake, as findings here have been mixed (e.g., Cooling & Blundell, 2001).
499 Here, we found that better fat discrimination between the skimmed and full fat milk samples,
500 indicated by a larger difference in fattiness ratings, was associated with DFS (diet) score.
501 Specifically, participants reporting greater intake of saturated fat were the poorest
502 discriminators. Liang et al., (2007) made a similar observation when correlating performance
503 on a laboratory fat discrimination task with self-reported food intake, in a much larger African-
504 American sample. Poorer discriminative performance was associated with greater intake of
505 high fat foods, sources of added fat and sugar, and reduced fat foods. It is currently unclear
506 whether these effects of fat discrimination are a consequence of dietary exposure or result from
507 pre-existing differences in sensory physiology. There is evidence for both, as controlled
508 exposure to low fat foods seems to improve various aspects of fat perception (threshold,
509 ranking of fat content; Stewart & Keast, 2012; and see Newman, Haryono & Keast, 2013),
510 while as noted above, PROP sensitivity, and certain receptor gene variants can also affect fat
511 perception (e.g., CD36 receptor; Keller et al., 2012). Finally, these types of individual
512 differences in fat perception may be practically significant, as a recent laboratory study found
513 that poorer fat perception was associated with greater food intake (Keast et al., 2014).

514 In examining the perceptual correlates of habitual diet there has been very little interest
515 in flavour perception. Here we focussed on two classes of flavour-related interaction effect,

516 both of which are psychologically based (Stevenson, 2009). The first concerned interactions
517 between taste and smell, and the ability of tastes to augment perception of certain odours and
518 vice versa (Frank & Byram, 1988; Von Sydow et al., 1974). The second, relates to the ability
519 of fats to suppress perception of odours in the mouth (Bult, de Wijk & Hummel, 2007). We
520 found no evidence for diet-related differences in odour-fat interactions, but we did find an
521 effect for sweetness taste enhancement (i.e., where, in the mouth, a sweet taste is judged to
522 taste sweeter in the presence of certain odours) - noting that this was only marginally
523 significant after partialling out the control variables. People reporting a Western-style diet
524 tended to demonstrate taste enhancement effects for the weak odour concentration, but not for
525 the strong, while participants with a healthier diet reported a small degree of enhancement for
526 the weak odour concentration and most for the strong concentration. One way of
527 understanding the origin of odour-taste interactions is that they are a product of learning (e.g.,
528 Stevenson, Boakes & Prescott, 1998). On this basis, when tastes are present in the mouth,
529 participants with Western-style diets may tend to experience typically weaker oral odour
530 sensations than participants with a healthier diet. Thus weaker smells may be more likely to
531 enhance tastes than stronger smells in frequent consumers of a Western-style diet. There are
532 two reasons to suspect that exposure to weaker oral olfactory percepts actually occurs in
533 habitual consumers of a Western-style diet. Not only may their greater fat intake suppress oral
534 odour perception they may also consume fewer foods that have high volatile contents (i.e.,
535 fruits and vegetables). Needless to say, we only tested one oral odourant, one tastant and one
536 fatty vehicle, but the observation of an effect here suggests that dietary associates of flavour
537 perception may not be hard to find.

538 Two observed effects related to taste and flavour hedonics. First, the salt solutions
539 were judged as less pleasant by more frequent consumers of a Western-style diet. This finding
540 is surprising as higher salt concentrations are a feature of this type of diet. However, saline
541 solutions are infrequently experienced outside of the laboratory and so there dislike could

542 reflect greater neophobia in more frequent consumers of a Western-style diet – something that
543 has been observed before (Siegrist et al., 2013). In addition, greater PROP sensitivity is also
544 associated with reduced liking for aqueous salt solutions (Hayes, Sullivan & Duffy, 2010).
545 Second, hedonic differences also emerged on the flavour analysis. Cherry odourant enhanced
546 the pleasantness of skimmed milk most successfully in the medium sugar level for those who
547 ate a healthier diet, and at the higher sugar level for those who consumed a Western-style diet.
548 As the higher sugar level generates both a sweeter taste and a fattier mouthfeel, this may be
549 more appealing to participants who consume fattier and sweeter foods.

550 Finally, we note that many elderly people have impaired chemosensory perception
551 (e.g., Doets & Kremer, 2016), yet this appears to have relatively little impact upon their
552 enjoyment of food (e.g., Arganini & Sinesio, 2015; Kremer et al., 2007). This might suggest
553 that sensory differences have little impact on dietary choice. We suspect this conclusion may
554 be less likely to apply to younger people. First, hedonic reactions to food have a learned
555 component. Elderly consumers - in contrast to younger ones - may be more reliant on such
556 learned reactions and thus less susceptible to the effects of sensory loss. Second, sensory
557 factors that favor unhealthy food choices may aid establishing a dietary pattern that then
558 becomes habitual, making it more resistant to change during ageing. Third, to the extent that
559 certain sensory differences are innate, these may affect parental dietary choice, which in turn
560 will shape the foods the child is exposed to. Importantly, we note that as yet there has been no
561 systematic examination of how broad sensory differences in young adults may affect food
562 choice – in contrast to the studies completed for the elderly (e.g., the European Union
563 Healthsense project).

564 In conclusion, we set out to examine chemosensory correlates of a Western-style diet.
565 We observed differences in odour identification ability, PROP sensitivity, fat discrimination,
566 sweetness taste enhancement, and taste and flavour hedonics, but no differences in odour
567 discrimination or threshold, in perception of taste intensity/quality (excluding PROP) or the

568 ability of fats to affect flavour perception. Most of the observed relationships were of small to
569 moderate effect size. While their manifestation in habitual consumers of a Western-style diet
570 would generally seem to work against eating a more healthful diet, whether they are a cause or
571 a consequence of dietary choices remains to be established for most of the effects reported
572 here.

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Acknowledgments

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577 The authors thank Lina Teichmann, Kristen Manchester, Kimberley Weldon, Andrea
578 Zuniga, Jade Jackson and Alessandra Teunisse for their assistance with this study. This work
579 was supported by a grant from the Australian Research Council [grant number DP150100105].

References

- 580
581 Akbaraly TN, Singh-Manoux A, Marmot MG, Brunner EJ, (2009). Education attenuates the
582 association between dietary patterns and cognition. *Dementia Geriatric Cog Disorders.*
583 27:147-154.
- 584 Appel L, Moore T, Obarzanek E, Vollmer W, Svetkey L, Sacks F, Bray G, Vogt T, Cutler J,
585 Windhauser M, Lin P, Karanja N. (1996). A clinical trial of the effects of dietary
586 patterns on blood pressure. *New Eng J Med.* 336:1117-1124.
- 587 Arganini C, Sinesio F. (2015). Chemosensory impairment does not diminish eating pleasure
588 and appetite in independently living older adults. *Maturitas* 82:221-244.
- 589 Attuquayefio T, Stevenson RJ, Boakes RA, Oaten MJ, Yeomans MR, Mahmut MK, Francis H.
590 (Submitted). A Western-style diet is associated with poorer state-dependent inhibition of
591 wanting for palatable snack foods: The role of hippocampal related processes.
- 592 Bachmanov A, Kiefer S, Molina J, Tordoff M, Duffy V, Bartoshuk L, Mennella J. (2003).
593 Chemosensory factors influencing alcohol perception, preferences and consumption.
594 *Alcoholism-Clin Exp Res.* 27:220-231.
- 595 Bartoshuk L, Duffy V, Hayes J, Moskowitz H, Snyder D. (2006). Psychophysics of sweet and
596 fat perception in obesity: Problems, solutions and new perspectives. *Phil Trans Royal*
597 *Soc B.* 361:1137-1148.
- 598 Bertino M, Beauchamp GK, Engelman K. (1982). Long-term reduction in dietary sodium
599 intake alters the taste of salt. *Am J Clin Nutr.* 36:1134-1144.
- 600 Bertino M, Beauchamp GK, Engelman K. (1986). Increasing dietary salt alters salt taste
601 preference. *Physiol Behav.* 38:203-213.
- 602 Beilharz J, Maniam J, Morris M. (2014). Short exposure to a diet rich in both fat and sugar or
603 sugar alone impairs place, but not object recognition memory in rats. *Brain Behav*
604 *Immun.* 37:134-141.

- 605 Bult JHF, de Wijk RA, Hummel T. (2007). Investigations on multimodal sensory integration:
606 texture, taste, and ortho- and retronasal olfactory stimuli in concert. *Neurosci Lett.* 411:6-
607 10.
- 608 Cicerale S, Riddell LJ, Keast RSJ. (2012). The association between perceived sweetness
609 intensity and dietary intake in young adults. *J Food Sci.* 71:31-35.
- 610 Cooling J, Blundell JE. (2001). High-fat and low-fat phenotypes: Habitual eating of high- and
611 low-fat foods not related to taste preference for fat. *E J Clin Nutr.* 55:1016-1021.
- 612 Cox D, Hendrie G, Carty D. (2016). Sensitivity, hedonics and preferences for basic tastes and
613 fat amongst adults and children of differing weight status. *Food Qual Pref.* 48:359-367.
- 614 Doets E, Kremer S (2016). The silver sensory experience – A review of senior consumers food
615 perceptions, liking and intake. *Food Qual Pref.* 48:316-332.
- 616 Dempsey RA, Stevenson RJ (2002). Gender differences in the retention of Swahili names for
617 unfamiliar odors. *Chem Sens.* 27:681-689.
- 618 Drewnowski A, Brunzell JD, Sande K, Iverius PH, Greenwood MRC. (1985). Sweet tooth
619 reconsidered: Taste responsiveness in human obesity. *Physiol Behav.* 35:617-622.
- 620 Duffy V, Hayes J. (2014). Smell, taste and oral somatosensation: Age-related changes and
621 nutritional implications. In (Ed) R. Chernoff, *Geriatric Nutrition* (pp.115-164). Jones and
622 Bartlett Learning, Burlington MA.
- 623 Duffy V, Hayes J, Sullivan B, Faghri P. (2009). Surveying food and beverage liking: A tool for
624 epidemiological studies to connect chemosensation with health outcomes. *Ann NY Acad*
625 *Sci.* 1170:558-568.
- 626 Feeney E. (2011). The impact of bitter perception and genotypic of TAS2R38 on food choice.
627 *Nutr Bull.* 36:20-33.
- 628 Ferris A, Duffy V. (1989). Effects of olfactory deficits on nutritional status: Does age predict
629 persons at risk? *Ann NY Acad Sci.* 561:113-123.

- 630 Francis H, Stevenson RJ. (2013). Validity and test-retest reliability of a short dietary
631 questionnaire to assess intake of saturated fat and free sugars: a preliminary study. *J Hum*
632 *Nutr Diet.* 26:234-242.
- 633 Frank RA, Byram J. (1988). Taste-smell interactions are tastant and odorant dependent. *Chem*
634 *Sens.* 13:445-455.
- 635 Frost M, Janhøj J. (2007). Understanding creaminess. *Int Dairy J.* 17:1298-1311.
- 636 Green B, Dalton P, Cowart B, Shaffer G, Rankin K, Higgins J. (1996). Evaluating the labelled
637 magnitude scale for measuring sensations of taste and smell. *Chem Sens.* 21:323-334.
- 638 Hayes J, Duffy V. (2007). Revisiting sugar-fat mixtures: Sweetness and creaminess vary with
639 phenotypic markers of oral sensation. *Chem Sens.* 32:225-236.
- 640 Hayes J, Duffy V. (2008). Oral sensory phenotype identifies level of sugar and fat required for
641 maximal liking. *Physiol Behav.* 95:77-87.
- 642 Hayes J, Feeney E, Allen A. (2013). Do polymorphisms in chemosensory genes matter for
643 human ingestive behaviour. *Food Qual Pref.* 30:202-216.
- 644 Hayes J, Sullivan B, Duffy V. (2010). Explaining variability in sodium intake through oral
645 sensory phenotype, salt sensation and liking. *Physiol Behav.* 100:369-380.
- 646 Hummel T, Sekinger B, Wolf S, Pauli E, Kobal G. (1997). 'Sniffin' sticks: Olfactory
647 performance assessed by the combined testing of odor identification, odor discrimination
648 and olfactory threshold. *Chem Sens.* 22:39-52.
- 649 Keast R, Azzopardi K, Newman L, Haryono R. (2014). Impaired oral fatty acid
650 chemoreception is associated with acute excess energy consumption. *Appetite* 80:1-6.
- 651 Keller KL, Liang LCH, Sakimura J, May D, van Belle C, Breen C, Chung WK. (2012).
652 Common variants in the CD36 gene are associated with oral fat perception, fat
653 preferences, and obesity in African Americans. *Obesity.* 20:1066-1073.
- 654 Kirkmeyer S, Tepper B. (2005). Consumer reactions to creaminess and genetic sensitivity to
655 PROP: A multidimensional study. *Food Qual Pref.* 16:545-556.

- 656 Kjolvik G, Evensmoen H, Brezova V, Haberg A. (2012). The human brain representation of
657 odor identification. *J Neurophysiol.* 108:645-657.
- 658 Kremer S, Bult J, Mojet J, Kroeze J. (2007). Food perception with age and its relationship to
659 pleasantness. *Chem Sens.* 32:591-602.
- 660 Liang LCH, Sakimura J, May D, Breen C, Driggin E, Tepper BJ, Keller KL. (2012). Fat
661 discrimination: A phenotype with potential implications for studying fat intake behaviors
662 and obesity. *Physiol Behav.* 105:470-475.
- 663 Lim K, Wood A, Green BG. (2009). Derivation and evaluation of a labelled hedonic scale.
664 *Chem Sens.* 34:739-751.
- 665 Lledo P, Alonso M, Grubb M. (2006). Adult neurogenesis and functional plasticity in neuronal
666 circuits. *Nat Rev Neurosci.* 7:179-193.
- 667 Mattes, RD. (1985). Gustation as a determinant of ingestion: methodological issues. *Am J Clin*
668 *Nutr.* 41:672-683.
- 669 McCartney K, Rosenthal R. (2000). Effect size, practical importance and social policy for
670 children. *Child Dev.* 71:173-180.
- 671 Mente A, de Koning L, Shannon H, Anand S. (2009). A systematic review of the evidence
672 supporting a causal link between dietary factors and coronary heart disease. *Arch Intern*
673 *Med.* 169:659-669.
- 674 Merkonidis C, Grosse F, Ninh T, Hummel C, Haehner A, Hummel T. (2015). Characteristics
675 of chemosensory disorders – results from a survey. *Eur Arch Otorhinolaryngol.*
676 272:1403-1416.
- 677 Miwa T, Furukaw M, Tsukatani T, Costanzo R, DiNardo L, Reiter E. (2001). Impact of
678 olfactory impairment on quality of life and disability. *Arch Otolaryngol Head Neck Surg.*
679 127:497-503.
- 680 Moskowitz H, Kumariah V, Sharma K, Jacobs H, Sharma S. (1975). Cross-cultural differences
681 in simple taste preferences. *Sci.* 190:1217-1218.

- 682 Murray P, Holmes P. (2011). An overview of BDNF and implications for excitotoxic
683 vulnerability in the hippocampus. *Int J Peptides*: e654085.
- 684 Newby P, Weismayer C, Akesson A, Tucker K, Wolk A. (2006). Long-term stability of food
685 patterns identified by use of factor analysis among Swedish women. *J Nutr*. 136:626-633.
- 686 Newman L, Haryono R, Keast R. (2013). Functionality of fatty acid chemoreception: A
687 potential factor in the development of obesity? *Nutrients* 5:1287-1300.
- 688 Pachucki M. (2012). Food pattern analysis over time: unhealthful eating trajectories predict
689 obesity. *Int J Obesity*. 36:686-694.
- 690 Poncelet J, Rinck F, Bourgeat F, Schaal B, Rouby C, Bensafi M, Hummel T. (2010). The effect
691 of early experience on odor perception in humans: Psychological and physiological
692 correlates. *Behav Brain Res*. 208:458-465.
- 693 Preacher K, Rucker D, MacCallum R, Nicewander W. (2005). Use of the extreme groups
694 approach: A critical re-examination and new recommendations. *Psychol Methods*.
695 10:178-192.
- 696 Prescott J, Ripandelli N, Wakeling I. (2001). Binary taste mixture interactions in PROP non-
697 tasters, medium-tasters and super-tasters. *Chem Sens*. 26:993-1003.
- 698 Richardson B, Vander Woude E, Sudan R, Thompson J, Leopold D. (2004). Altered olfactory
699 acuity in the morbidly obese. *Obesity Surg*. 14:967-969.
- 700 Rozin P, Bauer R, Catanese D. (2003). Food and life, pleasure and worry, among American
701 college students: Gender differences and regional similarities. *J Pers Soc Psychol*. 85,
702 132-141.
- 703 Sakai N, Kobayakawa T, Gotow N, Saito S, Imada S. (2001). Enhancement of sweetness
704 ratings of aspartame by a vanilla odor presented either by orthonasal or retronasal routes.
705 *Percept Mot Skills*. 92:1002-1008.
- 706 Siegrist M, Hartmann C, Keller C. (2013). Antecedents of food neophobia and its association
707 with eating behaviour and food choices. *Food Qual Pref*. 30:293-298.

- 708 Stafford L, Whittle A. (2015). Obese individuals have higher preference and sensitivity to odor
709 of chocolate. *Chem Sens.* 40:279-284
- 710 Stevenson RJ. (2009). *The Psychology of Flavour*. OUP, Oxford.
- 711 Stevenson RJ. (2014). Flavor binding: Its nature and cause. *Psychol Bull.* 140:487-510.
- 712 Stevenson RJ, Boakes RA, Prescott J. (1998). Changes in odor sweetness resulting from
713 implicit learning of a simultaneous odor-sweetness association: an example of learned
714 synesthesia. *Learn Motiv.* 29:113-132.
- 715 Stevenson RJ, Miller L, McGrillen K. (2013). The lateralization of gustatory function and the
716 flow of information from tongue to cortex. *Neuropsychologia* 51:1408-1416.
- 717 Stewart JE, Keast RSJ. (2012). Recent fat intake modulates fat taste sensitivity in lean and
718 overweight subjects. *Int J Obesity.* 36:834-842.
- 719 Thiebaud N, Johnson MC, Butler JL, Bell GA, Ferguson KL, Fadool AR., Fadool DA. (2014).
720 Hyperlipidemic diet causes loss of olfactory sensory neurons, reduces olfactory
721 discrimination, and disrupts odor-reversal learning. *J Neurosci.* 34:6970-6984.
- 722 Tucker RM, Mattes RD. (2013). Influences of repeated testing on nonesterified fatty acid taste.
723 *Chem Sens.* 38:325-332.
- 724 Von Sydow E, Moskowitz H, Jacobs H, Meiselman H. (1974). Odor-taste interaction in fruit
725 juices. *Lebensmittel-Wissenschaft und Technologie.* 7:18-24.
- 726 Van Toller S. (1999). Assessing the impact of anosmia: Review of questionnaire findings.
727 *Chem Sens.* 24:705-712.
- 728 Walsh T, Emerich D. (1988). The hippocampus as a common target of neurotoxic agents.
729 *Toxicol.* 49:137-140.
- 730 Yackinous CA, Guinard J-X. (2002). Relation between PROP (6-n-propylthiouracil) taster
731 status, taste anatomy and dietary intake measures for young men and women. *App.*
732 38:201-209.
- 733

734 Table 1: Descriptive statistics and Pearson correlations between participant characteristics,
 735 DFS (diet) score

737 Variable	Descriptive statistics	Correlation with DFS (Diet) score
740	<hr/>	
741	DFS (diet) score	M = 59.9, SD = 13.7, range 37-104
742	743 Gender	38 men/49 women
744	Age	M = 20.8, SD = 3.3, range 18-31
745	BMI	M = 22.3, SD = 3.1, range 16.0-32.7
746	Chemosensory problem score	
747	M = 1.0, SD = 1.0, range 0-4	0.18
748	<hr/>	
749		
750		

751 Table 2: Chemosensory correlates of a Western-style diet (WS-D) alongside the variance
 752 accounted for by each effect

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754 Test	755 Correlate	756 Variance accounted for	757 Finding
758 Sniffin Sticks	759 Odour identification	4.0%	Poorer in consumers of a WS-D
760 Gustatory tests	761 PROP sensitivity	4.0%	Greater in consumers of a WS-D
	762 PROP disliking	4.4%	Greater in consumers of a WS-D
	763 Salty taste disliking	6.8%	Greater in consumers of a WS-D
764 Flavour test	765 Sweetness enhancement	766 4.4%	767 Maximal enhancement at lower odourant level in consumers of a WS-D
	768 Fat discrimination	8.4%	Poorer in consumers of a WS-D
	769 Flavour hedonics	10.2%	770 Odour increased pleasantness of skimmed milk most at high sugar-levels in consumers of a WS-D

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