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1 **Identification of Psychological Correlates of Dietary Mis-Reporting under Laboratory**
2 **and Free-Living Environments**

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31 **ABSTRACT**

32 Errors inherent in self-reported measures of energy intake (EI) are substantial and well-
33 documented, but correlates of mis-reporting remain unclear. Therefore, potential predictors of
34 mis-reporting were examined. In Study One, 59 individuals (BMI=26.1±3.8kg/m²,
35 age=42.7±13.6yrs, females=29) completed a 14d stay in a residential feeding behaviour suite
36 where eating behaviour was continuously monitored. In Study Two, 182 individuals
37 (BMI=25.7±3.9kg/m², age=42.4±12.2yrs, females=96) completed two consecutive days in a
38 residential feeding suite and five consecutive days at home. Mis-reporting was directly
39 quantified by comparing covertly measured laboratory weighed intakes (LWI) to self-reported
40 EI (weighed dietary record; WDR, 24-hr recall, 7-day diet history, food frequency
41 questionnaire; FFQ). Personal (age, sex, %body fat) and psychological traits (personality,
42 social desirability, body image, IQ, eating behaviour) were used as predictors of mis-reporting.
43 In Study One, those with lower psychoticism (p=0.009), openness to experience (p=0.006) and
44 higher agreeableness (p=0.038) reduced EI on days participants knew EI was being measured
45 to a greater extent than on covert days. Isolated associations existed between personality traits
46 (psychoticism, openness to experience), eating behaviour (emotional eating) and differences
47 between the LWI and self-reported EIs, but these were inconsistent between dietary assessment
48 techniques and typically became non-significant after accounting for multiplicity of
49 comparisons. In Study Two, sex was associated with differences between LWI and the WDR
50 (p=0.009), 24-hr recall (p=0.002) and diet history (p=0.050) in the laboratory, but not home
51 environment. Personal and psychological correlates of mis-reporting identified displayed no
52 clear pattern across studies or dietary assessment techniques, and had little utility in predicting
53 mis-reporting.

54 1.0 INTRODUCTION

55 The relationship between energy and nutrient intake and disease prevalence is crucial in
56 understanding disease aetiology at the individual and population level. However, quantifying
57 true patterns of food intake in the free-living environment is severely limited by the under or
58 over-reporting of energy and nutrient intakes using self-report techniques. This has led to
59 suggestions that self-report dietary techniques are not only “useless” in elucidating diet-health
60 relationships, but may actually distort the true nature of the relationships upon which nutritional
61 health policies are based^(1; 2). Although this view has been refuted⁽³⁾, errors inherent in self-
62 reported intakes appear substantial⁽⁴⁾. Dietary mis-reporting with self-report techniques has
63 long been recognised⁽⁵⁾, but this has yet to lead to the development of techniques that i) detect
64 the extent of mis-reporting in self-reported dietary data, ii) identify or predict those likely to
65 mis-report using self-report techniques, and iii) correct for erroneous values in self-reported
66 data.

67
68 Previous studies suggest that under-reporting is more prevalent in women^(6; 7), older rather than
69 younger adults⁽⁷⁾, and those with higher BMIs^(4; 8). However, identification of consistent
70 correlates of mis-reporting across different self-reported dietary measurement techniques (e.g.
71 food frequency questionnaires, 24-hr dietary recalls, dietary records/diaries), study populations
72 (e.g. sex, age, ethnicity, social class and educational level) or environments (e.g. laboratory vs
73 free-living) has proved remarkably difficult. An array of psychological, personality and social
74 characteristics have been suggested as potential correlates, including dietary restraint^(9; 10),
75 social desirability and approval^(10; 11), social economic class and educational level^(12; 13; 14).
76 However, purported correlates are often not consistent between studies and typically only
77 explain a small proportion of the variance in under or over reporting^(7; 15). This failure to
78 identify robust correlates of mis-reporting may reflect the fact that previous studies have not
79 directly quantified mis-reporting (i.e. the discrepancy between what people actually eat and
80 report eating), but rather, use indirect estimates of low or high energy reporting based on
81 indices of energy balance (e.g. doubly labelled water^(6; 10) or the Goldberg cut-offs⁽¹⁶⁾) or
82 nitrogen balance (e.g. dietary to urinary nitrogen ratios⁽¹⁷⁾). Given the limitations associated
83 with these approaches in identifying mis-reporting at the individual level^(18; 19; 20), these indirect
84 estimates may lack sufficient sensitivity to detect correlates of under or over reporting.

85

86 Identification and prediction of dietary mis-reporting is further complicated by the fact that
87 mis-reporting is not a unitary phenomenon. Rather, it comprises of two separate but
88 synchronous processes, termed the observation effect and the reporting effect⁽²¹⁾, that summate
89 to determine overall mis-reporting. Based on covert measures of food intake during a 14 day
90 stay in residential metabolic facility, Stubbs et al.⁽²¹⁾ were able to directly compare actual food
91 intake to that self-reported by participants during their stay. Participants were shown to
92 decrease their energy intake (EI) by 5% when asked to record their food intake, which was
93 termed the observation effect. Self-reported EI was 5 to 21% lower than the actual intake,
94 depending on the reporting method used (termed the reporting effect). However, potential
95 correlates of the observation and reporting effects have yet to be examined in these data.

96

97 Therefore, the present paper examined the psychological correlates of mis-reporting in two
98 separate studies in which objective and self-reported food intake was measured to directly
99 quantify mis-reporting of EI under i) residential laboratory conditions in which energy balance
100 and feeding behaviour were measured continuously for 14 days (Study One; $n = 59$)⁽²¹⁾, and ii)
101 combined residential (two days) and free-living (four days) conditions in which laboratory
102 dietary intakes were compared to self-reported assessments made in the laboratory and home
103 environments (Study Two; $n = 182$)⁽¹⁹⁾. This approach allowed mis-reporting to be directly
104 quantified in a metabolic facility and under simulated conditions representative of the
105 environments in which EI is often estimated in dietary survey studies using self-report
106 techniques. These studies included commonly used self-report techniques (weighed dietary
107 records, 24-hr recall, food frequency questionnaire and diet history), and the validity of these
108 approaches has been discussed elsewhere⁽⁷⁾.

109

110 **2.0 METHODS**

111 Data from two separate studies are reported in which dietary mis-reporting was directly
112 quantified by comparing covertly measured food intake to self-reported intakes using four
113 commonly used methods (weighed dietary records, 24-hr recalls, 7-day diet history, food
114 frequency questionnaire). In Study One, 59 participants (age = 42.7 ± 13.6 years; BMI = 26.1
115 ± 3.8 kg/m²) completed a 14 day stay in a residential feeding behaviour suite during which
116 food intake was recorded for 12 consecutive days following a two day maintenance period. In
117 Study Two, 182 participants (age = 42.4 ± 12.2 years; BMI = 25.7 ± 3.9 kg/m²) completed
118 three consecutive days (one day maintenance and two days recording) in a residential feeding
119 behaviour suite and five consecutive days (one day maintenance and four days recording) in

120 their home environment in a randomised, and counter-balanced order. All data were collected
121 at the Rowett Institute, University of Aberdeen, United Kingdom, and participants were weight
122 stable (weight change of <2 kg in the previous three months), healthy, non-smokers, and not
123 taking medication known to influence metabolism or appetite. The true purpose of each study
124 was not explained to participants, who were informed that the studies examined the
125 relationships between diet and lifestyle. Written informed consent was obtained prior to the
126 start of each study. The studies were conducted according to the guidelines laid down in the
127 Declaration of Helsinki, and all procedures involving human subjects/patients were approved
128 by the Joint Ethical Committee of the Grampian Health Board and the University of Aberdeen.
129

130 **2.1 STUDY ONE- Participants and Design**

131 Fifty-nine participants (30 men and 29 women) were recruited, with participants stratified into
132 three age categories (20-35 years, 36-50 years and 51-65 years) and two BMI categories (BMI
133 20-25 kg/m² and BMI >25 kg/m²). Participant characteristics can be seen in Table 2. The
134 overall aim of this study was to develop a gold standard protocol for the measurement of food
135 intake against which common self-reported dietary intake methods could be evaluated. Primary
136 outcomes from this study relating to the nature and extent of dietary mis-reporting have
137 previously been reported⁽²¹⁾. The current novel analyses examined the personal and
138 psychological correlates of this mis-reporting.
139

140 **Figure 1 here**

141
142 Figure 1 describes the experimental protocol, and a detailed description of the procedures used
143 can be found elsewhere⁽²¹⁾. Participants completed a 14 day stay in a residential feeding
144 behaviour suite (Human Nutrition Unit at the Rowett Institute of Nutrition and Health) during
145 which energy balance and feeding behaviours were measured continuously. Resting metabolic
146 rate (indirect calorimetry) was measured on a screening visit prior to the start of the study. On
147 days 1-2, participants consumed a fixed diet designed to maintain energy balance, with EI
148 estimated at 1.5 and 1.6 times resting metabolic rate for women and men, respectively. The
149 proportion of energy contributed by fat, protein and carbohydrate to daily energy intake was
150 35%, 15% and 55%, respectively. Percentage body fat (skinfold thickness) was measured on
151 day 3. On days 3-14, food intake was covertly measured by trained research staff using a
152 laboratory weighed intake method (LWI) to establish actual energy and nutrient intake.
153 Participants were unaware that their food intake was being measured in this fashion using

154 covert LWI measures. Participants also self-reported their food intake using a weighed dietary
155 record method (WDR) and 24-hr recall during two, 3-day overt feeding periods during days 3-
156 14. On these overt feeding days, participants were aware that their food intake was being
157 measured using these self-report techniques, but they remained blinded to the fact that their
158 food intake was also being covertly measured using the LWI. As such, we refer to the days in
159 which self-reported measures of intake were conducted as overt days to reflect the participants
160 awareness that their food intake was being monitored. The order of these overt feeding periods
161 was randomized using a cross-over design. In total, six 24-hr recalls and six weighed dietary
162 records were completed by participants over the 14-day period, while food intake was covertly
163 measured for 12 days. A 7-day diet history was also conducted, between two-days and two-
164 weeks, before the start of the study, and two food frequency questionnaires (FFQ) were
165 completed. The first FFQ was completed on day 1 and related to the frequency of consumption
166 of specific foods over the preceding 2 to 3 months. On day 15, the same FFQ was completed
167 for a second time but pertained to their intake over the preceding 14 days in the residential
168 feeding suite (this is referred to as FFQ²).

169

170 Participants were able to move freely around the unit and associated grounds (under
171 supervision of a member of staff) and were free to leave the unit during the study (but were
172 accompanied and observed by a member of staff at all times). During the 14-day periods,
173 participants also completed a range of psychological questionnaires, and the specific timing of
174 their completion can be found in Table 1.

175

176 **2.2 STUDY TWO- Participants and Design**

177 Participants (n = 182; 86 men and 96 women) were recruited to cover a range of age (25-60
178 years) and BMIs (19-30 kg/m²) in a balanced design. Participant characteristics can be seen in
179 Table 3. This study was designed in parallel with Study One, and aimed to extend this study
180 by identifying the nature and extent of under-reporting in a larger sample of individuals under
181 laboratory and home environments. The plausibility of the self-reported EI relative to the LWI
182 in these data have previously been reported⁽¹⁹⁾. The current analyses are novel. The protocol
183 for Study Two can be seen in Figure 1, and a detailed description of the procedures used can
184 be found elsewhere⁽¹⁹⁾. In a randomised order, participants completed three consecutive days
185 (one day maintenance and two days recording) in the Human Nutrition Unit, Rowett Institute
186 of Nutrition and Health, five consecutive days (one day maintenance and four days recording)

187 in their home environment. Percentage body fat (skinfold thickness) was measured on day 1 of
188 the laboratory phase.

189

190 **2.2.1 Laboratory phase**

191 The laboratory phase consisted of two consecutive days (Friday and Saturday, or Sunday and
192 Monday), in consecutive order, with one day's maintenance diet beforehand. On each day, EI
193 was covertly measured by research staff using the LWI method. Participants also completed a
194 WDR on each day, and a 24-hr dietary recall was performed on the morning of the subsequent
195 day. Prior to the start of the study, participants also completed a FFQ and a 7-day diet history,
196 as in Study One.

197

198 **2.2.2 Home phase**

199 The home study consisted of a one-day maintenance followed by four consecutive days
200 consisting of two weekdays and two weekend days (days 1-4, Thursday to Sunday, or Saturday
201 to Tuesday). During this time participants conducted daily WDR in their home environment
202 (referred to as WDR-H), using the same method as the laboratory phase. No other measures of
203 food intake were taken during this home phase. During the laboratory and home phases of
204 Study Two participants also completed a range of psychological questionnaires, and the
205 specific timing of their completion can be found in Table 1.

206

207 **2.3 COMMON METHODOLOGICAL PROCEDURES**

208 **2.3.1 Resting Metabolic Rate**

209 Resting metabolic rate was measured following an overnight fast (12-hr) using an indirect
210 calorimetry device fitted with a ventilated hood (Deltatrac II, MBM-200, Datex
211 Instrumentarium Corporation, Finland). Resting metabolic rate was calculated from minute-
212 by-minute data using the mean of 15 minutes of stable measurements, with the first and last
213 five minutes excluded. The equations of Elia and Livesey⁽²²⁾ were used to calculate resting
214 metabolic rate. Details of calibration burns and repeatability testing have been described
215 previously⁽²³⁾.

216

217 **2.3.2 Anthropometry and Skinfold Thickness**

218 Height was measured to the nearest 0.5 cm using a portable stadiometer (Holtain Ltd.,
219 Crymych, Dyfed, Wales), while body weight was measured to the nearest 0.01 kg after voiding
220 (DIGI DS-410 CMS Weighing Equipment, London, UK). Skinfold thickness was also

221 measured at standardized anatomic locations (biceps, triceps, subscapular and supra-iliac)
222 using calibrated skinfold callipers (Holtain Ltd., Dyfed, Wales, UK), and the equations of
223 Durnin & Womersley⁽²⁴⁾ were used to estimate percentage body fat from skinfold thickness.

224

225 **2.3.3 MEASURES OF FOOD INTAKE**

226 **Laboratory Weighed Intake Method**

227 During the laboratory phases of Study One and Study Two, each participant had access to their
228 own individual kitchen, which consisted of a fridge, freezer and a cupboard containing pre-
229 selected foods and beverages. Between two-days and two-weeks prior to the start of each study,
230 a 7-day diet history was completed, and shopping receipts were collected. An inventory of
231 foods and beverages they typically consumed was purchased. Participants then had *ad libitum*
232 access to these foods and beverages during the laboratory phases of each study. If a participant
233 reported that a food or beverage usually consumed in their habitual diet had been omitted, this
234 item was subsequently purchased and made available. Participants were able to freely select
235 what and when they wanted to eat (based on their own foods and beverage items), and meals
236 were cooked by participants in their own kitchens. Access to these was restricted, with
237 participants only having key access to his/her own kitchen. Participants were instructed to leave
238 all food waste, peelings and packaging in special bins. Furthermore, any dishes/cooking
239 utensils used were placed in a specific section of their kitchen following meal/snack
240 consumption, and subjects were instructed not to wash any dishes/utensils.

241

242 On days in which the participants stayed in the residential feeding suite in Study One and Two,
243 measures of daily food intake were made using the LWI method. Participants were unaware
244 that their food intake was being measured in this fashion, and therefore we refer to these
245 measures of food intake as covert. Each morning, a researcher entered the kitchen before the
246 participants woke and re-weighed all the food items to the nearest 0.1 g (Soehnle model 820;
247 Soehnle-Waagen GmbH or Ravencourt model 333; Ravencourt), and the weights of any left-
248 overs, peelings and packaging found in their bins were also recorded. The laboratory-weighed
249 intakes were then used to calculate 24-hr food intakes, with EI calculated using dietary analysis
250 software (Diet 5, Robert Gordon University, Aberdeen). Nutritional information from
251 manufacturers was added to the Diet 5 database for processed foods. Each individual kitchen
252 contained a discrete unobtrusive video camera, while all parts of the unit were monitored via
253 video cameras (aside from the bathroom facilities and private rooms; participants were not
254 allowed to take food into these areas). Participants were informed that cameras were present

255 for security purposes, although they were not made overtly aware of the camera in their larders,
256 which resembled an infrared motion detector commonly used in burglar alarm systems. Video
257 data were used to ensure participants were adhering to the study procedures.

258

259 **Weighed Dietary Records**

260 Participants were instructed to carry out weighed dietary records⁽²⁵⁾ on the overt phases of
261 Study One and the laboratory phase of Study Two. Participants were asked to weigh and record
262 all food and drinks consumed and any leftovers, in a food diary. Participants used digital
263 portable weighing scales (Soehnle model 820), which were calibrated prior to use. Full written
264 and verbal information on how to conduct a WDR was given at the beginning of the study and
265 participants were trained in the use of the equipment.

266

267 **Twenty-Four Hour Recalls**

268 24-hr recalls were performed by trained member of staff based on the multiple pass method.
269 Each recall was conducted on the day after participants completed a WDR during the overt
270 phases of each study.

271

272 **7-Day Diet History**

273 Prior to taking part in each study participants completed a 7-day diet history with a trained
274 member of staff. The diet history was based on the multiple pass method. Participants were
275 asked to describe their usual food intake at different meal/snack occasions during the previous
276 week, and were asked to use household measures when recalling food items. This information
277 was also used to formulate a list of foods and beverages usually consumed by each participant,
278 which were made available to them during the laboratory phases of each study. Each diet
279 history was entered into a spreadsheet, and suitable portion sizes were used to convert the
280 household food portion sizes into grams using the UK Food Standard Agency book on average
281 portion sizes⁽²⁶⁾.

282

283 **Food Frequency Questionnaire**

284 The Aberdeen Food Frequency Questionnaire^(27; 28), which is a 150-item semi-quantitative
285 questionnaire, was used to assess the frequency of consumption of foods in the habitual diet of
286 participants in both studies and mean daily energy and nutrient intakes calculated. Full written
287 and verbal information on how to complete this questionnaire was provided.

288 2.3.4 Psychological Predictors

289 A range of common questionnaires to measure aspects of personality and eating behaviours
290 hypothesised to be of potential relevance to biased responding of food intake were completed
291 by participants in both studies to examine potential predictors of dietary mis-reporting^(7; 15). IQ
292 was measured using the National Adult Reading Test (NART)⁽²⁹⁾, the Alice Heim 4 (AH4)⁽³⁰⁾
293 and the Raven Standard Progressive Matrices⁽³¹⁾. The NART is a single word, oral reading test
294 in which participants read out 50 written words with irregular spellings graded in difficulty.
295 The AH4 is a two-part test with multi-choice answers. Part 1 is a 65-item test with verbal or
296 numerical bias that assesses mental arithmetic, vocabulary and reasoning by analogy, while
297 Part 2 is a 65-item test with a diagrammatic bias. The Raven Standard Progressive Matrices
298 tests problem solving ability using shapes and diagrams, and contains 60 problems requiring
299 participants to determine the relationships between abstract shapes. To measure mood, the
300 UWIST Mood Adjective Checklist⁽³²⁾ was used. This measures the average state of mood
301 experienced by the participants during the present day, with 24 separate feelings rated on a
302 scale of definitely to definitely not. Perceptions of body image were measured using the Body
303 Image Questionnaire⁽³³⁾, with participants presented with a series of schematic silhouettes of
304 different body sizes from which they selected the one most representing their own body shape.
305 Personality was measured using two questionnaires; the Eysenck-100 (EPQR)⁽³⁴⁾ and the
306 Neuroticism, Extraversion, Openness Personality Inventory-Revised (NEOPIR)⁽³⁵⁾. The EPQR
307 measures four personality traits (sociability, psychoticism, neuroticism and lie scale), with
308 participants responding true/false to 100 statements. The NEOPIR consists of 100 questions to
309 determine the big five personality traits; neuroticism, extraversion, openness, agreeableness
310 and conscientiousness. Social desirability was measured using the Marlowe Crowne Social
311 Desirability Scale⁽³⁶⁾, a 33-item questionnaire assesses whether or not respondents are
312 concerned with social approval, and the Balanced Inventory of Desirable Responding
313 (BIDR)⁽³⁷⁾, which is a 40 item questions that measure the tendency to give socially desirable
314 responses on self-reports (each item is scored 1 to 7 on a true or false scale). Psychometric
315 eating behaviours were assessed using the Dutch Eating Behaviour Questionnaire [DEBQ]⁽³⁸⁾.
316 The DEBQ is a 33-item questionnaire that uses a 5-point Likert scale ranging from 1 (seldom)
317 to 5 (very often) to assess three eating behaviour domains: restrained eating (10 items),
318 emotional eating (13 items) and the external eating (10 items).

319

320

Table 1 here

321

322 **2.5 Statistical Analyses**

323 Data are reported as mean \pm SD. Statistical analyses were performed using IBM SPSS
324 (Chicago, Illinois, Version 25). Two-sided paired t-tests were used to examine differences in
325 EI between the LWI method and self-report methods. Discrepancies between measured and
326 reported EIs were displayed using Bland-Altman plots (mean bias and upper and lower 95%
327 limits of agreement). In Study One, the effect of being observed on feeding behaviour (the
328 observation effect) was quantified by comparing LWIs during covert and overt phases using
329 two-sided paired t-tests. The difference between what people actually ate and what they
330 reported eating (the reporting effect), was quantified by comparing the difference between the
331 measured LWI during the overt days and the self-reported intakes using two-sided paired t-
332 tests.

333

334 A two-stage approach was taken to the analyses of the potential correlates of mis-reporting.
335 Firstly, we examined the associations between individual psychological traits and mis-
336 reporting using separate multiple regression models (while controlling for age, sex and
337 percentage body fat), and secondly, we included all of the individual predictors found to be
338 significant in a subsequent stepwise regression model to examine the overall predictive ability
339 of any significant predictors identified. Multiple linear regressions were used to examine if
340 mis-reporting (i.e. the discrepancy between actual food intake and reported food intake) was
341 associated with personal (age, sex and percentage body fat) and selected dimensions of
342 personality and eating behaviour traits (personality, social desirability, body image, IQ, mood,
343 and eating behaviours). To account for potential confounding, age, sex and percentage body
344 fat (% BF) were included in all models. Including BMI rather than percentage body fat did not
345 change any of the reported outcomes. Regression analyses are summarised in the Results
346 Section, and individual model parameters are reported in the Supplementary Materials
347 (Supplementary Tables S1-S24). Benjamini & Hochberg false discovery rate (FDR) adjusted
348 q-values⁽³⁹⁾ were calculated using the regression coefficients in models where significant
349 predictors were identified due to the multiplicity of comparisons presented (R Studio, Version
350 1.2.5042, RStudio, Inc.).

351

352 In Study One, to examine the predictors of the observation effect, differences between covert
353 and overt LWIs were regressed against personal and psychological characteristics (Section
354 3.2.1). To examine for predictors of the reporting effect, differences between the LWI on overt

355 days and each self-reported measure of intake were regressed against personal and
356 psychological characteristics (Section 3.2.2). For the laboratory phase of Study Two, the
357 discrepancy between the LWI and the self-reported intakes were regressed against personal and
358 psychological characteristics. In the home phase of Study Two, the discrepancy between the
359 WDR-H and the FFQ and diet history were regressed against personal and psychometric
360 characteristics (Section 3.3.1). The WDR-H was not compared to the 24-hr recall performed
361 during the laboratory phase as the timings of these measures differed. To examine the
362 predictive ability of the correlates identified in Study One and Two, data common to both
363 studies were combined, and stepwise regression was used in which all of the previously
364 identified correlates were entered as predictors (probability of F; 0.05 entry and 0.10 removal).
365 The differences between the LWI on overt days and each self-reported measure of intake were
366 used as the outcome variables (Section 3.3.2).

367

368 **3.0 RESULTS**

369 Descriptive characteristics of participants in Study One and Study Two can be found in
370 Tables 2 & 3.

371

Tables 2 & 3 here

372

373 **3.1 Extent of Dietary Mis-Reporting**

374 A summary of mean daily EI using measured and self-reported techniques can be found in
375 Table 4, and Bland-Altman plots displaying the deviations between intake measures at the
376 individual level can be found in Figure 2. When compared to the measured LWI, self-reported
377 EI was -0.6 ± 1.9 MJ/day lower ($p < 0.001$) using the WDR (Study One = -0.6 ± 1.3 MJ/day,
378 $p < 0.001$; Study Two = -0.6 ± 2.1 MJ/day, $p < 0.001$), -1.4 ± 2.3 MJ/day lower ($p < 0.01$) using
379 the 24-hr recall (Study One = -1.2 ± 1.5 MJ/day, $p < 0.001$; Study Two = -1.5 ± 2.4 MJ/day, p
380 < 0.001), -2.4 ± 3.7 MJ/day lower ($p < 0.001$) using the 7-day diet history (Study One = $-1.8 \pm$
381 2.4 MJ/day, $p < 0.001$; Study Two = -2.6 ± 4.0 MJ/day, $p < 0.001$), and -1.2 ± 4.2 MJ/day lower
382 ($p < 0.001$) using the FFQ (Study One = -0.3 ± 3.6 MJ/day, $p = 0.492$; Study Two = -1.4 ± 4.4
383 MJ/day, $p < 0.001$).

384

385

Figure 2 here

386

Table 4 here

387

388 **3.2 STUDY ONE OUTCOMES**

389 EI during the overt phase was significantly lower than the covert phase (10.9 ± 2.7 vs $11.6 \pm$
390 2.9 MJ/d; $p < 0.001$). This discrepancy, termed the observation effect, reflects the effect of
391 being observed on feeding behaviour. To quantify the difference between what people actually
392 ate and what they reported eating, the measured LWI during the overt days were compared to
393 self-reported intakes. This difference is referred to as the reporting effect. Compared to the
394 measured LWI, self-reported intake was significantly lower using the WDR (-0.6 ± 1.3 MJ/d;
395 $p < 0.001$), 24-hr recall (-1.2 ± 1.5 MJ/d; $p < 0.001$), 7-day diet history (-1.8 ± 2.4 MJ/d; $p <$
396 0.001), FFQ (-0.3 ± 3.6 MJ/d; $p = 0.492$) and FFQ² (i.e. intake over the 14 day residential
397 period; -1.2 ± 2.6 MJ/d; $p < 0.001$).

398 **3.2.1 Correlates of the Observation Effect**

399 After controlling for age, sex and %BF, those with lower EPQR psychoticism ($\beta = 0.389$; $p =$
400 0.009) reduced energy intake on overt days to a greater extent as compared to covert days.
401 However, the FDR correct p-value for EPQR psychoticism was non-significant ($q = 0.063$).
402 Those with higher NEO PIR agreeableness ($\beta = -0.303$; $p = 0.038$) and lower NEO PIR
403 openness to experience ($\beta = 0.440$; $p = 0.006$) also reduced EI on overt days to a greater extent
404 as compared to covert days. While the association between NEO PIR openness to experience
405 and the observation effect remained significant after FDR adjustment ($q = 0.048$), the NEO
406 PIR agreeableness adjusted p-value was non-significant ($q = 0.152$). Age, sex, %BF, eating
407 behaviour traits, body image, social desirability, IQ and mood were not associated with
408 observation effect (Supplementary Tables S1 to S8).

409

410 **3.2.2 Correlates of the Reporting Effect**

411 Lower NART performance IQ was associated with greater underreporting of EI using the WDR
412 as compared to the LWI after accounting for age, sex and % BF ($\beta = 4.072$; $p = 0.036$), but this
413 did not remain significant after FDR adjustment ($q = 0.288$). Sex ($\beta = -0.564$; $p = 0.001$), %
414 body fat ($\beta = -0.664$; $p = 0.001$) and DEBQ emotional eating ($\beta = -0.350$; $p = 0.044$) were
415 associated with the discrepancy between the LWI and 24-hr recall. Males, those with greater
416 %BF or emotional eating demonstrated greater underreporting of EI using the 24-hr recall as
417 compared to the LWI. Sex ($q = 0.001$) and % body fat ($q = 0.001$) remained significant after
418 FDR adjustment, but the FDR adjusted p-value for emotional eating was non-significant ($q =$
419 0.088). After accounting for age, sex and %BF, higher EPQR psychoticism ($\beta = -0.338$; $p =$
420 0.024) and NEO PIR openness to experience ($\beta = -0.335$; $p = 0.044$) were associated with

421 greater underreporting using the diet history as compared to the LWI. However, the FDR
422 adjusted p-value for EPQR psychoticism ($q = 0.168$) and NEO PIR openness to experience (q
423 $= 0.352$) were non-significant. Males also demonstrated greater underreporting using the FFQ²
424 as compared to the LWI ($\beta = -0.447$; $p = 0.012$), and this remained significant after FDR
425 adjustment ($q = 0.036$). No other significant associations were found for personal
426 characteristics, eating behaviour traits or personality traits, social desirability, body image, IQ
427 or mood (Supplementary Tables S9 to S16).

428

429 **3.3 STUDY TWO OUTCOMES**

430 During the laboratory phase of Study Two, the self-reported WDR was 0.6 ± 2.1 MJ/day lower
431 than the LWI ($t_{(181)} = 3.726$, $p < 0.001$). In turn, the WDR in the home phase was 1.0 ± 2.9
432 MJ/day lower than the WDR during the laboratory phase ($t_{(180)} = 4.620$, $p < 0.001$; Figure 3).
433 This difference in the WDR between laboratory and home environments was associated with
434 %BF ($\beta = 0.274$; $p = 0.010$). However, no further associations were seen between this
435 difference and sex, age, eating behaviour, body image, personality, social desirability, IQ or
436 mood.

437

438

Figure 3 here

439

440 **3.3.1 Correlates of Mis-Reporting under Laboratory and Home Environments**

441 **Laboratory Phase**

442 When the discrepancy between the measured LWI and the self-reported techniques was
443 regressed against personal characteristics, sex was associated with the discrepancy between the
444 LWI and WDR ($\beta = -0.214$; $p = 0.029$), 24-hr recall ($\beta = -0.297$; $p = 0.002$) and the 7-day diet
445 history ($\beta = -0.188$; $p = 0.050$), with mis-reporting greater in men than women (Supplementary
446 Tables S17 to S24). After FDR adjustment, the p-value for sex remained significant for the 24-
447 hr recall ($p = 0.006$), but not for the WDR ($p = 0.087$) or the 7-day diet history ($p = 0.150$).
448 Lower NEO PIR neuroticism ($\beta = 0.186$; $p = 0.022$), higher NEO PIR openness to experience
449 ($\beta = -0.218$; $p = 0.028$) and higher BDR self-deceptive enhancement ($\beta = -0.161$; $p = 0.048$)
450 were associated with a greater underreporting using the 24-hr recall as compared to the LWI
451 (after accounting for age, sex and %BF). However, NEO PIR neuroticism ($q = 0.077$), openness
452 to experience ($q = 0.077$) and BDR self-deceptive enhancement ($q = 0.144$) were not
453 significant after FDR adjustment. Higher EPQR extraversion ($\beta = -0.164$; $p = 0.032$) was
454 associated with greater underreporting using 7-day diet history as compared to the LWI, but

455 this did not remain significant after FDR adjustment ($q = 0.224$). After accounting for age, sex
456 and %BF, lower DEBQ external eating was associated with greater underreporting using the
457 FFQ as compared to the LWI ($\beta = 0.212$; $p = 0.028$), but this was not significant after FDR
458 adjustment ($q = 0.168$). IQ and mood were not associated with the discrepancy between the
459 LWI and any of the self-reported techniques (Supplementary Tables S17 to S24).

460

461 **Home Phase**

462 In the home environment, lower body image ($\beta = 0.223$; $p = 0.028$), lower DEBQ external
463 eating ($\beta = 0.214$; $p = 0.030$), higher emotional eating ($\beta = -0.231$; $p = 0.024$) and lower EPQR
464 social desirability ($\beta = 0.178$; $p = 0.024$) were associated with greater underreporting using the
465 FFQ as compared to the WDR-H. However, after FDR adjustment body image ($q = 0.089$),
466 DEBQ external eating ($q = 0.090$), emotional eating ($q = 0.090$) and EPQR social desirability
467 ($q = 0.168$) were not significant. No further associations were seen between personal
468 characteristics, personality traits, eating behaviour, social desirability, IQ or the discrepancy
469 between the WDR-H and the other self-report techniques (Supplementary Tables S17 to S24).

470

471 **3.3.2 Combined Analyses of Study One and Study Two.**

472 In order to examine the predictive ability of the correlates identified in Study One and Two,
473 data common to both studies were combined (sex, %BF, body image, external eating,
474 emotional eating, EPQR social desirability, psychoticism and extraversion, NEO PIR
475 neuroticism, agreeableness and openness to experience, BIDR self-deceptive enhancement,
476 NART performance IQ), and stepwise regression performed (probability of F; 0.05 entry and
477 0.10 removal). Sex was the only variable entered into the model when the discrepancy between
478 the LWI and the WDR ($\beta = -0.170$; $F_{(1, 225)} = 6.670$, $\text{adj-R}^2 = 0.025$, $p = 0.010$), 24-hr recall (β
479 $= -0.279$; $F_{(1, 225)} = 18.841$, $\text{adj-R}^2 = 0.073$, $p < 0.001$), and 7 day history ($\beta = -0.217$; $F_{(1, 224)} =$
480 11.033 , $\text{adj-R}^2 = 0.043$, $p = 0.001$) were examined. When the discrepancy between the LWI
481 and FFQ was examined, %BF was the only variable entered into the model ($\beta = 0.223$; $F_{(1, 224)}$
482 $= 11.717$, $\text{adj-R}^2 = 0.046$, $p = 0.001$).

483

484 **4.0 DISCUSSION**

485 The present paper examined the psychological correlates of mis-reporting under laboratory and
486 free-living conditions using two separate studies designed *a priori* to examine the nature and
487 extent of dietary mis-reporting^(19; 21). The design of these studies allowed the extent of under or
488 over-reporting to be directly quantified via comparisons between covertly measured food

489 intake and that self-reported using a range of common dietary assessment techniques. These
490 data were collected alongside a large amount of psychometric data under conditions more
491 rigorous than typically possible in free-living studies. Despite these methodological strengths,
492 there was little evidence of robust psychological correlates of mis-reporting. Sex and selected
493 personality and eating behaviour traits were correlated with mis-reporting, but these associated
494 were not consistent across studies or dietary assessment types, and explained little of the
495 variance in mis-reporting (typically <5%). The lack of robust and consistent correlates suggests
496 that personal or psychological characteristics have little utility in predicting the extent of mis-
497 reporting, even when mis-reporting is directly quantified.

498

499 **4.1 Effect of Measurement Technique and Study Environment on Energy Intake**

500 When food intake was measured under laboratory conditions in which energy balance and
501 feeding behaviour were measured continuously for 12 days (Study One), self-reported EI was
502 5-21% lower than measured intake depending on the self-report technique used. The extent of
503 under-reporting was greater for the dietary recall and the FFQ as compared to the WDR
504 method. While the mean bias using the FFQ was relatively small, examination at the individual
505 level indicated significant under and over reporting (Figure 3). In Study Two where mis-
506 reporting was measured under laboratory conditions and free-living environments, results
507 revealed the same degree of mis-reporting in the laboratory phase as in Study One. However,
508 relative to the laboratory, mis-reporting increased further in the home environment, with EI
509 lower in the home environment than reported in the laboratory environment.

510

511 **4.2 Correlates of the Observation and Reporting Effect (Study One)**

512 While the mis-reporting of energy and nutrient intake using self-report techniques has long
513 been documented⁽⁴⁰⁾, this has not led to *a priori* techniques that allow the identification of those
514 likely to mis-report or the extent to which an individual will mis-report. A number of purported
515 correlates of mis-reporting have previously been suggested, but these are inconsistent between
516 studies and typically have little explanatory value^(7; 10; 15). This may in part reflect the use of
517 proxy measures of mis-reporting (i.e. indices of energy requirements or expenditure to estimate
518 the degree of low or high energy reporting with the assumption that individuals are in energy
519 balance) rather than direct comparisons between ‘true’ and self-reported intake. To address
520 this, mis-reporting was directly quantified in the present study and potential correlates were
521 examined separately for the observation and reporting effect.

522

523 When the observation effect was examined, lower psychoticism and openness to experience
524 and higher agreeableness were associated with a greater reduction in EI on days when
525 participants knew food intake was being measured (i.e. overt vs. covert days). Age, sex and %
526 BF, or any of the other psychological measures, were not correlated with the observation effect.
527 Personality traits have previously been reported to correlate with dietary mis-reporting⁽⁷⁾, but
528 in the present study, the amount of variance in the observation effect explained by personality
529 traits was small and of little predictive value after adjusting for potential confounders (<5%).
530 Furthermore, these associations typically became non-significant after FDR adjustment. When
531 the reporting effect was examined, sex was found to be associated with the discrepancy
532 between the LWI and both the 24-hr recall and FFQ² (i.e. intake over the 14-day residential
533 period), with males under-reporting to a greater extent than females. No associations were seen
534 between sex and the WDR, 7-day diet history or FFQ. Isolated associations were also seen
535 between the LWI and selected self-report methods, but there appeared to be no consistency
536 between the self-reported measurement techniques. Furthermore, while some of the same
537 personality traits were correlated with both the reporting and observation effect (e.g.
538 psychoticism and openness to experience), it should be noted that the direction of these
539 associations differed between mis-reporting states, and again, these associations often became
540 non-significant after accounting for multiplicity of comparisons. The reported associations
541 should therefore be interpreted with caution as isolated values occurring amongst multiple
542 comparisons are likely of limited significance. Taken together, these data indicate that both the
543 reporting and the observation effect are difficult to predict from the personal and psychological
544 characteristics used in this study even under the controlled residential condition of Study One.
545

546 **4.3 Correlates of Mis-Reporting under Different Study Environments (Study Two)**

547 It was also interesting to note in Study Two EI using the WDR was lower in the home phase,
548 with the EI:RMR in the home environment 1.58 vs 1.75 the laboratory environment (using the
549 WDR as the reference values of EI). While this could be taken to suggest that mis-reporting
550 was greater in the home environment, it should be noted that i) the WDR measured in the
551 laboratory and home phases were measured at different time points, and, ii) ‘true’ intake was
552 not measured in this phase so a comparison between true intake and self-reported intake cannot
553 be made in the same way as Study One. While this limits direct comparison, it is possible that
554 the residential nature of the laboratory phase, with fewer of the usual day-to-day distractions,
555 may have increased the completeness of food recording during this phase of the study and
556 limited mis-reporting of EI in the laboratory. It is also note Therefore, future studies should

557 further examine the effect of the eating environment, as well as the dietary assessment tool, on
558 the extent of mis-reporting. As was the case in Study One, sex was found to be associated with
559 the degree of mis-reporting between the LWI and 24-hr recall, WDR and 7-day diet history in
560 the laboratory environment, with males mis-reporting to a greater extent than females.
561 Furthermore, several psychological traits, namely neuroticism, openness to experience,
562 agreeableness, extroversion and external eating, were related to mis-reporting in the laboratory
563 environment when EI was self-reported. Again however, caution must be taken when
564 interpreting these isolated associations given the size and complexity of the dataset, and the
565 multiplicity of comparisons. Indeed, these association often did not remain significant after
566 FDR adjustment, the extent to which these psychometric traits predicted mis-reporting in the
567 laboratory phase of Study Two was again extremely limited (typically <5% of the variance in
568 mis-reporting), and the correlates of mis-reported differed between the laboratory and home
569 environments as well as self-report measurement techniques.

570

571 It is interesting to note that in these data males mis-reported to a greater extent than females,
572 while there was also an apparent lack of association between personal characteristics such as
573 age and % body fat and mis-reporting. It has previously been reported that females and those
574 with a higher BMI, as a proxy measure of body fat, are more likely to under-report. However,
575 despite the wealth of studies examining both the extent, prevalence and correlates of mis-
576 reporting using self-reported techniques, results remain inconsistent^(7; 15). For example, while
577 some studies find that women under-report EI more often than men^(41; 42; 43; 44), others have
578 found under-reporting to be higher in males^(45; 46) or there to be no association with sex⁽⁴⁷⁾. It is
579 also worth noting that due to their greater body size, energy requirements in men was ~20%
580 higher than women. This was reflected in greater absolute EI in males, and therefore greater
581 mis-reporting (in absolute terms) may in part reflect a body size effect. Numerous studies have
582 reported an association between higher BMI and an increased likelihood of under-reporting
583 when compared to estimated energy requirements, such as estimated RMR. However, RMR is
584 often estimated using linear regression equations, which tend to over-estimate RMR at higher
585 body weights. Over-estimating RMR will lower the ratio of reported energy intake to RMR,
586 and result in subjects with higher BMIs being more likely to be incorrectly identified as under-
587 reporters than are lean subjects.

588

589 The apparent lack of associations between personal and psychological traits and mis-reporting
590 in the present study may also reflect the fact that participants in Study One and Study Two

591 were stratified for age, sex and BMI. This is of particular importance as potential psychological
592 correlates of mis-reporting (e.g. personality and eating behaviours traits) are known to covary
593 with age, sex and body weight/composition. Age and BMI are also often used as independent
594 predictors of mis-reporting, but in the populations concerned age and BMI almost always co-
595 vary. Given the large amount of psychometric data collected as part of Study One and Two,
596 these data suggest that mis-reporting behaviours do not appear to aggregate into discrete
597 clusters amongst people. When such factors are considered alongside the marked heterogeneity
598 in study design and populations used, and the methods used to assess both of dietary intake and
599 misreporting and the significant methodological limitations inherent to these, it is not perhaps
600 surprising previous findings are inconsistent.

601

602 **4.4 Can Mis-Reporting be predicted based Personal or Psychological Characteristics?**

603 Findings from the two studies presented here indicate that it is difficult to predict mis-reporting
604 based on either personal characteristics or psychological traits. While some correlates of mis-
605 reporting were seen, the strength of these associations was too low to enable reliable prediction.
606 Indeed, when data were combined across studies, the only consistent predictor across the
607 dietary assessment methods was sex, but only ~5% of the variance in the discrepancy between
608 the LWI and the WDR, 24-hr recall or 7-day history was accounted for by sex. It may be that
609 these variables truly contain no predictive value, or that their small effects are overwhelmed in
610 these studies by random variation in food intake. When this is considered alongside the fact
611 that mis-reporting is normally distributed, with virtually all participants exhibiting some degree
612 of mis-reporting⁽²¹⁾, mis-reporting as a phenomenon appears to be very difficult to predict at
613 the individual level even when all of its components are precisely and accurately measured
614 (which, in itself, is often very difficult under free-living conditions). Given the small amount
615 of variance the personal and psychological traits accounted for in the present study, and the
616 fact that associations differed between dietary assessment techniques, our interpretation is that
617 it is not possible to use these traits to develop models that will predict with any certainty who
618 will mis-report, and to what extent they will mis-report. It seems almost everyone exhibits mis-
619 reporting to some degree, and the underlying personal, behavioural and psychological traits do
620 not aggregate into discrete clusters amongst people, making them difficult to predict. While
621 subject traits are often related to either low energy reporting or mis-reporting (e.g. sex and
622 BMI), these relationships are often far too tenuous to use these traits to account for more than
623 a few percent of the variance in mis-reporting. It should be noted that socioeconomic level,

624 which has previously been shown to be associated with dietary mis-reporting⁽⁷⁾, was not
625 measured in-depth or included in the analysis of the present study.

626

627 **4.5 Limitations**

628 As compared to previous studies⁽⁷⁾, the extent and magnitude of under reporting in the present
629 study was smaller. This may reflect the design of the two studies, with the residential nature of
630 the laboratory phases reducing the usual day-to-day distractions and increasing the
631 completeness of food recording for example. Furthermore, in both studies the 24-hr recall was
632 performed the day after the WDR. As the 24-hr recall method is memory based, it is possible
633 that the WDR acted to prime participants and improve the accuracy of the subsequent 24-hr
634 recall. The analyses of the present paper were also limited to discrepancies in the reporting of
635 EI, with mis-reporting of specific nutrient intakes not considered here. While there is some
636 evidence of macronutrient specific mis-reporting^(7; 15; 41), and that some food groups tended to
637 be under-reported to a greater extent than did others in Study One⁽⁴⁸⁾, the personal or
638 psychological factors reported in the present data failed to predict mis-reporting of
639 carbohydrate, fat and protein intake (data not reported). It should also be acknowledged that
640 while the WDR and 24-hr recall techniques used in Study One and Two, and the FFQ² in Study
641 One, provided direct self-assessment of EI on the same days in which food intake was covertly
642 measured (LWIs), the 7-day diet histories and FFQ reflected a participant's habitual intake.
643 FFQs are more commonly used in dietary surveys to quantify patterns of dietary intake rather
644 than absolute energy or nutrient intakes. Thus, it is not perhaps surprising mis-reporting of EI
645 relative to the LWIs was evident with these tools. During the laboratory phases of each study
646 every effort was made to provide an environment in which participants habitual physical
647 activity and (eating patterns) could be replicated. Participants were able to move freely around
648 the unit and associated grounds (under supervision of a member of staff) and were free to leave
649 the unit during the study (but were accompanied and observed by a member of staff at all
650 times). Despite this, it is unlikely that physical activity and food intake reflected true free-living
651 habitual patterns. While participants were in a slight positive energy balance in both studies, it
652 is noted that in Study One total daily energy expenditure was measured using doubly labelled
653 water⁽²¹⁾ and the mean daily PAL was 1.69 x RMR. This is similar to those seen in modern
654 Western populations when energy expenditure is measured using doubly labelled water under
655 free-living conditions. By design, the home phase of Study Two was more representative of
656 their habitual feeding environment, but as a result this phase was less controlled, and it is
657 unknown whether illnesses or special events for example influenced the reported intakes.

658

659 **4.6 Conclusions**

660 While selected personal and psychological traits were associated with mis-reporting, these
661 associations displayed no clear pattern across studies or dietary assessment technique and had
662 little utility in predicting mis-reporting. Even when mis-reporting is directly quantified under
663 robust experimental conditions (that exceed the level of control likely to be achieved in free-
664 living studies), it appears difficult, if not impossible, to predict mis-reporting based on personal
665 or psychological characteristics. It is therefore recommended that wherever possible, EI should
666 be studied in the context of energy balance. Indeed, there is increasing focus on using intake-
667 balance methods and mathematical models to estimate energy intake from energy expenditure
668 and changes in stored energy. While not providing information on macro-nutrient intake, these
669 approaches provide the only current objective quantitative framework in which to measure the
670 impact of mis-reporting of EI, and avoids cross-validation of self-report techniques. It also
671 offers a context in which new biomarkers of energy and nutrient balance can be developed,
672 using metabolomic approaches, to further improve the measurement of energy and nutrient
673 balance.

674

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800 **FIGURE LEGENDS**

801

802 **Figure 1:** Schematic overview of Study One (Panel A) and Study Two (Panel B) design. MTD, maintenance diet. LWI, laboratory weighed intake. In Study One, covert measurement of food intake was made using the laboratory weighed intake method across all days, while participants self-reported food intake during overt phases only. Order of covert and overt phases was randomised. In Study Two, covert measures of food intake were made using the laboratory weighed intake and self-report methods during the laboratory phase. Food intake was measured using daily weighed dietary records in the home phase, and the order of the home and laboratory phases was randomised. LWI, laboratory weighed intake. WDR, weighed dietary record. FFQ, food frequency questionnaire. UWIST, UWIST Mood Adjective Checklist; IQ, intelligence quotient; NART, National Adult Reading Test; AH4, Alison Heim 4; Raven, Raven Standard Progressive Matrices; EPQR, Eysenck-100; NEOPIR, Neuroticism, Extraversion, Openness Personality Inventory-Revised; BIDR, Balanced Inventory of Desirable Responding.

815

816 **Figure 2:** Bland-Altman plots illustrating the difference between mean daily energy intake using the laboratory weighed intake method and the weighed dietary record method (a), 24-hr recall (b), 7-day diet history (c) and food frequency questionnaire (d) against the mean of the two measures. The dashed horizontal line represents the mean bias between the two methods, and the two dotted horizontal lines represent the upper and lower 95% limits of agreement. LWI, laboratory weighed intake. WDR, weighed dietary record. FFQ, food frequency questionnaire.

823

824 **Figure 3:** Effect of the study environment on reported energy intake measured using the weighed dietary record under laboratory and home environments of Study Two (n = 181; men = 86, women = 96). Data are mean \pm SD. *Significant difference (two-sided paired t-test) between energy intake measured using the weighed dietary record under laboratory and home environments (p < 0.05). WDR, weighed dietary record.

829 Table 1: Psychological questionnaires used in Study One and Two, and the specific day(s) of completion.

	Study One	Study Two
HOME PERIOD		
Mood		
- UWIST	-	2-5
LABORATORY PERIOD		
Mood		
- UWIST	3-14	2-3
IQ		
- NART	3	2
- AH4	4	2
- Ravens	3	3
Personality		
- Body image questionnaire	9	2
- EPQR	13	2
- NEO PIR	15	3
Social desirability		
- Marlowe Crowne Social Desirability Scale	7	2

- BIDR	7	3
Eating behaviour		
- Dutch eating Behaviour Questionnaire	1	1

830 UWIST, UWIST Mood Adjective Checklist; IQ, intelligence quotient; NART, National Adult Reading Test; AH4, Alison Heim 4; Raven,
831 Raven Standard Progressive Matrices; EPQR, Eysenck-100; NEOPIR, Neuroticism, Extraversion, Openness Personality Inventory-Revised;
832 BIDR, Balanced Inventory of Desirable Responding.
833

834 **Table 2:** Descriptive characteristics of subjects (Study One).

	Total Sample (n = 59)		Men (n = 30)		Women (n = 29)	
	Mean	SD	Mean	SD	Mean	SD
Age, yrs	42.7	13.6	42.9	13.1	42.5	14.3
Height, m	1.71	0.1	1.76	0.1	1.65	0.1
BMI, kg/m ²	26.1	3.8	26.7	4.0	25.4	3.5
Weight, kg	75.9	14.3	82.7	14.5	68.9	10.3
Body fat, %	32.2 ^a	7.0	28.4	6.2	36.2	5.4
RMR, MJ/d	6.56	1.23	7.20	1.17	5.90	0.91
Education	42% secondary		33% secondary		52% secondary	
Level (%)	58% tertiary		67% tertiary		48% tertiary	

846 RMR, resting metabolic rate. Note, % body fat estimated from skinfold thickness using the equations of Durnin & Womersley⁽²⁴⁾. ^an = 57.

847 **Table 3:** Descriptive characteristics of subjects (Study Two).

	Total Sample (n = 182)		Men (n = 86)		Women (n = 96)	
	Mean	SD	Mean	SD	Mean	SD
Age, yrs	42.4	12.2	41.2	12.1	43.3	12.3
Height, m	1.70 ^a	0.1	1.77 ^b	0.1	1.63 ^c	0.1
BMI, kg/m ²	25.7	3.9	26.1	3.7	25.4	4.0
Weight, kg	74.6	14.1	82.1	13.6	67.8	10.8
Body fat, %	30.2 ^a	8.2	24.9 ^b	7.0	34.8 ^c	6.0
RMR, MJ/d	6498	121	7286	1184	5755	845
Education	31% secondary		31% secondary		31% secondary	
Level (%)	69% tertiary		69% tertiary		69% tertiary	

859 RMR, resting metabolic rate. Note, % body fat estimated from skinfold thickness. ^an = 179. ^bn = 84. ^cn = 95.

860 Table 4: Measured and self-reported mean daily energy intake values for Study One, Study Two and the total sample combined.

	Total Sample (n = 241)		Study One (n = 59)		Study Two (n = 182)	
	Mean	SD	Mean	SD	Mean	SD
LWI overt phase (MJ/day)	11.6	3.8	10.9	2.7	11.8	4.1
Laboratory WDR (MJ/day)	11.0	3.5	10.3	2.6	11.2	3.7
Home WDR (MJ/day)	N/A	N/A	N/A	N/A	10.2	3.1
24-hr Recall (MJ/day)	10.2	3.3	9.7	2.3	10.3	3.6
7-day Diet history (MJ/day)	9.2 ^a	3.5	9.1	3.3	9.2 ^c	3.6
FFQ (MJ/day)	10.4 ^a	3.9	10.6	4.0 ^b	10.4	3.9

861 LWI, laboratory weighed intake. WDR, weighed dietary record. FFQ, food frequency questionnaire. N/A, measure not taken during this particular
 862 experimental phase. ^an= 240; ^bn = 58; ^cn = 181.