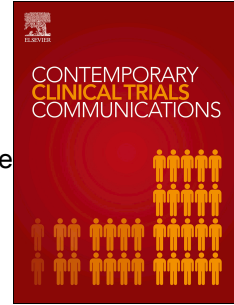


Journal Pre-proof

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PII: S2451-8654(21)00146-0

DOI: <https://doi.org/10.1016/j.conctc.2021.100846>

Reference: CONCTC 100846

To appear in: *Contemporary Clinical Trials Communications*

Received Date: 19 April 2021

Revised Date: 17 August 2021

Accepted Date: 6 September 2021

Please cite this article as: T. Redpath, F. Naseer, R.K. Price, A. Boyd, M. Martin, C.W. le Roux, A.C. Spector, M.B.E. Livingstone, Evaluation of the impact of gastric bypass surgery on eating behaviour using objective methodologies under residential conditions: Rationale and study protocol, *Contemporary Clinical Trials Communications* (2021), doi: <https://doi.org/10.1016/j.conctc.2021.100846>.

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Evaluation of the impact of gastric bypass surgery on eating behaviour using objective methodologies under residential conditions: rationale and study protocol

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Ethical approval: This study was approved by the West of Scotland Research Ethics Service (WoSRES) (REC 16/WS/0056, IRAS 200567) (Clinical trial number: NCT03113305)

Acknowledgements: Research reported in this publication was supported in part by the US-Ireland Research and Development Partnership program through the Health and Social Care

R&D Division of Northern Ireland and the Medical Research Council (STL/5062/14), Health Research Board of the Republic of Ireland (USIRL-2006-), and the National Institute of Diabetes and Digestive and Kidney Diseases of the National Institutes of Health (R01DK106112). The content is solely the responsibility of the authors and does not necessarily represent the official views of the funding agencies.

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Abstract

Gastric bypass surgery leads to significant and sustained weight loss and a reduction in associated health risks in individuals with severe obesity. While reduced energy intake (EI) is the primary driver of weight loss following surgery, the underlying mechanisms accounting for this energy deficit are not well understood. The evidence base has been constrained by a lack of fit-for-purpose methodology in assessing food intake coupled with follow-up studies that are relatively short-term. This paper describes the underlying rationale and protocol for an observational, fully residential study using covert, objective methodology to evaluate changes in 24-hr food intake in patients (n=31) at 1-month pre-surgery and 3-, 12- and 24-months post-surgery, compared to weight-stable controls (n=32). The main study endpoints included change in EI, macronutrient intake, food preferences, and eating behaviours (speed, frequency, and duration of eating). Other physiological changes that may influence EI and weight regulation including changes in body composition, circulating appetite hormones, resting metabolic rate, total energy expenditure and gastrointestinal symptoms were also evaluated. Understanding which mechanisms contribute to a reduction in EI and weight loss post-surgery could potentially help to identify those individuals who are most likely to benefit from gastric bypass surgery as well as those that may need more targeted intervention to optimise their weight loss post-surgery. Furthermore, clarification of these mechanisms may also inform targeted approaches for non-surgical treatments of obesity.

Key words: Gastric bypass; study protocol; objective validation; food intake

1. Introduction

Gastric bypass surgery is a safe, effective treatment for individuals with severe obesity [1] and leads to improvements in associated conditions including type 2 diabetes mellitus (T2DM) [2] and cardiovascular disease [3]. The most frequently performed procedure is the Roux-en-Y gastric bypass (RYGB), and more recently the One-Anastomosis gastric bypass (OAGB) which are equally effective for both weight loss [4] cardiovascular and quality-of-life outcomes [5] [6].

The exact mechanisms underlying the profound weight loss and subsequent weight maintenance remain elusive and involve a complex interaction between physiological, psychological, and behavioural factors. Although a decrease in energy intake (EI) is the main driver of weight loss [7] [8] [9] [10] this cannot be fully explained by purely restrictive and malabsorptive mechanisms [11] [12]. Other proposed mechanisms include changes in hunger and satiety [13] [14] caused by changes in circulating gut hormones [15] [16], changes in eating patterns such as reduced meal sizes without compensatory increases in meal frequency or duration [17] [18], or shifts in dietary energy density (ED) [19] resulting from changes in food selection and/or changes in food preferences [20] [21]. Change in macronutrient intakes and the associated impact on EI is a particularly contentious issue. Evidence from animal studies suggest that there is a post-surgical decrease in fat and sugar intakes [22] [23] [24] [25]. However, the evidence from human studies regarding changes in relative macronutrient intake in the short-term is equivocal [26], with studies variously reporting a decrease [20] [22] [27] or no change [18] [28] in the intake of high fat/high sugar foods.

The elucidation of the underlying mechanisms of post-operative weight loss has been severely hampered by inconsistencies in bariatric research methodology and further compounded by differences in the analysis, interpretation, and presentation of results [29] [30] [31]. In

particular, there has been overwhelming reliance on and acceptance of the purported validity of subjectively reported food intake and food preference data in bariatric research without proper acknowledgement that biased food intake data are a fundamental obstacle in understanding the dynamics of food selection and intake [32]. To date, only one research group has objectively observed food intake behaviour in a bariatric surgery population [33] [34] [35]. The observed reduction in EI was not macronutrient specific and was accounted for by consumption of smaller portion sizes of the same foods that were consumed pre-surgery. However, these potentially significant and independently validated findings are confined to one eating event which limits their extrapolation. Further verification of these findings is required across multiple eating events.

The integrity of the existing evidence base is further constrained by the frequency and duration of study follow-up. Most of the current evidence regarding shifts in post-operative EI is based on short-term (up-to 12 months post-surgery) and/or single time point studies. However, this is the stage when patients are losing significant weight and it is inconceivable that these studies will capture the dynamics of food intake behaviour and subsequent impact on the longer-term weight trajectory.

Full clarification of the mechanisms underpinning the dynamics of food selection and intake post-surgery can only be resolved by the application of fit-for-purpose methodology and reporting criteria. Consequently, the overall aim of this research was to evaluate the transition in food intake in patients pre- and post- gastric bypass surgery during a dynamic phase of weight change through covert and objective tracking of food intake and eating behaviours assessed under fully residential conditions. In addition, associations with Resting Metabolic Rate (RMR), free-living total energy expenditure (TEE), appetite hormones, and body composition were evaluated. It was hypothesised that the interplay between the various

dimensions of dietary intake, eating behaviour, energy expenditure, and gut hormone responses are key in driving both weight loss and longer-term weight regulation.

2. Methods

This observational study was conducted on patients scheduled to undergo gastric bypass surgery and time-matched, weight-stable controls under fully residential conditions at 4 time points (1-month pre-surgery and 3-, 12-, and 24-months post-surgery) to evaluate changes in energy and macronutrient intake, eating behaviours (eating speed (g/min, kJ/min), timing of eating) and food preferences over 2 years following surgery. Concurrent changes in circulating appetite hormones, body composition and RMR were also assessed and, in addition, free-living TEE and patterns of physical activity (PA) were evaluated in a subset of patients.

2.1. Participants

Patients scheduled to undergo gastric bypass surgery (n=34) and weight-stable controls (n=32) were recruited. Patients were referred for either RYGB or OAGB at several hospitals/health trusts across the United Kingdom (UK) and Republic of Ireland (ROI). Patients recruited in England were referred for surgery (provided by the National Health Service) by their General Practitioner whereas those recruited in Northern Ireland were self-referred and having their treatment privately. The referral criteria for both groups followed the UK guidelines, namely: a body mass index (BMI) $\geq 40\text{kg/m}^2$, or a BMI $\geq 35\text{kg/m}^2$ and an obesity-related condition (such as T2DM or high blood pressure) that might improve with weight loss and where previous weight loss methods have been unsuccessful.

Patients from ROI were recruited from a group who were clinically selected to undergo gastric bypass surgery as part of a pilot programme that offered the surgery primarily for the management of T2DM in individuals who were unable to manage the condition with lifestyle changes and medication.

Control participants were weight-stable (>6 months) individuals time-matched to the patient group and with no planned weight changes.

For all participants, the exclusion criteria were: <18years of age, pregnancy/lactation, food allergies/dietary restrictions and/or gastrointestinal conditions or medications that may affect food intake.

2.1.1. Recruitment strategy

Dietitians (UK), Ulster University researchers (UU), or researchers in Ireland (ROI) recruited patients at hospital clinics prior to surgery. During these initial screenings, a detailed explanation of the study protocol was provided and, following an expression of interest, full screening determined eligibility. The baseline (pre-surgery) study time point for the patient group was scheduled before the commencement of the requisite, energy-restricted diet prior to surgery (approximately 1-month pre-surgery).

The control group was recruited by UU researchers through word-of-mouth, social media, and via emails and posters within UU. The study time points for the control volunteers were matched with the patient group. Recruitment of all study participants took place from October 2016 to March 2018.

All participants provided fully informed written consent (REC 16/WS/0056, IRAS 200567).

To divert attention from the main purpose of the study, participants were informed that the primary purpose of the study was to measure changes in RMR following gastric bypass surgery.

At the conclusion of the study participants were debriefed on study outcomes.

2.1.2. Sample size

As this study protocol was both novel and intensive, there was no existing literature to inform a power calculation and so sample size was estimated using a randomised controlled trial (RCT) by le Roux *et al* [22]. This RCT, which assigned participants to undergo either RYGB or Vertical Banded Gastrectomy (VBG) and assessed dietary intake by self-report measures, detected significant differences in EI in 16 (VBG (n=7), RYGB (n=9)) participants at 6 years post-surgery. The sample size was calculated using the standard deviation (SD) associated with the change in dietary fat (% energy) intake from pre- to post-surgery and a 95% confidence interval as follows:

$$n=(\text{confidence level}*\text{standard deviation}/\text{margin of error})^2$$

$$n=(1.96*1.9/1)^2$$

$$n=14$$

Applying a 14% attrition rate as reported by Kenler *et al* [20] in which changes in self-reported dietary intake were reported at 2 years post-surgery it was estimated that a minimum of 16 patients should be recruited in the present study. However, given the intensity of the proposed protocol, possible participant attrition was accounted for by recruiting 32 patients scheduled to undergo gastric bypass surgery and 32 weight-stable control participants.

2.2. Study protocol

All participants were studied at 4 time points: at baseline (1-month pre-surgery) and 3 post-surgery time-points (3-, 12- and 24-months post-surgery). Figure 1 provides an overview of participant recruitment and progress.

At each of the 4 study time points, participants were required to undertake a 36hr fully residential period, starting late afternoon on day 1 and ending at lunchtime on day 3, in the

Human Intervention Studies Unit (HISU) within the Nutrition Innovation Centre for Food and Health (NICHE), Coleraine Campus, UU. This unit consists of 9 en-suite bedrooms, communal living and dining areas for participants and a closed-access kitchen (access for researchers only). All communal areas were monitored by closed circuit television cameras (CCTV) for verifying food intake and eating behaviours (timing/duration of eating, size/frequency of eating occasions, food interest, food selection, eating speed (g/min, kJ/min)). Participants were fully informed of and consented to the presence of CCTV monitoring within the unit.

All participants followed the same general protocol (Figure 2). Participants arrived at the HISU on the late afternoon/early evening of day 1 and a pre-set dinner (Spaghetti Bolognese) was provided if requested, followed by fasting from 10pm. All measurements, including the covert monitoring of 24hr food intake, began on the morning of day 2 (~7am) until bedtime (11pm). Participants remained in the unit for the duration of each study visit but had access to a range of sedentary activities including reading and crafts, with televisions in communal areas and bedrooms.

2.3. Food Provision

2.3.1. Food Choice Questionnaire

In order to ensure that the foods/beverages served were compatible with the usual food intake of each participant a food choice questionnaire was administered prior to the baseline visit. Participants rated their liking for 96 food options on a Likert scale (1-9; 1=dislike extremely, 9=like extremely) with food items listed in no particular order. These foods were chosen to be representative of 6 macronutrient (expressed as %energy) mix groups (high fat/low fat, high complex carbohydrate/low complex carbohydrate, high simple sugar/low simple sugar, high protein/low protein) (Table 1) (adapted from Geiselman *et al* [36]). Stated food choices were then used to establish the foods served to each participant. Individual participant menus

consisted of 9 food options from each of the 6 macronutrient mix groups for which participants had scored the highest hedonic response.

2.3.2. Participant menus

The **same** personalised menu of 54 foods was provided at each study time point. As far as possible the foods presented (n=54), including pack size and branding were consistent for each study visit. If foods were discontinued or modified over the course of the study, these were replaced with suitable alternatives that had similar macronutrient content and pack sizing. Sugar-sweetened and sugar-free beverages, tea, coffee, milk, and water were freely available to participants throughout the day, with condiments including salt, pepper, salad cream, tomato sauce, mayonnaise, butter, low-fat spread and mixed jams available. Foods and snacks were prepared and stored according to the manufacturer's instructions. An example of a participant menu is provided in Table 2.

A qualified chef prepared all composite evening dishes using modified (to meet macronutrient mix requirements) standardised recipes of popular savoury dishes (e.g. sweet and sour dishes, smoked haddock pie). These modified dishes underwent sensory testing prior to the start of the study to determine acceptability in relation to flavour, texture and colour. The side dishes, which accompanied the evening meal included pasta, potatoes (boiled), rice (white, boiled), salad and mixed vegetables with participants able to select any combination of these. Participants were also able to select sweet/dessert items representative of the 6 macronutrient mix groups from this menu. Desserts were recognisable, branded desserts (e.g., sugar-free jelly, apple pie) that participants were able to select in any combination. Double cream (served whipped or as pouring cream depending on participant specification) was available as an optional accompaniment for all desserts.

Foods were presented in different formats; hot and cold traditional 'breakfast' foods (n=6) were presented as a buffet, while lunch/snack foods (n=36) were available ad-libitum from each participant's assigned refrigerators and cupboard for storing non-perishable foods. Evening meals (n=12 dishes) were selected from individually tailored menus featuring hot savoury dishes (n=6) and desserts (n=6), with no restriction on the number of choices that could be made.

Participants were advised to consume only the foods provided to them and not to share food items. Researchers were not present while participants were eating. Meal and snack times were not researcher prescribed in advance, rather participants could select to eat at time(s) of their choosing.

2.4 Outcome Measures

2.4.1 Dietary Intake

The ad-libitum food intake of each participant was directly and covertly measured by weighing all foods before serving together with leftovers over a 24hr period (from participant wake up (~6am-8am) to 11pm) on day 2 of each study visit. Throughout this period, participants had ad-libitum access to foods and beverages stored in individually assigned fridges and store cupboards for non-perishable foods. Food intake was verified using the CCTV footage which also provided information on associated eating behaviours: frequency (n), duration (min), size (g), energy content (kJ) and distribution of eating occasions, eating rate (kJ/min, g/min), foods selected). All CCTV data were double-entered and checked, with a third member of the research team reconciling any discrepancies in the data entry.

Dietary intake data were calculated using a database developed specifically for this study. Estimated energy and nutrient content of foods were obtained from manufacturer websites, [37] and other food composition databases [38]. The main outcome measures were total EI

(kJ/d) and relative (%energy) macronutrient intake. Other outcome measures were macronutrient mix group contribution to overall EI (%EI), ratio of sugar:sugar free beverages, % contribution of energy-containing fluids to overall EI.

2:4:2. Dietary Energy Density

Dietary energy density (ED) (kJ/g), the amount of energy in food relative to weight, is primarily influenced by the fat and water content of foods [39]. Currently, there is no standardised definition of ED and methodological differences in its calculation can lead to inconsistencies in the interpretation of data, particularly if beverages are included in the calculation [40] [41].

This study evaluated how the application of different definitions impacted the measurement of dietary ED and study outcomes. Previous work [42] identified 8 methods for deriving ED, from which the following 4 definitions were identified as being most applicable to this study:

- Food only; solid/liquid items consumed as food.
- Food and milk; solid/liquid items consumed as food plus dairy beverages.
- Food and energy-containing beverages; solid/liquid items consumed as food plus beverages containing >21kJ/100ml.
- Food and all beverages: solid/liquid items consumed as food plus all beverages except water.

2:4:3 Definition of an eating occasion

By design, this protocol did not impose researcher- or participant-defined 'meals' and 'snacks', but instead applied the term 'eating occasion'. An eating occasion has been defined as 'an event which provides at least 210kJ with a separation in time from a preceding or following eating event of at least 15 minutes' [43]. However, this arbitrary definition has not been subjected to independent evaluation.

The CCTV data (patient (n=31) and control (n=30) group data merged) from the baseline study time point were used to determine both pause duration between eating occasions and energy content of eating occasions. A pause was operationally defined as ‘a pause in eating where a start and finish time can be clearly recorded, with termination defined as a break in eating for more than 5 seconds’. The minimum of 5 seconds was selected as this was the shortest pause time that could be clearly verified from the CCTV data. In total, 1577 pauses of ≥ 5 seconds were recorded for analyses.

All pauses were recorded on a scatter plot to examine any patterns in the data and plotted by frequency when grouped into minutes (>15 min). There were a high concentration of pauses within the first 100 seconds (63.1%), with the majority (83%) occurring within 300 seconds (5 min). A plateau occurred at approximately 5 min, suggesting that the previously defined 15-min pause may not be the most applicable for these data as eating occasions could merge. It was established that an interval of ≥ 5 min was more appropriate to apply to this data set.

To evaluate the efficacy of the definition that an eating event should provide >210 kJ, baseline CCTV data were evaluated in combination with directly measured available EI data for 61 participants (patients n=31, controls n=30). Nearly a fifth (18.7%) of all eating occasions (n=538) were <210 kJ and mainly consisted of sugar-free beverages and/or black tea or coffee.. Given the small proportion and energy content of eating occasions that fell below the 210kJ cut-off, it was concluded that only EOs ≥ 210 kJ should be analysed.

In summary, the proposed definition for an eating occasion employed in this study was ‘an eating event that provided at least 210kJ with a separation in time from a preceding or subsequent eating event of at least 5 minutes.’

2:4.4. Eating patterns

The distribution of EI across the measurement period was divided into four eating epochs: wake-up-11am, 11.01am-3pm, 3.01pm-7pm and 7.01pm-11pm. These eating epochs loosely represent the periods that would encompass traditional mealtimes in the UK and ROI (i.e., breakfast, lunch, dinner, supper) and were used to determine the distribution of eating occasions, EI, relative macronutrient intake and ED across the day.

CCTV data were also used to evaluate the frequency, duration, and size of eating occasions as well as eating rate (g/min), food interest (% of total number of visits to food storage areas when food was removed and consumed) and food selection (first food and its associated macronutrient mix composition from buffet table in eating epoch 1). Frequency/duration of eating occasions and eating rate were included in the analysis only if the start and finish time could be clearly observed.

2:4.5. Food preferences

Prior to leaving the HISU on day 3, 2 hr after breakfast and after all other dietary measurements had been completed, participants self-administered the Leeds Food Preference Questionnaire (LFPQ) [44]. This instrument has been validated in different populations [45] [46] and has previously been used to measure food preferences in individuals with obesity [47] [48].

The LFPQ is a computer-based measure of both explicit and implicit components of food preference and is a validated measure of food 'liking' (hedonic pleasure) and food 'wanting' (desire to consume). The LFPQ presented participants with a range of pre-validated pictures of common food items that are either high fat (>50% energy content) or low fat (<20% energy content) but similar in familiarity, palatability and sweet/savoury taste [49]. If participants expressed dislike for any of the foods presented these were substituted with a different food

with similar macronutrient content and taste (sweet/savoury). Any food picture substitutions were made at baseline only, then kept consistent at subsequent visits.

Explicit measures of food reward were determined by presenting participants with an image of a food item that is either high-/low-fat and sweet/savoury and requiring them to rate on a visual analogue scale either; '*How much would you like some of this food now?*' or '*How much do you want some of this food now?*'. Average responses to each category (n=4) were calculated, with a higher score representing higher explicit preference for that food category. Examples of the food pictures included chocolate (high fat/sweet), cheese (high fat/savoury), fruit salad (low fat sweet) and bread roll (low fat/savoury).

The LFPQ measured implicit wanting for food by presenting participants with a forced-choice paradigm which required them to choose between a high-fat vs. low-fat food and a sweet vs. savoury food. Participants were asked to respond quickly to the question '*Which food do you most want to eat now?*'. Responses and reaction times were subsequently used to calculate implicit wanting score, where selection and speed positively contribute to the score. Data were analysed using a frequency-weighted algorithm which has been developed to assess which foods have been avoided or selected, with non-selection negatively contributing to the implicit wanting score [49].

2:4:6. Body composition

Anthropometric measurements were made on day 2 of each study visit. Body weight was measured using the GE Lunar Dual X-Ray Absorptiometry (DXA) scanner (GE Healthcare). Participants were weighed while wearing light clothing and with no shoes or jewellery, and measurements were made to the nearest 0.1kg.

Height to the nearest 0.1cm was measured only at baseline under standardised conditions using a standing stadiometer. Participants were asked to stand on a base plate with their back against the stadiometer background with an upright spine and their feet flat and together.

BMI was calculated as $(\text{weight}(\text{kg})/\text{height}(\text{m})^2)$ and applying the BMI categories defined by using WHO cut offs (World Health Organisation, 2000).

Total weight loss (%TWL) was calculated using the following equation:

$$\%TWL = ((\text{Weight at baseline}(\text{kg}) - \text{weight at time point}(\text{kg}))/\text{baseline})*100$$

DXA scans were used to determine lean mass (LM) (kg, %) and total fat mass (FM) (kg, %) at each study time point, with the software additionally able to measure visceral fat (g, %) in participants who had a BMI $<40\text{kg}/\text{m}^2$. Bone mineral density was assessed at baseline and 24-months post-surgery.

Where body width exceeded the scanning area a half-body scan was used as a valid substitute for a whole-body scan [50] [51] to estimate body composition. Scans take between 7-15 min, depending on participant size.

Body composition was measured across multiple regions (arms, legs, trunk, android, gynoid). A qualified practitioner performed scans with outputs subsequently assessed by the same radiographer at each time point.

2:4:7. Energy expenditure

2:4:7:1. Free living total energy expenditure

Free living TEE was measured under free-living conditions over 14 consecutive days by the doubly labelled water (DLW) method [52] [53] [54] [55] in a subgroup of patients (n=7). TEE

is estimated by enriching the participant's body water with two stable isotopes: deuterium ($^2\text{H}_2$) and oxygen-18 (^{18}O) and determining the difference in elimination rate between both isotopes. The method is based on the principle that $^2\text{H}_2$ is eliminated as water, corresponding to water output, and ^{18}O exits the body as both water and expired CO_2 with the difference between the elimination rates providing a measure of CO_2 production from which TEE is calculated from classical indirect calorimetric equations.

The DLW dose for each participant was based on total body weight (0.07g/kg of 99% $^2\text{H}_2$ and 1.74g/kg of 10% ^{18}O , as advised by Iso-Analytical Limited, United Kingdom). The dose was administered orally (under supervision) followed by a 100ml regular water rinse to ensure all the labelled water was consumed. Five time-measured urine samples (5mls x 5) were collected from each participant: pre-dose, 4hr-, 24hr-, 7days-, and 14days post-dose. Samples were subsequently analysed at the Scottish Universities Environmental Research Centre (SUERC) (University of Glasgow, UK).

2:4:7:2. Resting Metabolic Rate

On waking on day 2, RMR was measured using indirect calorimetry in a well-ventilated room and with the participant in a supine position (ECAL, Metabolic Health Solutions). Participants were woken (~7am), asked to empty their bladder and rest a further 30 minutes before the measurement was made. Distractions such as use of mobile phones were not permitted. The first two minutes of the measurement period were automatically discarded by the ECAL software, with any other anomalous recordings (e.g. coughing, removal of mouthpiece) also discarded as 'false' readings. Data were recorded for a minimum of 5min, and was terminated after readings had been stable for 45 seconds.

2:4:7:3. Physical activity

Physical activity related energy expenditure (PAEE) was assessed both subjectively and objectively in the same subgroup of patients (n=7) whose TEE was measured by DLW.

Subjective assessment of PAEE was made using the Recent Physical Activity Questionnaire (RPAQ) [56] which is based on the European Prospective Investigation into Cancer and Nutrition (EPIC)-Norfolk Physical Activity Questionnaire (EPAQ2) [57] and has shown good validity for an assessing individual's PAEE [58]. The RPAQ assessed activities according to 4 domains of physical activity (home activity, work activity, commuting, leisure time) over the previous 4 weeks, with closed questions pertaining to the type, frequency, and duration of both physical and sedentary activities. This questionnaire was self-administered, and responses were coded and used to calculate the Metabolic Equivalent of Task (METs). One MET is equivalent to an individual's RMR, and METs are calculated based on activity level (sedentary (≤ 1.5 x RMR); light (1.5-2.99x RMR); moderate (3-5.99x RMR); vigorous (≥ 6.0 x RMR) [58] [59] [60], and applying the following standard formula:

PAEE = (METs x 3.5 x body weight(kg)/200) x duration of activity (minutes).

Objective assessment of PAEE was completed using an Actigraph (ActiGraphTM GT3X+, ActiGraph, Florida, USA). Actigraphs are small, lightweight activity monitors that measure triaxial activity (acceleration x magnitude x time). The monitor was worn for 7 consecutive days following each study time-point, and participants were included in the analysis if ≥ 10 hrs wear for ≥ 3 days was recorded. Measured activity was categorised into 4 intensities (sedentary, moderate, vigorous, very vigorous) [60]. Reported outcomes included daily step count, PAEE and duration per intensity category (hr/day).

2:4:8 Biochemistry

The following biochemical measurements were made to determine nutrient status (glucose, vitamin D, B6 and B12) and gut hormones (ghrelin and GLP-1). At all study time points fasted (28ml) and postprandial (8ml) blood samples (plasma EDTA and serum) were drawn on the morning of day 3 of each study visit. Postprandial blood samples were drawn 90 min after a standardised breakfast, during which participants were instructed to eat until they were comfortably full. Processing of plasma EDTA tubes was immediate (<5 min) in refrigerated centrifuges (15 min, 2600rpm, 4°C), with fasted blood glucose measurements (<15 min from draw) (Hemocue Hb 201+) and full blood counts completed on sample arrival at the laboratory (SYSMEX KX-21N). Serum tubes were coagulated at room temperature for 30 min prior to processing. All bloods samples were stored at -80°C until analysis.

2:4:9. Gastrointestinal symptoms

Self-reported gastrointestinal symptoms were assessed using a modified post-meal digestion questionnaire (PMDQ) which was developed by merging the Sigstad Clinical Diagnostic Index (SCDI) [61] and the more recent Dumping Symptom Rating Scale (DSRS) [19] [62]. The SCDI rates the intensity of 16 symptoms suggestive of Dumping Syndrome (DS). Two of the 16 symptoms (eructation/vomiting) are scored negatively to distinguish DS from other syndromes such as afferent loop syndrome or small stomach syndrome, but as vomiting is a common symptom following gastric bypass [63] [64], data were analysed with and without these negative scores. The DSRS evaluated severity and frequency of symptoms, and the assessment of symptom frequency was incorporated into the PMDQ (<once a week; once a week; several times per week; once a day; several times per day) along with additional questions on food avoidance. To determine if the gastrointestinal symptoms were experienced

during early or late DS, an additional question on the timing of symptoms (within 1hr of eating; 1-3hr after eating; both) was also incorporated into the PMDQ.

2:4:10. Medication

At each study time point medical information pertaining to any pre-existing medical conditions and details of any medications and dietary supplements taken were recorded. Information was obtained on dose, indication, frequency and start/stop dates of medications and supplements.

2:4:11. Qualitative Data

Patient experiences following gastric bypass surgery were assessed using a qualitative semi-structured interview at the final study time point. Based on the Socioecological Model (SEM) framework [65] [66] [67] [68], a series of open-ended interview questions was developed to explore the impact of gastric bypass surgery on post-operative health-related behaviours. The SEM categorises determinants of health behaviour into 3 groups: intra-personal, inter-personal and environmental. Questions were designed to elicit patient experiences following gastric bypass surgery, their current eating behaviours relative to pre-operative behaviour, the impact of surgery on personal relationships, personal and professional support received, their clinical experience and factors which were perceived to contribute to an individual's weight trajectory following surgery. Following audio-recorded interviews data were professionally transcribed verbatim and coded thematically [69] using NVivo 12 (QSR International Pty Ltd, Doncaster, Victoria, Australia). Research team members independently read and coded one randomly selected transcript to ensure the validity of the application of codes to the data.

2.5. Statistical Analysis

Statistical analyses were performed using SPSS (Statistical Package for the Social Sciences software, IBM). Available-case analysis was used. Where participants missed a study time point, missing value regression imputation was used where possible to predict results rather than exclude them from analysis. Imputed values were included only when the adjusted R square value was greater than 0.5. Continuous variables were presented as mean±SEM and categorical variables were presented as n (%) unless otherwise stated. Data were tested for normal distribution and log₁₀ transformed where necessary. Two-way mixed Analysis of Variance (ANOVA) was used to determine changes in overall group mean data (patients vs. weight-stable controls) following gastric bypass surgery and changes in mean data based on surgery type (RYGB vs OAGB). Subsequently, Bonferonni post-hoc tests (controlling for multiple comparisons) were conducted to explore all valid multiple pairwise comparisons within the dataset. Linear regression analyses were used to assess relationships between different outcome variables. Further in-depth analysis was conducted to determine individual differences in response to the surgery. Significance was considered at the $p \leq 0.05$ level.

3. Discussion

There are multiple mediators involved in weight loss following bariatric surgery. While a decrease in EI is the main driver of weight loss, the literature presents a complex and inconsistent picture of the consequences of bariatric surgery on macronutrient intake, food selection and food reward/aversion processes, all of which have been implicated to a greater or lesser extent in the diminution in EI.

From a methodological standpoint there are two plausible explanations for this confusion which were addressed in the proposed protocol. Firstly, there has been a paucity of follow-up studies with sufficiently robust methodology which acknowledge that food intake behaviour is

likely to transition over time as body weight decreases, then stabilizes and perhaps even rebounds following surgery. By studying patients and controls at 4 time points from 1-month pre-surgery to 24-months post-surgery it is more likely that potentially misleading conclusions about the dynamics of food intake behaviour following bariatric surgery based on single time point studies can be avoided.

Secondly, most of the studies have placed overwhelming and unquestioning reliance on the purported validity of self-reported food intake data. Unfortunately, all techniques for dietary assessment in current use are fraught with inherent and extrinsic methodological problems making accurate measurements of food intake data under free-living conditions one of the most intractable problems facing nutrition research [70] [71]. Independent validation of EI data using DLW estimates of TEE [70] [72] has conclusively demonstrated that people with obesity consistently underreport their EI by self-reported measures. By implication, under-reporting of EI also implies an under-reporting of dietary factors which may be food and/or macronutrient specific. However, the question of whether there is distortion in macronutrient reporting in patients after bariatric surgery has not been fully answered and is difficult to prove given that macronutrients are highly interrelated when expressed as %energy.

Consequently, an imperative in bariatric research must be to obtain objective and unbiased measures of appetite and eating behaviour. While the semi-naturalistic laboratory conditions of HISU can reduce participant self-awareness compared to standard laboratories [73] they cannot replicate the free-living situation. However, it can legitimately be argued that this is not their intention. Rather, this fully residential study permitted the isolation and systematic testing of specific variables associated with appetite and eating behaviour free from the constraints of external influences which are an inevitable part of a free-living scenario.

Another novel feature of this study protocol has been the opportunity to test the validity in a bariatric population of several definitions of ED [41] [42] and validate a definition of an EO which has been widely applied in the literature [43] [74] [75]. Other strengths of this protocol included the control over the period of fasting prior to the measurement period at each time point and the steps taken to facilitate ‘normal’, free-living behaviour including tailoring of menus to individual food preferences, the absence of researcher-imposed mealtimes, and researcher absence when food was being eaten. Finally, measurement of different outcomes including RMR, body composition and food preferences were standardised, with the LFPQ administered in the satiated state and after the covert measurements had been completed.

There were some limitations of this protocol which need to be acknowledged. While the aim was to encourage and mimic free-living behaviours, several factors including dietary advice delivered as standard post-operative care, social desirability bias and perceived negative and positive connotations associated with certain foods may have influenced food consumption decisions. Furthermore, the availability of a large variety of foods could have inadvertently heightened food interest and impacted eating behaviour. In addition, while recruitment was initially focused on RYGB patients, an increase in OAGB procedures in the U.K. led to the inclusion of patients having one or other of these procedures. However, there is evidence that the OAGB procedure may lead to greater malabsorption and adverse nutritional events compared to the RYGB procedure [76], which could potentially impact EI and food selection. In the patient cohort, differences in pre-operative care may also have an impact on the outcomes. For example, patients from the UK may be enrolled in weight management services for up-to 24-months prior to surgery, while patients in the ROI are clinically selected and not enrolled in pre-operative weight management programmes.

4. Conclusions

A robust methodology to assess the various components of eating and other associated behaviours is imperative for understanding the causal mechanisms underlying changes in food intake after bariatric surgery. While the proposed study design represented a compromise between the demands of external and internal validity it may help to fill a critical void in understanding the dynamics of food selection and intake behaviour following bariatric surgery which, hitherto, has suffered from overreliance on and uncritical acceptance of the purported integrity of self-reported food intake data. However, while objective laboratory observations of food intake are essential for providing crucial experimental data to complement free-living studies it is essential that laboratory and field research in this area should advance together to fully understand the clinical significance of changes in food selection and intake following gastric bypass surgery.

Understanding the underlying mechanisms and post-operative eating behaviours that contribute to individual variability in the reduction of EI and body weight following surgery could help identify those who are most likely to benefit from gastric bypass surgery and provide more individually targeted approaches to optimise sustained weight regulation. It may also have the potential to inform the development of more targeted approaches for the majority of people with obesity who will manage the condition by non-surgical treatments and allow patients to make more informed decisions regarding their treatment approaches.

Acknowledgements

Authors' Contributions:

The study was initiated by ACS, CWleR and MBEL. The study design was conceived and developed by ACS, CWleR, MBEL and RKP. Collection of data and data analysis were carried out by MBEL, RKP, TR, FN, AB and MM. All authors contributed to the writing of and approved the final manuscript.

Funding:

This research was supported by a US-Ireland Research and Development Partnership grant (R01DK106112) through the Health and Social Care R&D Division of Northern Ireland (STL/5062/14) and the UK Medical Research Council (MC_PC_16017), Health Research Board of the Republic of Ireland (USIRL-2006), and the National Institute of Diabetes and Digestive and Kidney Diseases of the National Institutes of Health (R01DK106112).

The content of the manuscript is solely the responsibility of the authors and does not necessarily represent the official views of the funding agencies.

Conflicts of interest:

The authors declare no competing interests

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Table 1. Macronutrient paradigm for the foods served to study participants.

	High Simple Sugar (%energy)	High Complex Carbohydrate (%energy)	High Protein (%energy)
High Fat (%energy)	n=9 Fat >40% energy Sugar >30% energy e.g. chocolate muffin, caramel chocolate bar, ice cream	n=9 Fat >40% energy CCHO >30% energy e.g. croissant, steak pies, apple pies	n=9 Fat >40% energy Protein >13% energy e.g. salted peanuts, smoked bacon, mature cheddar cheese
Low Fat (%energy)	n=9 Fat <20% energy Sugar >30% energy e.g. banana, grapes, sugar-free meringues	n=9 Fat <20% energy CCHO >30% e.g. sesame bagel, white bread, sugar-free jelly	n=9 Fat <20% energy Protein >13% energy e.g. turkey bacon, crumbed ham, fat- free cottage cheese

Macronutrient mix groups adapted from Geiselman et al., [36]
CCHO Complex carbohydrate

Table 2: Example of a participant menu of foods and beverages (n=84^a) served at each study time point

	HFHSS foods	HFHCCO foods	HFHP foods	LFHSS foods	LFHCCO foods	LFHP foods	Additional foods
Buffet	Granola Fruit and Nut	Croissant	Sausages (fried)	Banana	Porridge	Turkey rashers (fried)	Orange juice Apple juice White bread Brown bread
Fridge/Cupboard	Blueberry Muffins Chocolate cake bars KitKat Snickers Nutella Peanuts and Raisins	Steak & gravy pie Pain au chocolat Crisps Cheese crackers Shortbread Apple pie	Quiche Lorraine Mature cheddar Peanut butter Salami Boiled eggs Salted peanuts	Apple Grapes Pears Salad tomatoes Marshmallows Fruit pastel sweets	Baked Potato Minestrone soup Lentil soup Baked beans White bread Brown bread	Ham Tuna in brine Quorn turkey slices Fat-free cottage cheese Turkey slices 0%fat protein yoghurt	Coca cola Diet coca cola Fanta Fanta Zero Sugar-free cordial Tea Coffee Milk Sugar Sweetener White bread Brown bread Honey Jam Butter Margarine Salad Cream Ketchup Salt Pepper
Menu Options	Pork medallions in a cider jus Sticky toffee pudding	Vegetable loaf with tomato sauce Mini eclairs	Poached chicken coconut curry Almonds	Sweet and sour chicken Raspberry sorbet	Smoked haddock pie Low-fat rice pudding	Braised beef with vegetables and red wine sauce Vanilla soya dessert	Rice Pasta Boiled Potatoes Salad Mixed vegetables Cream

HFHSS; High fat, High Simple Sugar **HFHCCO**; High Fat High Complex Carbohydrate **HFHP**; High Fat, High Protein **LFHSS**; Low Fat, High Simple Sugar **LFHCCO**; Low Fat, High Complex Carbohydrate **LFHP**; Low Fat, High Protein

^a n=84 foods comprised of 54 foods based on the macronutrient mix groups [36] identified using the participant food choice questionnaire administered prior to time point 1 and 30 additional food items (e.g. tea, coffee, sauces) that were available to every participant.

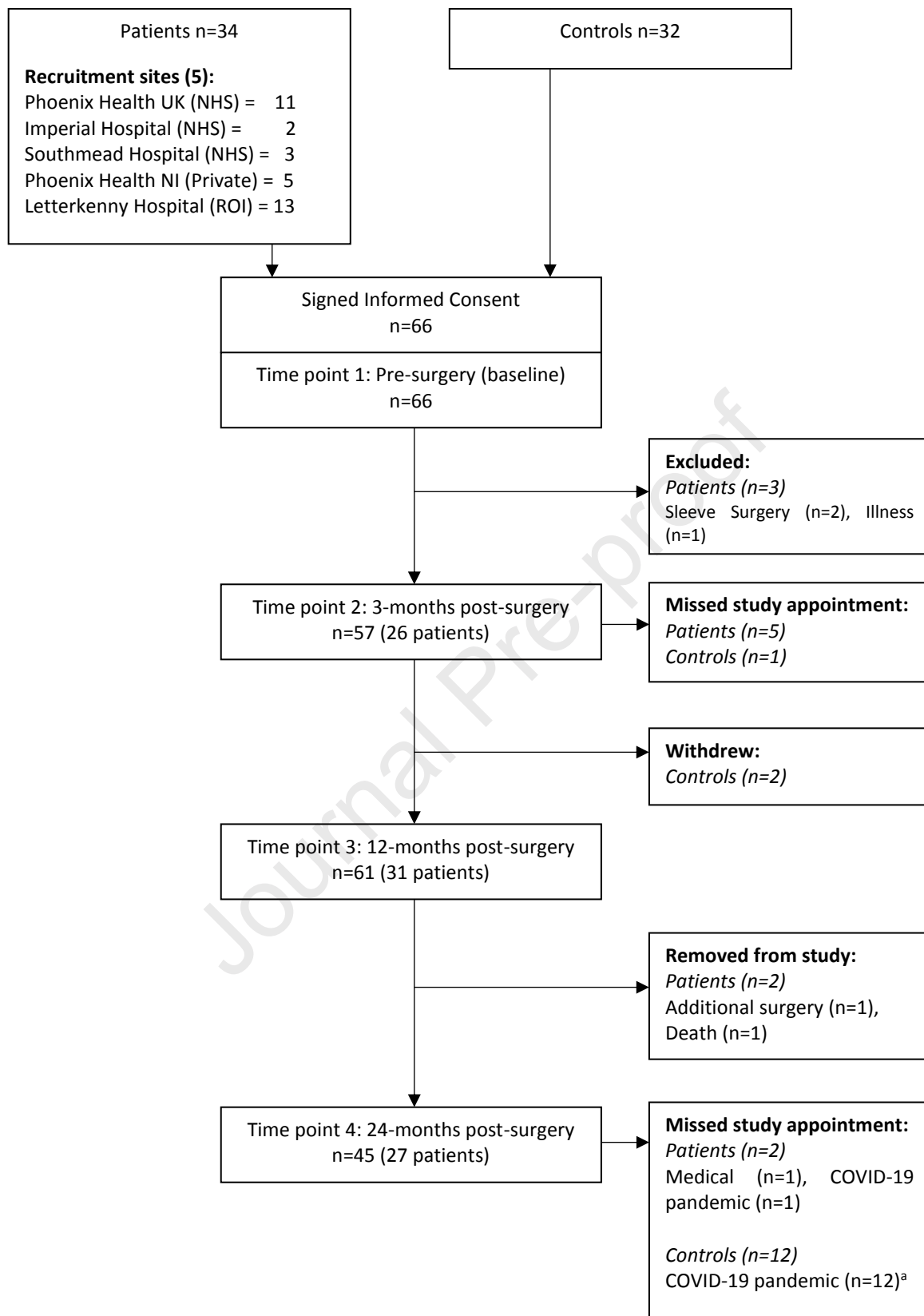


Figure 1: Overview of participant recruitment, progression and retention

NHS National Health Service **ROI** Republic of Ireland. ^a Body weight, gastrointestinal symptoms and medication data collected via telephone.

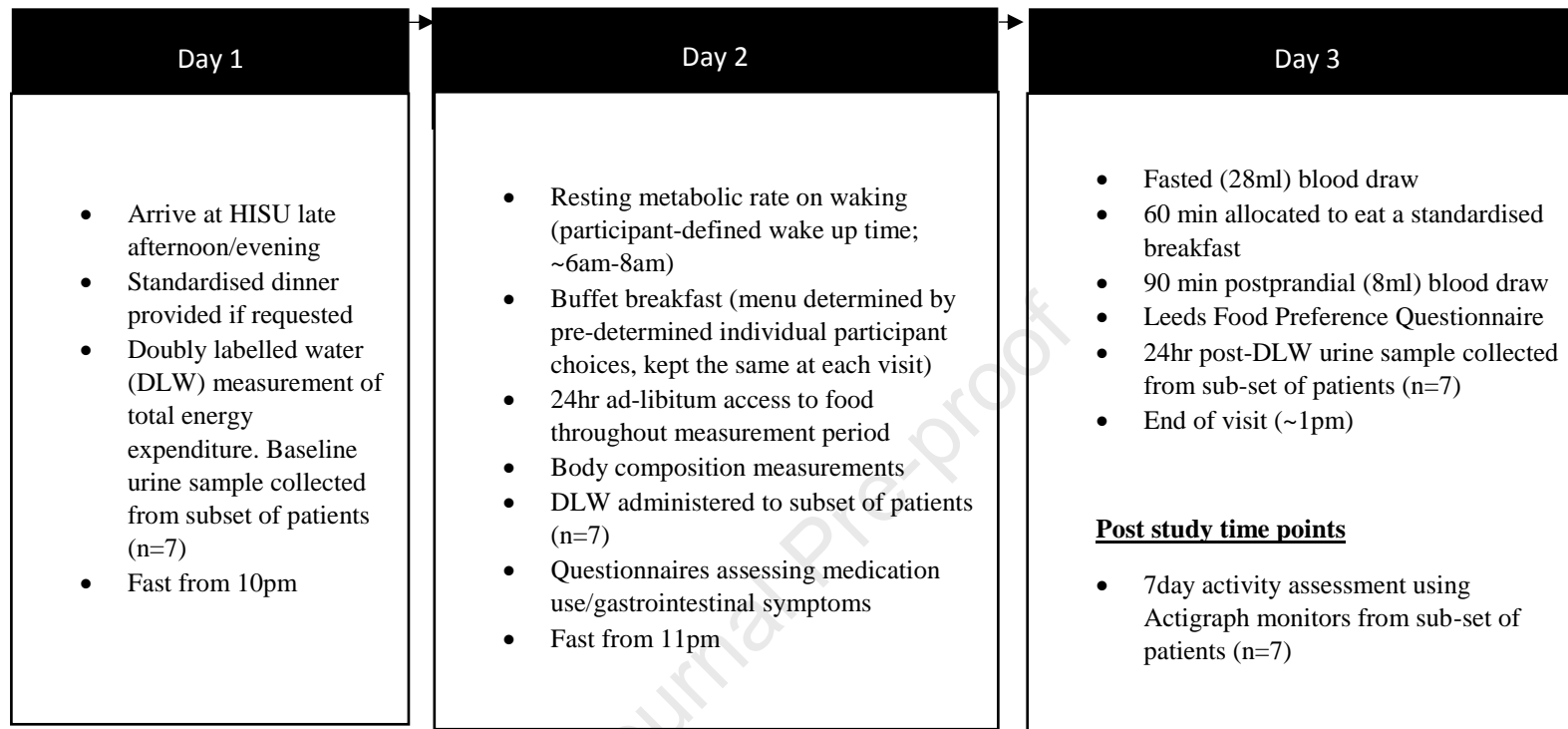


Figure 2. Residential study protocol with scheduled measurements.

AUTHOR DECLARATION

- 1) We wish to confirm that there are no known conflicts of interest associated with this publication and there has been no significant financial support for this work that could have influenced its outcome.
- 2) We confirm that the manuscript has been read and approved by all named authors and that there are no other persons who satisfied the criteria for authorship but are not listed. We further confirm that the order of authors listed in the manuscript has been approved by all of us.
- 3) We confirm that neither the entire paper nor any of its content has been submitted, published, or accepted by another journal. The paper will not be submitted elsewhere if accepted for publication in the Journal.
- 4) We confirm that we have given due consideration to the protection of intellectual property associated with this work and that there are no impediments to publication, including the timing of publication, with respect to intellectual property. In so doing we confirm that we have followed the regulations of our institutions concerning intellectual property.
- 5) We confirm that any aspect of the work covered in this manuscript that has involved either experimental animals or human patients has been conducted with the ethical approval of all relevant bodies and that such approvals are acknowledged within the manuscript.
- 6) We understand that the Corresponding Author is the sole contact for the Editorial process (including Editorial Manager and direct communications with the office). He/she is responsible for communicating with the other authors about progress, submissions of revisions and final approval of proofs.

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16-04-21

Journal Pre-proof