Endogenous plasma glucagon-like peptide-1 following acute dietary fibre consumption

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Abstract

SCFA resulting from the microbial fermentation of carbohydrates have been linked to increased glucagon-like peptide-1 (GLP-1) secretion from the gastrointestinal tract in cell and animal models; however, there is little direct evidence in human subjects to confirm this. The present study was designed to investigate whether endogenous plasma GLP-1 concentrations increase following acute consumption of 48 g dietary fibre (as resistant starch (RS)) from high-amylose maize type 2 RS (HAM-RS2) compared with matched placebo. In a randomised double-blind cross-over study, endogenous plasma GLP-1 concentrations were measured in 30 healthy males following a standardised breakfast and lunch meal. Changes to GLP-1, glucose, insulin and C-peptide were assessed half hourly for 7 h. Following the breakfast meal, plasma GLP-1 concentrations were lower with HAM-RS2 compared with the placebo (P = 0.025). However, there was no significant difference between the supplements following the lunch meal. Plasma insulin concentrations were significantly lower following the lunch meal (P = 0.034) with HAM-RS2 than with the placebo, but were not different after breakfast. Plasma glucose and C-peptide concentrations did not differ at any point. These results suggest that increased dietary fibre intake, in the form of HAM-RS2, does not acutely increase endogenous GLP-1 concentrations in human subjects. Further fibre feeding studies are required to determine whether GLP-1 concentrations may increase following longer-term consumption.

Key words: Resistant starch; Dietary fibre; Glucagon-like peptide-1

Glucagon-like peptide-1 (GLP-1) is secreted from the endocrine L cells in the small and large intestine. GLP-1 7–36 (the most biologically active form) is rapidly broken down by protease enzymes, such as dipeptidyl peptidase-IV, to GLP-1 9–36. GLP-1 has been well characterised as having two main roles as a satiety hormone and as an incretin. As a satiety hormone, GLP-1 has been shown to both enhance satiety and reduce food intake (12) and increased satiety (13,14). The only potential candidate that could fulfil all of these roles would be GLP-1.

GLP-1-secreting L cells in the colon (2). Therefore, the products of the fermentation of fibres may cause an increase in endogenous GLP-1 concentrations and thus may potentiate the effects of GLP-1 as an incretin as well as a satiety signal. Resistant starch (RS), a non-viscous fermentable dietary fibre, has been shown in many animal studies to increase GLP-1 production (3–5). Studies in human subjects where GLP-1 has been measured following increased RS feeding are limited (6–8), with only one of these being an acute feeding study (8), and these have predominately shown no effect of RS on GLP-1 concentrations. However, the beneficial effects that have been observed following increased RS consumption, ranging from 24 h to 12 weeks, have been an improvement in insulin sensitivity (6,7,9,10), insulin secretion (11), a reduction in food intake (12) and increased satiety (13,14). The only potential candidate that could fulfill all of these roles would be GLP-1.

Our group has previously demonstrated a significant reduction in food intake following increased RS intakes (from high-amylose maize type 2 RS (HAM-RS2)) (12). The present study therefore aimed to follow a similar protocol but specifically measuring the endogenous secretion of GLP-1 and postprandial insulin and glucose concentrations.

Abbreviations: GLP-1, glucagon-like peptide-1; HAM-RS2, high-amylose maize type 2 resistant starch; iAUC, incremental AUC; RS, resistant starch.

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The study aimed to determine whether the previously observed reductions in food intake and changes to postprandial insulin concentrations could be due to increases in endogenous GLP-1 concentrations.

**Experimental methods**

**Participants**

A total of thirty healthy adult males, with a mean age of 24·5 (SEM 0·7) years and a mean BMI of 26·2 (SEM 0·9) kg/m², participated in the present study. Participants were excluded if they had a history of gastrointestinal or endocrine disease. The present study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human subjects were approved by the University of Surrey Ethics Committee. Written informed consent was obtained from all participants.

**Study design**

The study design was based on that described previously[12]. The study was a single-blind, cross-over study in which participants visited the clinical unit on two occasions at least 1 week apart. On these clinical visits, participants consumed either 80 g Hi-Maize (comprising 60% type 2 RS (HAM-RS2) and 40% rapidly digestible starch (RDS), therefore providing 48 g HAM-RS2 as measured by the Association of Official Analytical Chemists for total dietary fibre method 991.43) or 32 g of placebo (Amioca, comprising 100% RDS) supplying no dietary fibre. Both starches were manufactured and supplied by Ingredion Inc. Half of the starch was provided at breakfast and the other half at lunch, incorporated into a mousse, therefore providing 24 g dietary fibre or 0 g dietary fibre at each meal.

On arrival at the clinical unit, a cannula was placed into the antecubital fossa and two fasted blood samples were taken. At time zero, participants consumed the standardised breakfast (1587 kJ, 9·6 g protein, 67·1 g carbohydrate, 8·0 g fat and 24·5 g dietary fibre (24·5 g dietary fibre for the HAM-RS2 supplement)). Following the breakfast, blood samples were taken every 30 min for 3 h. Participants were then provided with the standardised lunch (3888 kJ, 36·1 g protein, 112·1 g carbohydrate, 37·0 g fat and 5·0 g dietary fibre (29·0 g dietary fibre for the HAM-RS2 supplement)). Further blood samples were collected for the following 4 h. Bowel habit and symptom diaries were completed on the day of the study and the following day to assess gastrointestinal tolerance of the starches.

**Biochemistry**

Whole blood was collected into fluoroxyalate tubes for glucose analysis, potassium EDTA tubes for insulin analysis and potassium EDTA tubes with 200 kallikrein-inhibiting units aprotinin per ml of blood for C-peptide and total GLP-1 concentrations. All samples were centrifuged for 10 min at 3000 rpm and plasma stored at −20°C until batch analysis. Plasma glucose concentrations were measured enzymatically using a commercially available kit for the ILab650 (Instrumentation Laboratory) and inter-assay variation was <2%. Plasma insulin and C-peptide concentrations were measured by RIA using commercially available kits (Millipore) with inter-assay variations of 18·2 and 17·7%, respectively, and intra-assay variations of 9·2 and 9·5%, respectively. Plasma total GLP-1 concentrations were measured by ELISA using a commercially available kit (Millipore) with an inter-assay variation of 15·0% and an intra-assay variation of 4·5%.

**A priori power calculation and statistics**

The thirty participants in the present cross-over study (α = 0·05) would yield an 80% probability of detecting a 540 pmol/l per 420 min change in postprandial GLP-1 response (AUC, based on a treatment sd of 1018). All post-prandial time course data were analysed using repeated-measures ANOVA. The incremental AUC (iAUC) was calculated for each meal using the trapezoid method, and then the iAUC was compared between the treatments using
paired t tests. All statistical analyses were carried out using SPSS version 18 for Windows (SPSS, Inc.) with significance assumed as \( P < 0.05 \). All results presented are means with their standard errors.

Results

Both HAM-RS2 and placebo were well tolerated by the participants with a similar frequency of bowel movements recorded and no adverse gastrointestinal effects reported on the day of the study or the following day. Participants’ weight did not differ between the two study days.

Glucagon-like peptide-1 concentrations

There was a significant treatment \( \times \) time interaction between HAM-RS2 and placebo for postprandial GLP-1 concentrations \( (P = 0.020) \), Fig. 1).

The iAUC for each meal showed a significantly lower GLP-1 response with HAM-RS2 compared with placebo following breakfast \( (P = 0.025); \) Fig. 2(a)). However, following the lunch meal, there were no differences between the starches (Fig. 2(a)).

Postprandial metabolite concentrations

Over the whole 7 h postprandial period, there were no differences between HAM-RS2 and placebo for concentrations of plasma glucose, insulin or C-peptide.

The iAUC for insulin following breakfast (0–180 min) were not different between HAM-RS2 and placebo; however, after lunch (180–300 min), there was a significantly lower response following HAM-RS2 than with placebo \( (P = 0.034) \); Fig. 2(c)).

There were no significant differences in iAUC between the RS and placebo for glucose or C-peptide (Fig. 2(b) and (d), respectively).

Discussion

To our knowledge, this is the first study that has investigated the effects of increased HAM-RS2 intake on endogenous GLP-1 concentrations when compared with a placebo matched for both energy and glycaemic load, in human subjects. The present study has shown that during the 3 h following a breakfast meal containing HAM-RS2, there was a significant reduction in GLP-1 concentrations compared with placebo. However, after a further HAM-RS2-containing meal, there was no difference in GLP-1 concentrations between the supplements.
In the present acute postprandial study, increased intakes of 48 g HAM-RS2, a non-viscous fermentable fibre, did not increase plasma GLP-1 concentrations and, indeed, following the breakfast meal, GLP-1 concentrations were significantly lower with the RS than compared with the placebo. There is little previous direct evidence for an effect of RS on GLP-1 in human subjects. A postprandial study comparing 27 g RS (from raw potato starch) with a placebo (that was not matched for glycaemic carbohydrate) also found lower GLP-1 with the RS(18). In the present study, there was a significant treatment × time interaction, translating as the RS-containing meal giving a lower acute GLP-1 response with higher post-lunch levels, when compared with placebo (Fig. 1). This may suggest that the increase in GLP-1 concentrations was due to the fermentation of HAM-RS2, and investigating the effects later on in the day or even the subsequent morning may have produced significant differences in GLP-1 concentrations. However, one study in which 60 g HAM-RS2 was consumed for 24 h before the investigation compared with the same placebo found no effect of HAM-RS2 on GLP-1 concentrations(19). Another study looking at carry-over effects of evening meals varying in indigestible carbohydrate content found no effect of evening meals on fasting GLP-1 concentrations the following day, but did demonstrate a significantly higher GLP-1 AUC (0–120 min) following one treatment (barley kernels containing 20·2 g of total dietary fibre) compared with the control product (3·9 g of total dietary fibre)(19). Additionally, a study in which 30 g HAM-RS2 was consumed daily for 4 weeks compared with a matched glycaemic carbohydrate placebo again found no effect of HAM-RS2 on GLP-1(17), although, unlike the present study, this was not an acute response to the fibre. Overall, the data from the present study and from the previous studies would suggest that HAM-RS2 does not cause an increase in endogenous GLP-1, particularly when HAM-RS2 is consumed acutely.

While the GLP-1 data in human subjects following increased RS intake are limited, there is much animal work that has shown consistent increases in GLP-1 blood concentration following higher levels of RS feeding as well as increases in proglucagon gene expression in the caecum and colon. However, it is difficult to directly translate these findings as the studies in rodents have not been acute (days rather than hours), and therefore it may be that prolonged exposure to HAM-RS2 in human subjects is required to affect GLP-1 concentrations. A previous study in human subjects has shown an increase in GLP-1 concentrations following wheat fibre intake, although a year of increased wheat fibre feeding was required before the effect was observed.

Other acute studies using increased fibre feeding have shown inconsistent results. A 2 h postprandial study feeding a galactose and guar gum-mixed drink compared with a control drink found increased GLP-1 concentrations another 6 h postprandial study comparing an insulin drink with high-fructose maize syrup drinks found GLP-1 concentrations to be higher with insulin at 30 min. However, other studies where dietary fibres have been given acutely, as in the present study, have found no effect on GLP-1 concentrations.

Overall, a recent review has concluded that the current research on the effects of fibres on GLP-1 demonstrates that fibres do not increase GLP-1 concentrations compared with control; however, the authors have acknowledged that there are few studies that have actually assessed the effects of fibre on gut hormones directly.

Previously, we have shown that acute ingestion of HAM-RS2 reduced postprandial insulin concentrations, which we have demonstrated again in the present study following the second dose of HAM-RS2, but not following the first dose, or over the whole 7 h period. In the present study, the lower insulin response observed after the second dose was found in conjunction with a higher GLP-1 concentration, which is not in agreement with GLP-1 acting as an incretin following fibre feeding. This would therefore suggest that following increased RS intakes, GLP-1 is unlikely to have a role as an incretin. In many experimental designs, RS has replaced portions of digestible starch, or ratios of amylose: amylopectin have been manipulated (changing glycaemic index and/or glycaemic load), thus making comparisons problematic. A recent review of the literature related to RS and glycaemia/insulinaemia highlighted this disparity. In our previous work, we hypothesised that a reduced postprandial insulin response often attributed to RS intake may be due to increased hepatic insulin clearance; however, we found no evidence to support this in the present study.

The limitations of the present study are that we only measured total as opposed to active GLP-1 concentrations and therefore the effects of GLP-1 as a satiety signal may be masked. As recently reviewed, gut hormones may not act independently, and it may be the additive effects of several gut hormones that result in observed changes to satiety. Indeed, it is established that the postprandial levels of GLP-1 attained during feeding would have minimal effects on food intake, when considered in isolation. However, GLP-1 may still have a role as an incretin, as some studies have suggested that it may reduce glycaemia to a small degree independently to any effects on insulin secretion. Therefore, the relevance of measuring total GLP-1 concentrations needs to be considered in an attempt to understand the health benefits of dietary fibre intake.

Overall, it would appear from the present study and that of the literature that GLP-1 is not a candidate signal for the acutely reduced food intake and enhanced insulin sensitivity previously attributed to HAM-RS2.

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and analysis of the data. L. S. was involved in the recruitment of the volunteers, and the conducting of the study days and analysis of the data. M. D. R. was involved in the designing of the study protocol, assistance on the study days, analysis of the data, statistical analysis and refining of the paper.

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