Clinical Study

Consumption of Cross-Linked Resistant Starch (RS4_{XL}) on Glucose and Insulin Responses in Humans

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Objective. The objective was to compare the postprandial glycemic and insulinemic responses to nutrition bars containing either cross-linked RS type 4 (RS4_{XL}) or standard wheat starch in normoglycemic adults ($n=13$; age $=27\pm5$ years; BMI $=25\pm3$ kg/m$^2$).

Methods. Volunteers completed three trials during which they consumed a glucose beverage (GLU), a puffed wheat control bar (PWB), and a bar containing cross-linked RS4 (RS4_{XL}) matched for available carbohydrate content. Serial blood samples were collected over two hours and glucose and insulin concentrations were determined and the incremental area under the curve (iAUC) was calculated.

Results. The RS4_{XL} peak glucose and insulin concentrations were lower than the GLU and PWB ($P<.05$). The iAUC for glucose and insulin were lower following ingestion of RS4 compared with the GLU and PWB trials.

Conclusions. These data illustrate, for the first time, that directly substituting standard starch with RS4_{XL}, while matched for available carbohydrates, attenuated postprandial glucose and insulin levels in humans. It remains to be determined whether this response was due to the dietary fiber and/or resistant starch aspects of the RS4_{XL} bar.

1. Introduction

Consumption of whole grains has been recommended to improve insulin sensitivity and lower serum glucose and insulin concentrations. Whole grain consumption of three servings or more per day was among changes that were included in the 2005 dietary guidelines to reduce the risk of acquiring chronic diseases [1]. Whole grains are major sources of dietary fiber (DF), yet typical DF consumption patterns do not meet the recommended 25–35 g per day. Therefore, eating more grain-based fiber-rich foods is warranted to help optimize health and potentially manage some chronic metabolic conditions.

At present, resistant starches (RSs) have drawn broad interest for their health benefits and functional properties [2, 3]. Initial clinical studies demonstrated that RS has properties similar to soluble fiber, shows promising physiological benefits in humans, and may prevent disease. Several potential physiological benefits ascribed to RS include attenuation of blood glucose and insulin levels in both healthy and diabetic individuals, positive effects on large bowel health and prevention of colonic cancer, increased absorption of minerals, serving as a prebiotic, and increased fat oxidation [3–9]. There are four basic “types” of RS. Type 1 (RS1) is composed of starch granules embedded in indigestible plant material. Type 2 (RS2) is native granular starch with a B-type x-ray pattern, such as found in potato and high-amylose maize. Type 3 (RS3) is crystallized starch and maltodextrins made by alternate cooking/cooling processes on starchy materials. Type 4 (RS4) is chemically modified starch typically through esterification, crosslinking (RS4_{XL}), or transglycosylation.

The majority of human clinical trials have been conducted using only RS2 or RS3, which tend to illustrate decreased blood glucose following consumption of foods with these starches added [3, 6, 10–15]. It is difficult to fully understand the beneficial capacity of RS due to the methods used in the human clinical trials that tested the efficacy of RS. For example, one clinical trial failed to control both the amount and source of all the ingredients [16].
In a study investigating the effects of esterified RS type 4 (RS4OSA), Heacock et al. [17] reported that RS4OSA decreased peak glucose and insulin levels when it was administered in water, but RS is typically consumed in foods. Also, in studies by Behall et al. [18, 19], the amount of available carbohydrate differed, which limits the capacity to determine if the attenuation of glucose and insulin was due to the RS or the fact there was less available carbohydrate. Furthermore, Robertson et al. [6] provided packets containing RS2 for volunteers to sprinkle on their food thereby not illustrating the effects that might be achieved when provided in the food supply. Taken together, the available data illustrate that RS has the potential to lower blood glucose. However, few clinical trials testing the effects of RS have controlled ingredients and the amount of available carbohydrates to better delineate the role of RS in affecting the insulin and glucose responses, and no published clinical trials investigating the glucose lowering potential of RS4XL exist.

Therefore, the aim of this study was to investigate the acute effects of consuming RS4XL incorporated into a nutrition bar, while controlling for nonstarch ingredients and available carbohydrates, on postprandial glucose and insulin responses in young adults with a randomized clinical trial (NCT00687960, clinicaltrials.gov).

2. Materials and Methods

2.1. Subjects. The Institutional Review Board of Kansas State University approved the study, and written informed consent was obtained from all volunteers prior of the study. Inclusion criteria were no diagnosis of acute or chronic metabolic diseases, free of gastrointestinal disorders, body mass index of 23–30 kg/m², and nonsmokers. Volunteers were screened for glucose tolerance using a two hour 75 g glucose tolerance test prior to enrollment to ensure each volunteer had normal glucose tolerance. Based on BMI values, these volunteers all have similar risks for glucose tolerance using a two hour 75 g glucose tolerance test was performed prior to enrollment to ensure all volunteers were either contraceptive pills or progesterone injections. Regardless, the females were scheduled to perform all the trials during the follicular phase of their menstrual cycle and each served as their own control.

2.2. Oral Glucose Tolerance Test. Prior to enrollment, volunteers arrived after a 12 hours overnight fast. Blood samples were drawn by finger stick at baseline and 120 minutes after ingesting 75 g of glucose in solution (296 mL; Sun-Dex 75 g, Fisher Scientific, Houston, Tex, USA). Samples were analyzed in duplicate for glucose concentration (YSI 2300 STAT, Yellow Springs, Ohio, USA). The oral glucose tolerance test was used to confirm the absence of prediabetes or diabetes.

2.3. Study Design. All trials were performed at the Human Metabolism Laboratory at Kansas State University. Volunteers completed three trials via a controlled randomized crossover design. During each trial, volunteers consumed one of the following: dextrose solution (198 mL of a standard 75 g oral glucose tolerance beverage; GLU), a control bar containing pumped wheat (65 g; PWB), and a bar containing cross-linked RS4XL bar (80 g; RS4XL) (Table 1). All treatments were designed to provide 50 g of available carbohydrate (Table 2). All trials were completed by each volunteer with at least a seven-day washout between testing days. This was a quasiblinded experiment in that one treatment was a beverage and the other two were in bar form, which were randomly administered using a Latin Square design. Some female volunteers did not use oral contraceptives, but others used either contraceptive pills or progesterone injections. Regardless, the females were scheduled to perform all the trials during the follicular phase of their menstrual cycle and each served as their own control.

2.4. Experimental Bars. The only altered ingredients between bars were either pumped wheat or RS4XL (Table 1). Briefly, the nutrition bars were prepared by adding pumped wheat or cross-linked RS4 to wheat germ. The dose of the RS4XL (27.2 g) was intended to be close to the dose used previously [6], whereby a significant improvement in insulin sensitivity was observed. The remaining ingredients (water, corn syrup, brown sugar, gum acacia, and Panodan 150 K) were heated to 85°C over 4 minutes, poured over the dry ingredients, and then manually mixed quickly until dry ingredients were evenly distributed throughout the mixture. The mixture was scooped into a metal pan, pressed evenly throughout the pan, and allowed to cool for 20 minutes before cutting into bars. Crude nutrient analysis was determined by proximate analysis, while total dietary fiber was assessed independently (Medallion Labs, Minneapolis, Minn) (Table 2). Available carbohydrate was calculated as the difference between total carbohydrate and dietary fiber as used previously [9]. Controlling for available carbohydrate in this fashion has been shown to affect the glycemic response, while controlling for RS content does not necessarily affect the glycemic response [21].

<table>
<thead>
<tr>
<th>Table 1: Ingredients and their concentrations by relative weight (% total) in the test bars.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Puffed Wheata</td>
</tr>
<tr>
<td>Resistant Starch type 4b</td>
</tr>
<tr>
<td>Corn Syrupc</td>
</tr>
<tr>
<td>Wheat Germd</td>
</tr>
<tr>
<td>Brown Sugare</td>
</tr>
<tr>
<td>Waterf</td>
</tr>
<tr>
<td>Gum Acaciag</td>
</tr>
<tr>
<td>Panodan 150Kh</td>
</tr>
<tr>
<td>--------------------------------------------------</td>
</tr>
<tr>
<td>a Quaker Oats</td>
</tr>
<tr>
<td>b Fibersym RW; MGP Ingredients, Inc.</td>
</tr>
<tr>
<td>c Karo light corn syrup</td>
</tr>
<tr>
<td>d Kretschmer Original Toasted</td>
</tr>
<tr>
<td>e CS H Pure Cane Sugar; golden brown</td>
</tr>
<tr>
<td>f Tap water (Manhattan, Kan)</td>
</tr>
<tr>
<td>g TIC Gums</td>
</tr>
<tr>
<td>h Danisco</td>
</tr>
</tbody>
</table>

* Samples were analyzed in duplicate for glucose concentration (YSI 2300 STAT, Yellow Springs, Ohio, USA). The oral glucose tolerance test was used to confirm the absence of prediabetes or diabetes.

* The only altered ingredients between bars were either pumped wheat or RS4XL (Table 1). Briefly, the nutrition bars were prepared by adding pumped wheat or cross-linked RS4 to wheat germ. The dose of the RS4XL (27.2 g) was intended to be close to the dose used previously [6], whereby a significant improvement in insulin sensitivity was observed. The remaining ingredients (water, corn syrup, brown sugar, gum acacia, and Panodan 150 K) were heated to 85°C over 4 minutes, poured over the dry ingredients, and then manually mixed quickly until dry ingredients were evenly distributed throughout the mixture. The mixture was scooped into a metal pan, pressed evenly throughout the pan, and allowed to cool for 20 minutes before cutting into bars. Crude nutrient analysis was determined by proximate analysis, while total dietary fiber was assessed independently (Medallion Labs, Minneapolis, Minn) (Table 2). Available carbohydrate was calculated as the difference between total carbohydrate and dietary fiber as used previously [9]. Controlling for available carbohydrate in this fashion has been shown to affect the glycemic response, while controlling for RS content does not necessarily affect the glycemic response [21].
Table 2: Nutrient composition of each treatment per dose (GLU = 198 mL; PWB = 65 g; RS4 = 80 g).

<table>
<thead>
<tr>
<th></th>
<th>GLUa</th>
<th>PWBb</th>
<th>RS4XLc</th>
<th>Δd</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total Energy (kcal)</td>
<td>200</td>
<td>261</td>
<td>326</td>
<td>(65 kcal, 125%)</td>
</tr>
<tr>
<td>Carbohydrate (g)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>50</td>
<td>56</td>
<td>71</td>
<td>(15 g, 127%)</td>
</tr>
<tr>
<td>Availablec</td>
<td>50</td>
<td>51</td>
<td>51</td>
<td>(0 g, 0%)</td>
</tr>
<tr>
<td>Total Dietary Fiber (g)d</td>
<td>—</td>
<td>5</td>
<td>20</td>
<td>(15 g, 400%)</td>
</tr>
<tr>
<td>Fat (g)</td>
<td>—</td>
<td>1</td>
<td>2</td>
<td>(1 g, 200%)</td>
</tr>
<tr>
<td>Protein (g)</td>
<td>—</td>
<td>7</td>
<td>6</td>
<td>(1 g, 86%)</td>
</tr>
</tbody>
</table>

a glucose tolerance test beverage (Sun-Dex, Fisher Scientific, Houston, Tex)
b Crude nutrient composition was determined by proximate analysis (total energy, total fat, total protein, total carbohydrate).
c derived by subtracting total dietary fiber from total carbohydrate.
d dietary fiber analysis was performed by Medallion Laboratory (Minneapolis, Minn).
e difference (subtraction value, % value) between bars.

2.5. Food Tolerance Test. The postprandial test (two-hours with seven blood samples) was modified from Flammang et al. [22]. During each test, volunteers arrived to the laboratory after a 10–12 hour overnight fast. An indwelling catheter (Terumo, 22gx1, Terumo Medical Corporation, Elekton, Md) was inserted into a forearm vein. The line was kept patent with 0.9% isotonic saline solution (0.9% sodium chloride USP, B. Braun Medical Inc., Irvine, Calif). Ten minutes after inserting the IV catheter, the fasting blood sample was collected. Thereafter, the volunteers consumed the assigned food item for that day. The solution (GLU) or bar was consumed within 10 minutes. The treatment bars were served with 198 mL of water (to match the fluid that was consumed during the GLU trial). Relative to taking the first bite of food, blood samples were collected at −10, 10, 20, 30, 60, 90, and 120 minutes.

2.6. Blood Samples. After collection, blood samples were centrifuged at 2,500 rpm for 15 minutes at 4°C. The plasma was extracted from the vacutainer and immediately analyzed in duplicate for glucose (YSI 2300, Yellow Springs Instruments, Yellow Springs, Ohio) with the remainder stored at −80°C. The frozen samples were analyzed in duplicate for insulin using an endocrine assay (LINCOplex kit, St. Charles, Mo), and measured by LumineX100 (Austin, Tex) instrumentation. Once the samples were analyzed, the highest value attained over the 120 minutes was determined to be the peak value, with the difference between baseline and peak determined to measure the change from fasting to peak.

Glucose and insulin areas under the curve (iAUC) were determined using the trapezoid method (GraphPad Prism v 5.02, La Jolla, Calif). This approach was adapted from a previous RS feeding study [18]. Fasting insulin and glucose values were also used to calculate the homeostasis model assessment (HOMA) [23].

2.7. Diet and Physical Activity Records. A diet record was completed by the volunteers prior to the first test of the study. Volunteers were instructed on how to record their intakes. They were requested to then eat the same foods from that diet recorded the day before the second and third trials. This procedure was used previously [24]. Also, volunteers were requested to record their physical activity for the day before testing and perform that same activity (or inactivity) the day prior to the subsequent tests.

3. Data Analysis

Sample size estimation was calculated using PASS software (NCSS 2007 and PASS 2005, Kaysville, Utah). Based on results from previously published data using RS2 [6], six volunteers were determined necessary (power > 0.80, and P < .05) to detect significant differences in glucose and insulin responses. A repeated measures analysis of variance (SPSS version 11.5, Chicago, Ill) was used to determine significant main effects with significance set at P = .05. Paired t-tests (SPSS version 11.5, Chicago, Ill) were used to determine differences between and among trials for peak, change from baseline to peak and iAUC values for glucose and insulin. The comparisons of interest were primarily between the two bars, with the glucose treatment providing a standard point of reference.

4. Results

The three meals provided practically the same amount (50-51 g) of available carbohydrate (Table 2). Both nutrition bars contained similar amounts of protein and were low in fat, but the RS4XL bar contained four times the dietary fiber (20 g versus 5 g) compared to the PWB. Approximately one-fifth of the total dietary fiber in the RS4XL bar was from gum arabic (∼4.4 g) with the remainder (∼15.6 g) being from RS4XL. The meals differed in their calculated food-energy contents (200–326 kcal, Table 2) because their compositions differed and because their weights to deliver 50 g of available carbohydrates differed.

The commercially available RS4 XL used in this study contains 0.4% phosphors, 10.6% moisture, 91.9% total dietary fiber by AOAC-International Method 991.43, and 83.3% RS by a modified Englyst method [25]. At the time of preparing the 80 g bar of RS4XL, the amount of RS added was ∼20 g (dry solids) according to the formula in Table 1 and the composition of Fibersym RW, while the amount of gum arabic added was 4.4 g (dry solids). At the time of analyzing the finished 80 g bar, the bar contained 20 g of total dietary fiber (Table 2). Assuming no loss of gum arabic in the preparation of the RS4XL bar, the 4.4 g loss of dietary fiber could be attributed to some damage to the RS, which increased its digestibility. Assuming the ratio of RS to dietary fiber remains constant at 0.9 in partially damaged RS4XL, then the 80 g bar of RS4XL fed to the subjects contained ∼14 g RS, implying ∼70% was maintained following preparation of the bar.

The RS4 bar elicited decreased glucose and insulin concentrations at several time points (Figure 1) during the 120 minutes postprandial period as compared with GLU and PWB. Also, consumption of the RS4 bar led to an attenuation
of peak glucose and insulin, and significant differences from baseline to peak and iAUC values glucose and insulin when compared with GLU and PWB (Table 3). The PWB bar attenuated the peak glucose and insulin responses and the iAUC for glucose and insulin compared with GLU. The percent increase from baseline to peak was not different (P = .068) from GLU for glucose, while it was for insulin.

5. Discussion

These data, for the first time, indicate that eating RS4XL from wheat in place of standard wheat starch significantly decreased postprandial insulin and glucose responses. These results are in line with others investigating the insulin and/or glucose lowering effects when RS (typically RS2) is added to foods or incorporated in the diet [6, 16, 18, 26, 27], while a few reported no effect of RS2 or RS3 on glycemia [7, 9]. Results from several other clinical trials reported RS decreased the glycemic response, but those studies had volunteers sprinkle RS onto the food instead of it being an ingredient in the food, mixed only with water, ate large (up to 388 g) portion sizes, failed to control for available carbohydrate, and/or the food eaten contained di...
Table 3: Values for the incremental areas under the curves of glucose and insulin concentrations during each trial. Mean ± SE; different letters within a row indicates significant difference (P < .05).

<table>
<thead>
<tr>
<th></th>
<th>GLUC</th>
<th>PWB</th>
<th>RS4XL</th>
</tr>
</thead>
<tbody>
<tr>
<td>iAUC (mmol/L • 2 hr)</td>
<td>140 ± 31A</td>
<td>84 ± 17B</td>
<td>28 ± 11C</td>
</tr>
<tr>
<td>Peak (mmol/L)</td>
<td>7.30 ± 0.5A</td>
<td>6.33 ± 0.3B</td>
<td>5.40 ± 0.2C</td>
</tr>
<tr>
<td>Increase (%)</td>
<td>60.5 ± 10A</td>
<td>42.7 ± 6A</td>
<td>20.4 ± 3B</td>
</tr>
<tr>
<td>iAUC (pM • 2 hr)</td>
<td>17,575 ± 2,236A</td>
<td>8,758 ± 1,132B</td>
<td>3,659 ± 974C</td>
</tr>
<tr>
<td>Peak (pM)</td>
<td>344 ± 36.7A</td>
<td>211.5 ± 20.1B</td>
<td>162.3 ± 22.6C</td>
</tr>
<tr>
<td>Increase (%)</td>
<td>335 ± 53.2A</td>
<td>243.0 ± 49.3B</td>
<td>126.3 ± 45.8C</td>
</tr>
</tbody>
</table>

contained within the RS4XL bar did not take into account the energy provided via fermentation of the RS4XL into short chain fatty acids. Lastly, we cannot determine the mechanism for the effect observed. It is reasonable to conclude that the RS4XL caused the effects, as it was the only ingredient difference, but it is not possible to determine how much of the change was attributable to the dietary fiber or the RS that is contained within RS4XL.

In conclusion, this is the first published randomized clinical trial to investigate the glucose and insulin lowering potential of RS4XL. Additionally, this is one of a few clinical studies where the treatments were matched for available carbohydrate and the RS was substituted directly for standard carbohydrate and the RS was substituted directly for standard studies where the treatments were matched for available chain fatty acids. Lastly, we cannot determine the mechanism of RS4XL. Additionally, this is one of a few clinical trial to investigate the glucose and insulin lowering potential of RS4XL.

**Abbreviations**

- DF: Dietary fiber
- GLU: Glucose control solution
- iAUC: Incremental area under the curve
- HOMA: Homeostasis Model Assessment of insulin sensitivity
- PWB: Control bar made with puffed wheat
- RS: Resistant starch
- RS4XL: Treatment bar made with cross-linked resistant starch type 4.

**References**


