Impact of overweight and glucose tolerance on postprandial responses to high- and low-glycaemic index meals

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Abstract
The beneficial effects of a low-glycaemic index (GI) meal on postprandial glucose and insulin levels have been demonstrated. However, limited data are available on the impact of overweight and glucose tolerance on postprandial responses to different GI meals. Our aim was to study the effects of physiological characteristics on postprandial glucose, insulin and lipid responses and the relative glycaemic response (RGR) of a low-GI (LGI) and a high-GI (HGI) meal. We recruited twenty-four normal-weight and twenty-four overweight subjects, twelve with normal glucose tolerance (NGT) and twelve with impaired glucose tolerance (IGT) in each group. Both test meals were consumed once and the glucose reference twice. Blood glucose and insulin were measured in the fasting state and over a 2 h period after each study meal, and TAG and NEFA were measured in the fasting state and over a 5 h period. The glucose responses of subjects with IGT differed significantly from those of subjects with NGT. The highest insulin responses to both meals were observed in overweight subjects with IGT. Physiological characteristics did not influence TAG or NEFA responses or the RGR of the meals. The LGI meal resulted in lower glucose (P<0.001) and insulin (P<0.001) responses, but higher TAG responses (P<0.001), compared with the HGI meal. The GI of the meals did not affect the NEFA responses. In conclusion, the LGI meal causes lower glucose and insulin responses, but higher TAG responses, than the HGI meal. The RGR of the meals does not differ between normal-weight and overweight subjects with NGT or IGT.

Key words: Glycaemic index: Overweight: Glucose tolerance: Postprandial response

The concept of glycaemic index (GI) was originally introduced by Jenkins et al.(1). It is a classification of the blood glucose-raising potential of carbohydrate foods. GI is defined as the incremental area under the blood glucose curve of a test food, expressed as the percentage of the response to a reference food consumed by the same subject on a different day(2). When glycaemic responses to meals are investigated, the incremental area under the blood glucose curve of a test meal is divided by the response to a reference food, which is called the relative glycaemic response (RGR) of the test meal. Since GI was first introduced, many studies have investigated the potential health benefits of a low-GI (LGI) diet, as reviewed by Livesey et al.(3). A high-GI (HGI) meal, digested and absorbed rapidly, initially results in a high glycaemic response and increased need for insulin secretion, followed by a relative hypoglycaemia, increased counter-regulatory hormone secretion and increased serum NEFA concentrations(4). The beneficial effects of a LGI meal on postprandial hyperglycaemia and insulin levels have been demonstrated in many studies(5–9), but the data on the effects of meals with different GI values on TAG and NEFA responses are limited. A few studies(9,10) have shown an increased level of lipid response after a HGI meal, while others have failed to show any lipid differences(11,12). To our knowledge, there are only a few studies(13,14) that have assessed the acute effect of physiological characteristics, overweight and glucose tolerance on
postprandial glucose, insulin and lipid responses to LGI and HGI meals.

While it is generally recommended that GI test subjects should be healthy\(^1\)\(^{15}\), many studies have shown no effect of BMI\(^1\)\(^{16–18}\), glucose tolerance\(^1\)\(^{19}\), sex\(^1\)\(^{16,18}\), age\(^1\)\(^{16,18}\) and ethnicity\(^1\)\(^{16,18,20}\) on the measured GI values. Interestingly, there is little knowledge about the effect of body weight and glucose tolerance on the RGR of the meals.

The primary aim of the present study was to examine the effects of overweight and glucose tolerance on the glucose, insulin and lipid responses to a HGI and LGI meal. Furthermore, the second aim was to study the effect of BMI and glucose tolerance on the RGR of the meals.

**Experimental methods**

**Subjects**

We studied twenty-four normal-weight (BMI 20–24.9 kg/m\(^2\)) and twenty-four overweight (BMI 27.5–34.9 kg/m\(^2\)) subjects aged 62–72 years. Both groups included twelve subjects with normal glucose tolerance (NGT) and twelve with impaired glucose tolerance (IGT) based on a 75 g, 2 h oral glucose tolerance test, where the 2 h glucose values were impaired glucose tolerance (IGT) based on a 75 g, 2 h oral glucose tolerance test, where the 2 h glucose values were <7.8 and 7.8–11.0 mmol/l, respectively. Exclusion criteria included smoking, milk allergy, regular medication that would have an effect on glucose or lipid metabolism (e.g. antidiabetic drugs, lipid-lowering drugs), gastrointestinal disease influencing absorption or a first-degree family history of diabetes mellitus.

Diet, health and lifestyle data were assessed by questionnaires. The subjects’ mean energy intake was calculated on the basis of their BMR, taking into account their daily physical activity\(^2\)\(^{21}\).

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 Ethics Committee of the Hospital District of Helsinki and Uusimaa. Written informed consent was obtained from all subjects.

**Postprandial study**

The subjects were advised to follow their usual diet during the study. In addition, they were also advised to consume at least 150 g carbohydrates/d during the 3 d before the test mornings. The subjects were not allowed to drink alcohol and were asked to avoid strenuous exercise and sauna for 24 h before each study day. The day before the study day, they were asked to eat an evening meal, in accordance with instructions, that would provide 15% of the calculated daily energy requirement. The carbohydrate content of the evening meals was 55% energy. The subjects were also asked to fast for 10–12 h after their standardised evening meal.

In the clinic, body weight was measured. Changes of up to 2 kg in weight were allowed during the study. An intravenous cannula was inserted into an antecubital vein in the forearm, and a finger-prick capillary blood sample (0.5 ml/sample) and an intravenous blood sample (8 ml/sample) were drawn. Thereafter, the subjects consumed the test meal within 10 min. After the start of the meal, finger-prick capillary blood samples were taken at 15, 30, 45, 60, 90 and 120 min, and intravenous blood samples were collected at 30, 60, 120, 180, 240 and 300 min.

**Study meals**

All subjects consumed two different test meals, a HGI meal (calculated GI 81) and a LGI meal (calculated GI 33), and twice a reference meal, a glucose solution, in randomised order at 1-week intervals. The GI of the study meals was calculated using the recommended method\(^2\)\(^{22}\), and the GI values of each component of the meals were based on the GI database of the National Institute for Health and Welfare\(^2\)\(^{22}\). The test meals and the reference meal contained 50 g of available carbohydrate. Both the test meals contained the same amount of energy, protein, fat and fibre. The foodstuffs and the nutrient composition of the study meals are shown in Table 1. The energy nutrient contents of the meals were in accordance with the Nordic Nutrition Recommendation\(^2\)\(^{23}\) and included 55% energy as carbohydrate, 15% energy as protein and 30% energy as fat. All meals were served with a 150 ml drink of the subjects’ choice (water, coffee or tea); the selected drink was the same for all test meals. Water was chosen by twelve (three with normal-weight NGT, four with overweight NGT, two with normal-weight IGT and three with overweight IGT), coffee by twenty-seven (eight with normal-weight NGT, five with overweight NGT, six with normal-weight IGT and eight with overweight IGT) and tea by nine (one with normal-weight NGT, three with overweight NGT, four with normal-weight IGT and one with overweight IGT) subjects.

**Chemical composition of the test meals**

The chemical composition of the test meals was analysed by AnalyCen Laboratory (Lidköping, Sweden). The protein content of the meals was estimated by the method of Kjeldahl\(^2\)\(^{24}\), and the fat content by a modified method of Schmid–Bondzynski–Ratzlaff\(^2\)\(^{25}\). Free sugars (glucose, fructose, lactose, maltose and sucrose) were determined by the Dionex ion chromatograph system, and the starch contents of the test meals were analysed by the modified Aman & Hesselman method\(^2\)\(^{26}\). Total fibre was analysed by an enzymatic gravimetric procedure (Association of Official Analytical Chemists 45.4.07)\(^2\)\(^{27}\). The amount of available carbohydrate was calculated as the sum of free sugars and enzymatically available starch.

**Laboratory analysis**

Capillary blood glucose was analysed directly by using a glucose meter (Glucose 201 meter; HemoCue Limited, Espoo, Finland). The HemoCue Glucose system is based on a glucose dehydrogenase method. The results were automatically converted to express plasma values. A quality-control solution...
Impact of overweight and glucose tolerance

Samples were transferred to a refrigerator (samples were clotted 10 min at room temperature. Thereafter, insulin samples, and the separated serum was stored at 270°C until analysis. Blood for serum insulin was measured by a microparticle enzyme enzymatic method (Abbott Laboratories). The fasting total cholesterol at baseline was analysed using an enzymatic colorimetric method (Wako Chemicals, Neuss, Germany) and an enzymatic glycerol-3-phosphate oxidase method (Abbott Laboratories), respectively. During the course of the study, the mean CV of the different control levels between days for insulin, NEFA and TAG measurements were 6·8 (sd 0·7), 1·8 (sd 0·2) and 1·4 (sd 0·5)%., respectively.

**Statistical analysis**

The 2 h incremental area under the glucose and insulin response curve (IAUC), the 5 h incremental area under the TAG response curve and the incremental area over the NEFA response curve were calculated using a trapezoidal method for each test meal. The RGR of the meals was calculated from the 2 h incremental glucose area using glucose as a reference.

Individual RGR values >2 standard deviations from the mean were excluded from the results of the test meals according to the same standard praxis compared with the GI value measurement. Therefore, two RGR values were excluded for each test meal. Insulin curves that included at least one strongly haemolysed serum sample or more than two mildly haemolysed serum samples were excluded from the analysis. Only two insulin curves were excluded from the statistical analysis. Insulin resistance was measured by the homeostasis model assessment of insulin resistance, and it was calculated as described by Matthews et al.

The independent sample t test with Bonferroni’s corrections was used for testing the differences between study groups. Insulin responses were non-normally distributed; statistical significance was therefore assessed by using the non-parametric Wilcoxon test. All statistical analyses were done using SPSS for Windows version 15·0 (SPSS Inc., Chicago, IL, USA). The level of significance was P<0·05. Results are expressed as means with their standard errors or standard deviations.

**Results**

**Subjects’ characteristics**

Baseline characteristics of the subjects are illustrated in Table 2. Fasting serum glucose, TAG and NEFA levels did not differ between the groups, whereas fasting insulin levels (P<0·001) and the homeostasis model assessment of insulin resistance (P<0·001) were significantly higher among overweight subjects with IGT than among normal-weight subjects with NGT.

**Glucose and insulin responses**

Mean glucose concentrations during the 2 h after the LGI and HGI meals are shown in Fig. 1, and the postprandial responses

<table>
<thead>
<tr>
<th>Low-GI meal</th>
<th>Components</th>
<th>g/portion</th>
<th>ACHO/ component</th>
<th>GI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rye bread*</td>
<td>30-0</td>
<td>10·6</td>
<td>50</td>
<td></td>
</tr>
<tr>
<td>Margarine 70 %†</td>
<td>7·5</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cheese 24 %‡</td>
<td>13-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cucumber</td>
<td>40-0</td>
<td>0·6</td>
<td>39</td>
<td></td>
</tr>
<tr>
<td>Barley porridge§</td>
<td>160-0</td>
<td>19·7</td>
<td>35</td>
<td></td>
</tr>
<tr>
<td>Water</td>
<td>30-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Home-made raspberry juice‖</td>
<td>170-0</td>
<td>19·4</td>
<td>23</td>
<td></td>
</tr>
<tr>
<td>Coffee¶ or tea** or water</td>
<td>150-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>Available carbohydrate</td>
<td>50-3</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Protein</td>
<td>14-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fat</td>
<td>12-1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total fibre</td>
<td>7-2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Energy (KJ)</td>
<td>1527-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Calculated GI</td>
<td>33</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>High-GI meal</th>
<th>Components</th>
<th>g/portion</th>
<th>ACHO/ component</th>
<th>GI</th>
</tr>
</thead>
<tbody>
<tr>
<td>White bread††</td>
<td>36-0</td>
<td>18-7</td>
<td>79</td>
<td></td>
</tr>
<tr>
<td>Margarine 70 % †</td>
<td>6-5</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cheese 24 % †</td>
<td>24-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cucumber</td>
<td>40-0</td>
<td>0-6</td>
<td>39</td>
<td></td>
</tr>
<tr>
<td>Wheat porridge‡‡</td>
<td>160-0</td>
<td>14-1</td>
<td>76</td>
<td></td>
</tr>
<tr>
<td>Water</td>
<td>30-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rye fibre</td>
<td>5-0</td>
<td>1-3</td>
<td>40</td>
<td></td>
</tr>
<tr>
<td>Home-made raspberry juice§§</td>
<td>135-0</td>
<td>15-4</td>
<td>93</td>
<td></td>
</tr>
<tr>
<td>Coffee¶ or tea** or water</td>
<td>150-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>Available carbohydrate</td>
<td>50-1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Protein</td>
<td>14-2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fat</td>
<td>11-9</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total fibre</td>
<td>7-1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Energy (KJ)</td>
<td>1523-0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Calculated GI</td>
<td>81</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

GI, glycaemic index; ACHO, available carbohydrate.

* Whole-grain rye bread (REAL-ruisleipa®, Fazer Limited, Vantaa, Finland).
† Flora margarine 70 % (Unilever Limited, Helsinki, Finland).
‡ Edam cheese 24 % (Valio Limited, Helsinki, Finland).
§ Barley porridge prepared with lactose-free (1·5 % fat) milk.
‖ Filtered coffee (Juhlamokka; Gustav Paulig Limited, Helsinki, Finland).
¶ Flora margarine 70 % (Unilever Limited, Helsinki, Finland).
** Instant tea (Lipton Yellow Label; Unilever Limited).
§§ Sweetened by 13·5 g glucose.

Table 1. Foodstuffs and energy nutrient content of the test meals
expressed as IAUC are shown in Table 3. Overweight subjects with NGT had a similar 2 h glucose IAUC to normal-weight subjects with NGT after both the LGI and HGI meals, whereas among subjects with IGT, the 2 h glucose IAUC after the test meals were significantly greater than that among normal-weight subjects with NGT (Table 3). The LGI meal produced a glucose response that was significantly smaller – about half – than that for the HGI meal among all of the study groups (P < 0.001; Table 4). Neither did the BMI nor the glucose tolerance had a significant effect on the RGR of the meals (Table 3). The mean RGR for the HGI meal was 74, and that for the LGI meal was 36.

When comparing the 2 h insulin responses of the meals between normal-weight subjects with NGT and the other study groups, only the insulin responses of the overweight subjects with IGT differed significantly (Table 3). The LGI meal produced smaller insulin responses than the HGI meal in all of the study groups (P < 0.001; Fig. 2 and Table 3). After the LGI meals, the insulin IAUC was about 30–40% lower than the insulin IAUC after the HGI meals (Table 4). The HGI meal increased the postprandial glucose and insulin responses in the same ratio among normal-weight and overweight subjects with NGT or IGT (Table 4).

TAG and NEFA responses

The mean TAG and NEFA postprandial responses during 5 h after the test meals are presented in Table 3. Neither did the BMI nor the glucose tolerance have a significant effect on the TAG and NEFA responses (Table 3). The TAG responses were higher after the LGI meal than after the HGI meal in all of the study groups (P < 0.001; Table 4). After the LGI meals, the TAG concentration was about 30–40% lower than that after the HGI meals. The mean TAG concentration after the LGI meal was 36.

When comparing the 2 h insulin responses of the meals between normal-weight subjects with NGT and the other study groups, only the insulin responses of the overweight subjects with IGT differed significantly (Table 3). The LGI meal produced smaller insulin responses than the HGI meal in all of the study groups (P < 0.001; Fig. 2 and Table 3). After the LGI meals, the insulin IAUC was about 30–40% lower than the insulin IAUC after the HGI meals (Table 4). The HGI meal increased the postprandial glucose and insulin responses in the same ratio among normal-weight and overweight subjects with NGT or IGT (Table 4).

TAG and NEFA responses

The mean TAG and NEFA postprandial responses during 5 h after the test meals are presented in Table 3. The TAG responses were higher after the LGI meal than after the HGI meal in all of the study groups (P < 0.001), but no statistically significant differences were observed between the groups (Table 3 and Fig. 3). After the test meals, NEFA concentration...
Our findings therefore suggest that the LGI meal is more beneficial in preventing elevated postprandial glucose and insulin levels than the HGI meal. The subjects’ physiological characteristics did not modify the results. Thus, the HGI meal increased the postprandial responses at the same ratio among normal-weight and overweight subjects with NGT or IGT.

In the present study, the test meals were served with a drink of the subjects’ choice (water, coffee or tea). Caffeine in coffee and tea may increase the glucose responses and decrease insulin sensitivity. However, in our previous study, coffee modified postprandial glucose and insulin responses only modestly (unpublished results), an observation that is consistent with the study of Aldughpassi & Wolever. Therefore, it is unlikely that the drink served influenced the results.

In the present study, neither the subjects’ BMI nor their glucose tolerance affected the RGR of the test meals, although the glucose responses were higher among subjects with IGT than among subjects with NGT. The present results are in line with previous studies where BMI, glucose tolerance, sex, age, and ethnicity did not have a major effect on the GI values of foods.

### Discussion

In the present study, we investigated the influence of overweight and glucose tolerance on the postprandial glucose, insulin, TAG and NEFA responses to the HGI and LGI meals. In addition, we studied the effect of the test subjects’ BMI and glucose tolerance on the postprandial glucose, insulin, TAG and NEFA responses to the HGI and LGI meals.

levels than the HGI meal. The subjects’ physiological characteristics did not modify the results. Thus, the HGI meal increased the postprandial responses at the same ratio among normal-weight and overweight subjects with NGT or IGT.

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### Table 3. Glucose, insulin and TAG incremental area under the curve (IAUC) and NEFA incremental area over the curve (IAOC) after the low-glycaemic index (GI) and the high-GI meals and the relative glucose responses of the test meals

<table>
<thead>
<tr>
<th>Test meal</th>
<th>Normal-weight NGT</th>
<th>Overweight NGT</th>
<th>Normal-weight IGT</th>
<th>Overweight IGT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glucose 2 h IAUC (mmol\texttimes\text{min/l})</td>
<td>80* 14</td>
<td>100* 10</td>
<td>2* 0·24</td>
<td>139* 13</td>
</tr>
<tr>
<td>Insulin 2 h IAUC (pmol \times \text{min/l})</td>
<td>12307* 1730</td>
<td>26340* 10248</td>
<td>0·21</td>
<td>15718* 1638</td>
</tr>
<tr>
<td>TAG 5 h IAUC (mmol \times \text{min/l})</td>
<td>38* 8</td>
<td>51* 10</td>
<td>0·36</td>
<td>59* 9</td>
</tr>
<tr>
<td>NEFA 5 h IAOC (mmol \times \text{min/l})</td>
<td>80 16</td>
<td>87 21</td>
<td>0·78</td>
<td>62 20</td>
</tr>
<tr>
<td>Relative glycaemic response</td>
<td>37*§ 4</td>
<td>29§ 2</td>
<td>0·12</td>
<td>36* 3</td>
</tr>
<tr>
<td>Glucose solution</td>
<td>183 22</td>
<td>252 50</td>
<td>0·22</td>
<td>303 23</td>
</tr>
<tr>
<td>Insulin 2 h IAUC (pmol \times \text{min/l})</td>
<td>19737 2690</td>
<td>33678 9477</td>
<td>0·18</td>
<td>21966 1951</td>
</tr>
<tr>
<td>TAG 5 h IAUC (mmol \times \text{min/l})</td>
<td>27 10</td>
<td>22 8</td>
<td>0·66</td>
<td>33 9</td>
</tr>
<tr>
<td>NEFA 5 h IAOC (mmol \times \text{min/l})</td>
<td>63 18</td>
<td>83 13</td>
<td>0·39</td>
<td>56 10</td>
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<tr>
<td>Relative glycaemic response</td>
<td>79 5</td>
<td>69 5</td>
<td>0·17</td>
<td>73§ 5</td>
</tr>
</tbody>
</table>

**Table 4. Proportion of the low-glycaemic index (GI) meal glucose and insulin postprandial responses to the high-GI meal responses**

<table>
<thead>
<tr>
<th>Test meal</th>
<th>Normal-weight NGT</th>
<th>Overweight NGT</th>
<th>Normal-weight IGT</th>
<th>Overweight IGT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glucose (%)</td>
<td>45 6</td>
<td>51 9</td>
<td>47 5</td>
<td>53 4</td>
</tr>
<tr>
<td>Insulin (%)</td>
<td>64 5</td>
<td>73 7</td>
<td>72 4</td>
<td>71 5</td>
</tr>
</tbody>
</table>

**IP address:** 54.70.40.11, on 26 Jan 2019 at 12:12:33, subject to the Cambridge Core terms of use, available at https://www.cambridge.org/core/terms. https://doi.org/10.1017/S0007114510005477
Overweight is typically associated with a pattern of dyslipidaemia characterised by elevated levels of fasting and postprandial TAG and NEFA levels\(^\text{a29,30}\). In the present study, however, neither fasting concentrations nor postprandial responses of TAG and NEFA differed between normal-weight and overweight subjects. Because the previous day’s eating and alcohol consumption modify TAG concentrations, and therefore daily variations in fasting TAG levels are common\(^\text{a37}\), in the present study, the subjects were asked to follow their usual diet during the study, and they were not allowed to drink alcohol for 24 h before each study day. In addition, they were advised to eat a standardised evening meal preceding the study day. In the present study, the intra- and inter-individual variation of fasting TAG concentrations were modest. Only two subjects had a 1 mmol/l difference in fasting TAG concentrations between the test meals. These modest variations may have had an impact on the results.

Ingestion of the LGI meal resulted in elevated TAG levels in comparison with the HGI meal, which was not expected. It has been proposed that LGI meals have a beneficial effect on postprandial TAG responses\(^\text{a6}\). There are, however, studies that have not detected any effect of the GI of meals on postprandial TAG levels\(^\text{a11,13}\). In the present study, the fructose-sweetened juice may explain the higher TAG responses after the LGI meal. This finding is consistent with an earlier postprandial study\(^\text{a38}\). The harmful effect of fructose consumption may be caused by its ability to induce hepatic de novo lipogenesis and thus TAG production\(^\text{a39}\). It is, however, unknown whether small amounts of fructose, such as 17 g in our LGI meal, can induce higher postprandial TAG responses\(^\text{a40}\).

It has been suggested that after the glucose peak of the HGI meal, the glucose level decreases below the fasting level, which in turn causes an increased secretion of counter-regulatory hormones, e.g. cortisol and adrenaline. Counter-regulatory hormones stimulate the release of NEFA

Fig. 2. Mean responses in serum insulin after the consumption of (a) the low-glycaemic index (GI) meal and (b) the high-GI meal in normal-weight subjects with normal glucose tolerance (NGT; ——, n 12); overweight subjects with NGT (—•—, n 10); normal-weight subjects with impaired glucose tolerance (IGT; — Δ—, n 12); overweight subjects with IGT (—■—, n 12).

Fig. 3. Mean responses in serum TAG after the consumption of (a) the low-glycaemic index (GI) meal and (b) the high-GI meal in normal-weight subjects with normal glucose tolerance (NGT; ——, n 12); overweight subjects with NGT (—•—, n 12); normal-weight subjects with impaired glucose tolerance (IGT; — Δ—, n 12); overweight subjects with IGT (—■—, n 12).
from adipose tissue and thus increase NEFA concentrations\(^4\). Previous studies have replicated these findings in young lean and obese subjects\(^7,10\). In addition, Wolever et al.\(^3,13\) have shown that the GI and amount of carbohydrate of the meal have an effect on NEFA responses. However, in the present study, the GI of the test meals did not have a significant effect on the NEFA responses. Therefore, our findings are in agreement with previous studies\(^6,11,12,41\), suggesting that the GI of a meal has little or no effect on postprandial NEFA responses.

In conclusion, the present study shows that postprandial glucose responses are affected only by glucose tolerance, whereas overweight and IGT occurring simultaneously have an impact on insulin responses. Overweight and glucose tolerance do not have an effect on postprandial lipid responses. The present study confirms that the LGI meal causes lower postprandial glucose and insulin responses than the HGI meal. The LGI meal may cause higher TAG responses, especially when the meal contains fructose. The subjects’ physiological characteristics do not modify these effects. The RGR of the meals does not differ between normal-weight and overweight subjects with NGT or IGT.

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