Lipid profile and insulin sensitivity in rats fed with high-fat or high-fructose diets

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Abstract
The occurrence and severity of obesity- and insulin resistance-related disorders vary according to the diet. The aim of the present longitudinal study was to examine the effects of a high-fat or a high-fructose diet on body weight (BW), body fat mass, insulin sensitivity (IS) and lipid profiles in a rat model of dietary-induced obesity and low IS. A total of eighteen, 12-week-old male Wistar rats were divided into three groups, and were fed with a control, a high-fat (65 % lipid energy) or a high-fructose diet (65 % fructose energy) for 10 weeks. BW, body fat mass (H2O dilution method), IS (euglycaemic–hyperinsulinaemic clamp technique), plasma glucose, insulin, NEFA, TAG and total cholesterol were assessed before and at the end of 10-week period. Cholesterol was measured in plasma lipoproteins separated from pooled samples of each group and each time period by using fast-protein liquid chromatography. All rats had similar BW at the end of the 10-week period. Body fat mass was higher in the high-fat group compared to the control group. There was no change in basal glycaemia and insulinaemia. The IS was lower in the high-fat group and was unchanged in the high-fructose group, compared to the control group. Plasma TAG concentration and cholesterol distribution in lipoproteins did not change over time in any group. Plasma NEFA concentration decreased, whereas plasma TAG concentration increased over time, regardless of the diet in both cases. The 10-week high-fat diet led to obesity and low IS, whereas rats fed with the high-fructose diet exhibited no change in IS and lipidaemia. The high-fat diet had more deleterious response than high-fructose diet to induce obesity and low IS in rats.

Key words: insulin resistance; obesity; dyslipidemia; high-fat diet; high-fructose diet; rat

There is a growing prevalence of obesity and low insulin sensitivity (IS), often called insulin resistance, in human subjects. Thus, there is a need for an animal model to study the time course of these metabolic disturbances as well as their unhealthy consequences. Different animal models have been used to study obesity and IS, notably the rat, in which obesity can be caused by genetic mutations or induced by nutritional interventions. As human obesity is mainly due to nutritional habits, animal models of obesity and low IS induced by specific diets may be preferable to genetic models.

Various nutritional interventions have been used to induce obesity, low IS and dyslipidaemia in rats. High-fat diets have been shown to cause these metabolic disorders in previous studies, but there has been a large variability in the intensity of the metabolic changes(1–3). High-fructose diets have also been shown to lower IS and promote mild-to-severe dyslipidaemia(6–9). The differences in nutritional interventions, such as diet composition and interventional duration, have complicated the comparisons of these studies. Therefore, it is difficult to define the best nutritional intervention to induce obesity in an animal model that closely mimics the human disease. Longitudinal studies could be useful in determining the best diet to induce obesity and related disorders.

To our knowledge, this type of approach has never been conducted in rats. Here, we aimed to perform a longitudinal study to compare the effects of a high-fat diet and a high-fructose diet on body weight (BW), IS and plasma lipid profiles in rats.
Materials and methods

Animal groups and diets

Male Wistar rats (12 weeks old; Janvier, Le Genest Saint-Isle, France) were randomly separated into three groups (six per group): control, high fat or high fructose. According to their groupings, the rats were fed with a control diet (39.7% maize starch, 20% dextrose, 5.8% sunflower oil and 20.5% casein by weight), a high-fat diet (12.7% maize starch, 6.5% dextrose, 3.9% sunflower oil, 31.3% lard and 28.6% casein by weight) or a high-fructose diet (59.7% fructose, 5.8% sunflower oil and 20.5% casein by weight) for 10 weeks. The rats were housed in individual cages, with free access to the feed and water. The rats were maintained under a 12-h light–12-h dark cycle and a temperature of 22°C. The animals were housed at Oniris (National College of Veterinary Medicine, Food Science and Engineering, Nantes, France), according to the regulations for animal welfare of the French Ministry of Agriculture. The experimental protocol adhered to the European Union guidelines and was approved by the local animal use and care advisory committee.

Body weight and body fat mass

BW was recorded weekly. The body fat mass was determined by isotope dilution (2H2O; Eurisotop, Gif-sur-Yvette, France) on week 1 (before the dietary intervention) and at the end of the 10-week diet period (week 11). Blood samples (1 ml) were collected before and 2 h after a 2H2O injection (500 mg/kg BW). Plasma 2H2O concentrations were measured using Fourier-transformed IR spectroscopy (Bruker SA, Wissembourg, France).

Euglycaemic–hyperinsulinaemic clamp technique

The euglycaemic–hyperinsulinaemic clamp technique was performed before and at the end of the 10-week diet period. The catheter was inserted under anaesthesia into the jugular vein of an animal that was not fed overnight. Insulin (Actrapid; Novo Nordisk, Bagsvaerd, Denmark) was perfused [72 mU/kg (500 pmol/kg) for 1 min, then 18 mU/kg per min (125 pmol/kg per min) for 3 h] and glycaemia was measured at every 5 min. Glucose (20%; Braun Medical SAS, Boulogne Cedex, France) was perfused at a variable rate. The glucose infusion rate (mg/kg per min) was adjusted to attain and maintain the basal glycaemia. In hyperinsulinaemic conditions, GIR measures the insulin-mediated glucose uptake and is considered as a good reflection of insulin sensitivity.

Plasma lipid profiles

The basal plasma concentrations of total cholesterol, NEFA and TAG were assayed before and at the end of the 10-week diet period using enzymatic methods (Cholestech, was performed before and at the end of the 10-week diet period. The catheter was inserted under anaesthesia into the jugular vein of an animal that was not fed overnight. Insulin (Actrapid; Novo Nordisk, Bagsvaerd, Denmark) was perfused [72 mU/kg (500 pmol/kg) for 1 min, then 18 mU/kg per min (125 pmol/kg per min) for 3 h] and glycaemia was measured at every 5 min. Glucose (20%; Braun Medical SAS, Boulogne Cedex, France) was perfused at a variable rate. The glucose infusion rate (mg/kg per min) was adjusted to attain and maintain the basal glycaemia. In hyperinsulinaemic conditions, GIR measures the insulin-mediated glucose uptake and is considered as a good reflection of insulin sensitivity.

Table 1. Body weight, body fat mass, insulin sensitivity and plasma lipid profiles of rats in the control, high-fat and high-fructose groups

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>High fat</th>
<th>High fructose</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Week 1</td>
<td>Week 11</td>
<td>Week 1</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>SEM</td>
<td>Mean</td>
</tr>
<tr>
<td>Body weight (g)</td>
<td>401</td>
<td>6</td>
<td>526</td>
</tr>
<tr>
<td>Body fat mass (g)</td>
<td>35·6</td>
<td>3·3</td>
<td>63·3</td>
</tr>
<tr>
<td>Basal glycaemia (mmol/l)</td>
<td>5·51</td>
<td>0·11</td>
<td>4·93</td>
</tr>
<tr>
<td>Basal insulinaemia (pmol/l)</td>
<td>402</td>
<td>8·4</td>
<td>400</td>
</tr>
<tr>
<td>Total cholesterol (mmol/l)</td>
<td>1·89</td>
<td>0·15</td>
<td>2·01</td>
</tr>
<tr>
<td>TAG (mmol/l)</td>
<td>0·46</td>
<td>0·07</td>
<td>0·30</td>
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BW, body weight; GIR, glucose infusion rate.

* Mean values were significantly different among groups for the interaction between time and the type of diet (fixed-effects model analysis revealed that values in the high-fat group is significant at week 11, compared to the control group; P<0.05).
† Mean values were significantly different for the interaction between time and the type of diet; the effect of time and type of diet were not considered (P<0.05).
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§ n=5 for GIR (in hyperinsulinaemic conditions, GIR measures the insulin-mediated glucose uptake and is considered as a good reflection of insulin sensitivity).
|| At week 1, plasma total cholesterol concentration was different among groups.

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The plasma samples were pooled for each group and each time period (weeks 1 and 11), and the plasma lipoproteins were separated using a fast-protein liquid chromatography system (UNICORN 520; GE Healthcare, Pittsburgh, PA, USA). The cholesterol concentration was measured in each fraction.

Statistical analysis

Data analysis was performed using Statview software (version 5.0; SAS Institute Inc., Cary, NC, USA) and R software (version 2.10, lme4 package; R Foundation for Statistical Computing, Vienna, Austria). Data were expressed as mean values with the standard error of the mean. A linear mixed-effects model has been performed in order to study the effect of time, type of diet and the interaction between them for each variable. The mixed-effects models are the most efficient way to analyse repeated measurements data. A multiple comparison of means procedure with Tukey contrasts, adapted to the mixed-effects models, has been used when the interactions between time and the type of diet were significant. A significant difference has been considered for P value <0.05.

Results

Body weight and body fat mass

Table 1 gives the values of BW and body fat mass of the control, the high-fat and the high-fructose groups before and at the end of the 10-week period. The BW gain was similar for all groups. At week 11, the high-fat group had significantly higher (P<0.05) body fat mass compared to the control group. At week 11, compared to initial values from week 1, the high-fat diet caused a 393 (SEM 23)% increase in body fat mass and high-fructose diet caused a 139 (SEM 23)% increase.

Insulin sensitivity

No differences were observed in basal glycaemia and insulinaemia among groups. The high-fat group had lower IS compared to the control group at week 11, assessed by significantly lower (P<0.05) glucose infusion rate value. The high-fructose group had no difference in IS compared to the control group (Table 1).

Plasma lipid profiles

Plasma TAG concentration was not different among groups at any time. Plasma NEFA concentration decreased over time in all groups, regardless of the diet. At week 1, the plasma total cholesterol concentration was different among groups. Indeed, it was higher in the control group than in others. There was a 38 (SEM 12)% increase in total cholesterol concentration in rats fed with high-fat diet and a 15 (SEM 8)% increase in rats fed with high-fructose diet, but the difference did not reach level of significance (Table 1). Nevertheless, plasma total cholesterol concentration increased over time, regardless of the diet.

Discussion

The aim of the present study was to compare the effects of a high-fat and a high-fructose diet on obesity-related disturbances in rats by using a longitudinal approach. In the present study, all the basal values were in the range of normal values of rats. In the literature, various concentrations of plasma lipids have been measured in control rats. However, we did not find any study of obesity and IS in rats with the baseline values of all variables. Therefore, we used a longitudinal approach and composed the groups randomly. The plasma total cholesterol concentration was significantly different among groups at baseline level (week 1). The randomisation process used to compose the groups had been expected to abolish all the differences among groups, and all cholesterol concentrations have been measured at the same time, i.e. at the end of the study, to guarantee same analytical conditions. Unfortunately, the randomisation process failed to reach its objectives, at least in this regard (there was no other difference among the groups at week 1). The baseline values were then taken into account in the statistical analysis in order to nullify the pre-existing differences. Another factor is the age at which exposure to a high-fat or high-fructose diet started. It has been shown that adult rats fed with high-fructose diet produce signs of metabolic syndrome, but young rats do not(11). We therefore chose adult rats to conduct the present experiment.

We observed similar BW at the end of the 10-week period in all groups. However, rats fed with high-fat diet had higher body fat mass compared to the control rats. Increase in adiposity has previously been described in rats fed with a high-fat diet(2–5,12), and it was associated with increase in weight gain. An increase in BW alone does not necessarily represent obesity, but other factors have to be taken into account, such as changes in body composition. In the present study, the increase in body fat mass reflected the obesogenic property of the high-fat diet, which was due to lard being used as the source of dietary fat. This excess fat was stored in the form of adipose tissue in the body. In rats fed with high-fructose diet, we observed no change in BW and body fat mass compared to the control rats. Previous studies showed that administering high-fructose (60 % by weight) diet for about 10 weeks increased BW(6,8,13) and epididymal and retroperitoneal fat depots(13). However, de Moura et al.(13) observed no difference in BW in adult rats administered fructose, despite higher retroperitoneal, mesenteric and subcutaneous fat depots weights. This suggests that high-fructose diet does not necessarily lead to obesity.

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IS, as it measures the insulin-mediated glucose uptake under hyperinsulinaemic conditions) compared to the control group, as previously shown in several studies. Moreover, low IS has been shown to be associated with an increase in epididymal fat, and the present results of higher body fat mass in rats fed with high-fat diet are consistent with the previous results. However, basal glycemia and insulinemia were unchanged. Some studies using high-fat diet (lard or safflower oil) reported low IS, associated with hyperinsulinaemia and hyperglycaemia. We think that we could have assessed low IS by euglycaemic–hyperinsulinaemic clamp technique before the appearance of hyperglycaemic and hyperinsulinemic states, and that the present experiment could have been long enough to develop insulin resistance but not diabetes.

On the other hand, we found no change in IS, basal glycemia and insulinemia in rats fed with high-fructose diet. Many previous studies have reported hyperglycaemia, but Nakagawa et al. did not show any modification in glycemia after administering high-fructose diet. The higher hepatic glycochen content that has been described in rats fed with high-fructose diet could prevent hyperglycaemia. Previous studies have also shown that the high-fructose diet developed low IS and was associated with increased plasma TAG and NEFA concentrations. These increased concentrations in response to high-fructose diet could have an important role in the development of low IS by reducing insulin signalling pathway (reviewed in Tappy & Le). However, we did not find dyslipidemia, which is in accordance with the unchanged IS. Moreover, in a study in rats, high-sucrose diet has been shown to induce low IS in the liver before muscle. We assessed the IS by using the euglycaemic–hyperinsulinaemic clamp technique, which is the gold standard method for direct assessment of the whole-body IS. We suggest that high-fructose diet could have caused minor impairment of insulin action in the liver, but could not have caused whole-body low IS.

We observed no difference in plasma total cholesterol concentration between the groups, but there was a significant increase over time, regardless of the diet. This increase could have reflected an effect of age, whereas the absence of dietary cholesterol could possibly be the cause for unchanged cholesterol concentration.

Plasma basal NEFA concentration was not different among groups. Variable changes in plasma NEFA concentration have been described in response to a high-fat diet. The present results in rats fed with high-fructose diet were not consistent with some previous reports. We hypothesise that the liver could have higher hepatic TAG storage, by capturing a bulk amount of plasma NEFA, which would explain that no difference in plasma NEFA concentration was observed.

We found no change in plasma basal TAG concentration in any group. Some studies showed an increase in plasma TAG concentration at the end of a high-fat diet, whereas others with nearly the same fat content did not. Variable response on plasma basal TAG concentration has also been reported in rats fed with high-fructose diet. A few studies have reported higher postprandial TAG concentration in rats fed with high-fructose diet, but no change in basal TAG concentration for a short period (2 weeks) or for a long period (11 months). In the present study, we measured only the basal TAG concentration. We suggest that the rats in our study could have had higher postprandial TAG concentration that could not have been observed in overnight unfed rats.

On the basis of the present findings, feeding 12-week-old rats a high-fat diet for 10 weeks induces obesity and low IS. In contrast, the high-fructose diet produced no change in obesity-related disorders. In the future, it will be useful to study the specific effects of different fats, oils or fatty acids that account for the discrepancy between these studies.

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References


