Polyphenol-rich extract of pomegranate peel alleviates tissue inflammation and hypercholesterolaemia in high-fat diet-induced obese mice: potential implication of the gut microbiota

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Abstract

Pomegranate extracts have been used for centuries in traditional medicine to confer health benefits in a number of inflammatory diseases, microbial infections and cancer. Peel fruit are rich in polyphenols that exhibit antioxidant and anti-inflammatory capacities in vitro. Recent studies strongly suggest that the gut microbiota is an environmental factor to be taken into account when assessing the risk factors related to obesity. The aim of the present study was to test the prebiotic potency of a pomegranate peel extract (PPE) rich in polyphenols in a nutritional model of obesity associated with hypercholesterolaemia and inflammatory disorders. Balb/c mice were fed either a control diet or a high-fat (HF) diet with or without PPE (6 mg/d per mouse) over a period of 4 weeks. Interestingly, PPE supplementation increased caecal content weight and caecal pool of bifidobacteria. It did not significantly modify body weight gain, glycaemia, glucose tolerance and inflammatory markers measured in the serum. However, it reduced the serum level of cholesterol (total and LDL) induced by HF feeding. Furthermore, it counteracted the HF-induced expression of inflammatory markers both in the colon and the visceral adipose tissue. Together, these findings support that pomegranate constitutes a promising food in the control of atherogenic and inflammatory disorders associated with diet-induced obesity. Knowing the poor bioavailability of pomegranate polyphenols, its bifidogenic effect observed after PPE consumption suggests the involvement of the gut microbiota in the management of host metabolism by polyphenolic compounds present in pomegranate.

Key words: Pomegranates; Microbiota; Obesity; High-fat diet; Inflammatory disorders; Prebiotics; Polyphenols

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Abbreviations: COX-2, cyclo-oxygenase 2; CT, control; HF, high-fat; MAPK, mitogen-activated protein kinase; OGTT, oral glucose tolerance test; PPE, pomegranate peel extract.

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complex microbial ecosystem comprising a considerable metabolic versatility, using biological pathways that humans have not evolved. This capacity of the gut microbiome, encoded by the collective genomes of the gut microbiota or gut metagenome, includes the metabolism of indigestible polyphenols derived from fruit and vegetables. This last consideration is of particular interest knowing that the gut microbiota is increasingly considered as a symbiotic partner for the maintenance of health. Several data suggest that the activity of the gut microbiota is a factor to be taken into account when assessing the risk factors related to obesity, and associated disorders, such as dyslipidaemia, inflammation, insulin resistance and diabetes. Indeed, alterations in the composition of the gut microbiota – known as dysbiosis – have been proposed to contribute to the development of obesity, thereby supporting the potential interest of nutrients acting on the gut microbes to produce beneficial effects on host energy metabolism.

The purpose of the present study was to determine whether the oral administration of a polyphenol-rich extract of pomegranate peel to mice could modulate the gut microbiota composition and offset increases in body weight, cholesterol profile, glucose intolerance, lipid storage and inflammation occurring in mice fed a high-fat (HF) diet. We decided to perform this in vivo study on Balb/c mice since it was demonstrated that (1) a HF diet induces hepatic lipid accumulation at a higher extent in Balb/c mice than in C57BL6j mice and (2) Balb/c mice exhibit a higher capacity to respond to inflammatory stimulus (higher increase in plasma and hepatic levels of cytokines/chemokines upon caecal ligation and puncture inducing septic peritonitis) than C57BL6j mice.

**Experimental methods**

**Animals and diet**

A total of eighteen male Balb/c mice (9 weeks old at the beginning of the experiment, Charles River Laboratories) were housed in groups of three per cage in a controlled environment (12 h daylight cycle, lights off at 18.00 hours) with free access to food and water. After 1 week of acclimatisation, the mice were divided into three groups (six per group): a control (CT) group, fed a control diet (AO4, SAFE), a group fed a HF diet and a group fed the same HF diet and receiving pomegranate peel extract (HF-PPE) at a dose of 0.2% in tap water (resulting in average consumption of 6 mg/d per mouse). The full composition of both the HF diet (D12492, Research Diets) and the AO4 standard diet is given in Table S1, available online. Extract (OXYLENT GR®, Steranon S.A.) used in the study was derived from pomegranate peel in which polyphenol content reached 30% (Folin–Ciocalteu method, equivalent gallic acid) and the concentrations of punicalagin and ellagic acid were 8 and 5% (ultra-performance liquid chromatography method with diode array detection), respectively. Food intake was recorded taking into account spillage twice a week during 4 weeks. After 4 weeks and a 6 h period of fasting, mice were anaesthetised (ketamine/xylazine intraperitoneal, 100 and 10 mg/kg, respectively) and blood samples were harvested for further analysis. Liver, adipose tissues, caecal content and intestinal tissues (proximal colon and caecum) were carefully dissected and immersed in liquid N2 before storage at −80°C. The animal experiments were approved by the local ethics committee and housing conditions were as specified by the Belgian Law of 6 April 2010 on the protection of laboratory animals (agreement no. LA 1230514).

**Oral glucose tolerance test**

After 3 weeks of treatment, an oral glucose tolerance test (OGTT) was performed on 6 h-fasted mice. Glucose was administered orally (3 g/kg body weight, 66% glucose solution) and blood glucose levels were determined using a glucose meter (Roche Diagnostics) on 3.5 μL of blood collected from the tip of the tail vein both before (~30 min and 0 min) and after glucose administration (15, 30, 60, 90 and 120 min).

**Microbial analysis of the caecal contents**

At the end of the experiment, the total caecum content was collected and weighed before storage at −80°C. Quantitative PCR for total bacteria, *Bifidobacterium* spp., *Lactobacillus* spp., *Bacteroides–Prevotella* spp. and *Roseburia* spp. were performed as reported by Neyrinck et al. using Mesa Fast qPCR® (Eurogentec). Real-time PCR were performed with the StepOnePlus™ real-time PCR system and software (Applied Biosystems). The cycle threshold of each sample was then compared with a standard curve (performed in triplicate) made by diluting genomic DNA (5-fold serial dilution of *Bifidobacterium animalis* for *Bifidobacterium* spp., *Bacteroides fragilis* for *Bacteroides–Prevotella* spp., *Lactobacillus acidophilus* for *Lactobacillus* spp. and total bacteria) (BCCM/LMG, Ghent, Belgium). Cell counts before DNA extraction were determined by culture.

**Blood parameters**

Plasma insulin concentrations were determined using an ultrasensitive ELISA kit (Alpco™ immunoassay, Alere Healthcare). The insulin resistance index was calculated by multiplying the AUC for glucose, and the AUC for insulin, calculated from −30 min until 15 min after glucose challenge. Concentrations of IL-1α, IL-1β, IL-6, monocyte chemoattractant protein 1 (MCP-1), TNFα, IL-10 and IL-15 were determined in 15 μL of plasma using a multiplex immunoassay kit (BioPlex Cytokine Assay, Bio-Rad) and measured using Luminex® technology (BioPlex®, Bio-Rad). Plasma TAG, cholesterol and NEFA concentrations were measured using kits coupling enzymatic reaction and spectrophotometric detection of reaction end-products (Diays Diagnostic and Systems). HDL-cholesterol concentration was measured enzymatically after VLDL, chylomicron and LDL-cholesterol antibody precipitation (Diays Diagnostic and Systems). LDL was estimated by the Friedewald formula.
**Lipid analysis in the liver**

TAG and cholesterol were measured in the liver tissue after extraction with chloroform–methanol, as described by Neyrinck et al.\(^{19}\).

**Expression of selected genes in tissues**

Total RNA was isolated using the TriPure isolation reagent kit (Roche Diagnostics Belgium). Complementary DNA was prepared by reverse transcription of 1 μg total RNA using the Kit Reverse transcription System (Promega). Real-time PCR were performed with the StepOnePlus™ real-time PCR system and software (Applied Biosystems) using SYBR Green for detection according to the manufacturer’s instructions. RPL19 RNA was chosen as the housekeeping gene. Primer sequences for the targeted mouse genes are available on request (audrey.neyrinck@uclouvain.be). All samples were run in duplicate in a single ninety-six-well reaction plate and the data were analysed according to the 2^(-ΔΔCt) method. The identity and purity of the amplified product were checked through analysis of the melting curve carried out at the end of amplification.

**Statistical analysis**

Results are presented as means with their standard errors. Statistical analysis was performed by the Mann–Whitney test (GraphPad Software). \( P<0.05 \) was considered as statistically significant. A two-way analysis on repeated measures was performed for the evolution of body weight and the evolution of glycaemia upon OGTT.

**Results**

**Oral administration of pomegranate peel extract promotes the growth of Bifidobacterium spp. in the caecal content of mice upon high-fat feeding**

HF feeding decreased the weight of caecal content as compared to the control condition (Fig. 1(a)). Quantitative analysis performed in caecal content showed that HF diet decreased the number of total bacteria, the Gram-positive lactobacilli and Roseburia spp. as well as the Gram-negative bacteria such as Bacteroides–Prevotella spp. without effect on bifidobacteria content (Fig. 1). Surprisingly, the PPE treatment significantly increased the weight of caecal content and the caecal pool of Bifidobacterium spp. without modifying significantly other bacteria as compared to HF-fed mice. Of note, the increase of Bacteroides–Prevotella spp. due to PPE supplementation \( v \) the HF-fed group was nearly of significance (\( P=0.07 \)).

**The oral administration of pomegranate peel extract does not modify the high-fat-induced body weight gain, adiposity and glucose intolerance**

A drop in body weight occurred within the first 3 d of treatment in both groups receiving the HF diet, signalling an adaptation to the dietary changes (Fig. 2(a)). The body weight evolution did not reveal any significance about time, treatment and interaction (time \( \times \) treatment) after a repeated-measures analysis (two-way ANOVA). However, the HF diet significantly increased the body weight gain of mice when considering the difference between day 3 and day 30 (end of the experiment) (Fig. 2(b)). This effect was accompanied by fat accumulation as shown by the weight of the adipose tissues.
Oral administration of pomegranate peel extract reduces the high-fat-induced hypercholesterolaemia

HF feeding negatively affected cholesterol content in the serum (increase in total cholesterol, LDL-cholesterol and HDL-cholesterol), but decreased triacylglycerolaemia as compared to the CT group (Table 1). Interestingly, the administration of PPE was able to counteract the HF-induced hypercholesterolaemia, in particular total cholesterol and LDL-cholesterol without provoking any change in HDL-cholesterol. The lipid content in the liver was not different between the groups (Table 1).

Oral administration of pomegranate peel extract decreases high-fat-induced expression of inflammatory markers in the colon and the visceral adipose tissue

In contrast to what happens in the liver, a higher expression of inflammatory markers was observed both in the colon and the visceral adipose tissue in mice fed a HF diet v. control mice with a significant P value for IL-1β (Fig. 4). PPE administration significantly down-regulated COX-2 induction in colonic and adipose tissues but not in the liver. Moreover, PPE decreased the mRNA levels of IL-6 and IL-1β, with a significant P value for IL-6 mRNA in the colon and IL-1β mRNA in the visceral adipose tissue. Several inflammatory markers measured in the serum, including of IL-6 and IL-1β, were neither significantly affected by the HF diet nor by PPE administration (Data S1, available online).

Discussion

The health-promoting effect of plant constituents and extracts is subject to interesting developments, a phenomenon explaining that their consumption is on the rise in the Western world[1,2,25]. The health benefits of pomegranate fruit and/or juice consumption have recently received considerable scientific focus. PPE used in this study is a source of polyphenol (30%), with punicalagin concentration reaching 8%. When added in the tap water of HF-fed mice during 4 weeks, PPE did not significantly modify body weight, adiposity and glucose tolerance. However, it decreased the HF-induced inflammatory tone not only in the gastrointestinal tract but also in the adipose tissue. It has been proposed that the anti-inflammatory effect of the pomegranate extract was dependent on
both the inhibition of p38-mitogen-activated protein kinase (MAPK) pathway and inhibition of the activation of transcription factor NF-κB, as demonstrated in vivo in mouse skin exposed to 12-O-tetradecanoylphorbol-13-acetate, and in vitro in human chondrocytes (26,27). The activation of p38-MAPK and NF-κB is intimately associated with the increased gene expression of TNF-α, IL-1β, IL-6, monocyte chemoattractant protein 1 (MCP-1), inducible NO synthase (iNOS) and COX-2 that are critical mediators of inflammation and the pathogenesis of inflammatory diseases (28,29). In particular, COX-2-mediated inflammation in visceral fat plays a pivotal role in the development of metabolic disorders associated with obesity induced by HF feeding (30). In the present study, we have shown that PPE consumption down-regulated IL-1β, IL-6 and COX-2 in the colon and the visceral adipose tissue as compared to HF-fed mice without effect on the expression of those markers in the liver. These data demonstrate that bioavailable constituents of PPE or metabolites

![Fig. 3. Oral glucose tolerance test performed in mice fed a control (CT; (a), (b), (c), (d)) diet, a high-fat (HF; (a), (b), (c), (d)) diet or a HF diet with pomegranate peel extract (HF-PPE; (a), (b), (c), (d)) in tap water for 3 weeks. (a) Plasma glucose levels after the oral glucose load, (b) area under the curve (AUC) of the glucose excursion, (c) plasma insulin levels 30 min before and 15 min after the oral glucose load, (d) insulin resistance index. * Values were significantly different from those of CT (P<0.05). † Values were significantly different from those of CT (−30 min) (P<0.05).](image-url)

### Table 1. Lipid contents in the serum and the liver‡

<table>
<thead>
<tr>
<th></th>
<th>CT</th>
<th>HF</th>
<th>HF-PPE</th>
</tr>
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<tbody>
<tr>
<td><strong>Serum lipids (mM)</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>TAG</td>
<td>0·98 ± 0·13</td>
<td>0·59* ± 0·06</td>
<td>0·53* ± 0·03</td>
</tr>
<tr>
<td>NEFA</td>
<td>0·67 ± 0·07</td>
<td>0·47 ± 0·11</td>
<td>0·35 ± 0·06</td>
</tr>
<tr>
<td>Total cholesterol</td>
<td>1·51 ± 0·07</td>
<td>2·26* ± 0·07</td>
<td>1·92† ± 0·03</td>
</tr>
<tr>
<td>LDL-cholesterol</td>
<td>0·28 ± 0·06</td>
<td>0·79* ± 0·06</td>
<td>0·44† ± 0·08</td>
</tr>
<tr>
<td>HDL-cholesterol</td>
<td>0·79 ± 0·02</td>
<td>1·20* ± 0·07</td>
<td>1·23† ± 0·06</td>
</tr>
<tr>
<td><strong>Liver lipid content (nmol/mg protein)</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>TAG</td>
<td>283 ± 29</td>
<td>357 ± 45</td>
<td>378 ± 35</td>
</tr>
<tr>
<td>Total cholesterol</td>
<td>207 ± 21</td>
<td>210 ± 54</td>
<td>176 ± 22</td>
</tr>
</tbody>
</table>

CT, control; HF, high-fat; PPE, pomegranate peel extract.
* Values were significantly different from those of CT (P<0.05).
† Values were significantly different from those of HF (P<0.05).
‡ Mice were fed a CT diet, a HF diet or a HF diet with PPE in tap water for 4 weeks.
Health benefits of pomegranate extracts

Fig. 4. mRNA levels of inflammatory markers in (a) the colon, (b) the visceral adipose tissue and (c) the liver of mice fed a control (CT) diet, a high-fat (HF) diet or a HF diet with pomegranate peel extract (HF-PPE) in tap water for 4 weeks. Values are expressed relative to CT group (set at 1). * Values were significantly different from those of CT (P<0.05). † Values were significantly different from those of HF (P<0.05).

coming from the gut bacteria may reach host tissue such as visceral adipose tissue to exert their anti-inflammatory effects.

In fact, after drinking pomegranate juice containing punicalagins, ellagic acid was detected in plasma, suggesting acid hydrolysis of at least some of the ellagitanins releasing free ellagic acid, which is absorbed directly from the stomach or the proximal small intestine(32). Ellagitannin has also been detected in the plasma of rats after oral administration(31). When the ellagitanins and/or ellagic acid reach the distal part of the small intestine and the colon, they are metabolised by the gut microbiota producing urolithins A and B, which are then absorbed along with ellagic acid(6,32,33). Once the metabolites are absorbed, glucuronidation occurs in the intestinal cells and glucuronides are the main metabolites found in portal vein plasma(34). The metabolites are further metabolised in the liver to produce diglucuronides, and/or sulphates, to produce a whole combination of metabolites secreted in the bile and in the urine. It seems that urolithins are absorbed preferentially, as their lipophilicity increases with plasma and that they undergo active enterohepatic circulation(32,33).

Interestingly, it was shown that urolithins had anti-inflammatory action on colon fibroblasts activated with IL-1β(35). Results obtained by Gonzalez-Sarrias et al.(37) pointed out that kinase signalling pathways (MAPK) may be involved in the response of Caco-2 cells to urolithins. Therefore, it suggests that the gut microbiota could contribute to the anti-inflammatory effects of PPE through their capacity to metabolise polyphenols.

It is known that phenolic components of common foods readily contribute to gut bacteria modulations(30,38,39). Bifidobacteria served as a model for the concept of prebiotics, which has been defined as "the selective stimulation of growth and/or activity(ies) of one or a limited number of microbial genus(era)/species in the gut microbiota that confer(s) health benefits to the host"(40). Interestingly, the number of bifidobacteria was inversely correlated with the development of fat mass, glucose intolerance, adipose tissue inflammation and lipopolysaccharide level(37,41,42). Our data support that the gut microbiota may be an important actor in the health beneficial action of PPE. Indeed, we had demonstrated that PPE increased the caecal pool of bifidobacteria. Although, the potential prebiotic activity of pomegranate products has not been recognised yet, a recent study has shown that the pomegranate by-product, which is rich in oligomers composed of 2–10 repeating units of gallic acid, ellagic acid and glucose in different combinations, enhanced the growth of total bacteria, in particular Bifidobacterium spp., as well as the concentrations of SCFA using pH-controlled, stirred,
batch culture fermentation systems reflective of the distal region of the human large intestine in the fermentation medium\(^{(35)}\). It is worth noting that punicalagins did not affect the growth of bacteria in this in vitro system.

In addition to changes in the gut microbiota, the most important effect of PPE supplementation was to blunt HF-induced hypercholesterolaemia (total and LDL-cholesterol). Recently, we and other authors provided evidence that modulation of the gut microbiota-host metabolic interrelationship by dietary intervention has the potential to improve cholesterol homeostasis, which has relevance for cardiovascular health\(^{(44,45)}\). Martinez et al.\(^{(44)}\) have highlighted through correlation analysis that the Bifidobacterium/Coriobacteriaceae equilibrium was important for plasma cholesterol levels in hamsters, with bifidobacteria being beneficial and coriobacteria being detrimental. In fact, the gut microbiota may affect host cholesterol metabolism through the modulation of the enterohepatic circulation of bile acids\(^{(46)}\). We postulate that the modulation of the gut microbiota induced by PPE supplementation observed in the present study can be involved in its hypocholesterolaemic effects.

In conclusion, the present study has shown that oral administration of a PPE rich in polyphenols is able to modulate the gut microbiota in favour of bifidobacteria. This prebiotic effect was accompanied by a lower expression of key inflammatory genes in the colon and the visceral adipose tissue. The prebiotic effect of PPE supplementation observed in the present study can be involved in its hypocholesterolaemic effects.

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Supplementary table S1 and data are available online at http://www.journals.cambridge.org/bjn

**References**


