Blueberry intervention improves vascular reactivity and lowers blood pressure in high-fat-, high-cholesterol-fed rats

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Abstract
Growing evidence suggests that intake of flavonoid-containing foods may exert cardiovascular benefits in human subjects. We have investigated the effects of a 10-week blueberry (BB) supplementation on blood pressure (BP) and vascular reactivity in rats fed a high-fat/high-cholesterol diet, known to induce endothelial dysfunction. Rats were randomly assigned to follow a control chow diet, a chow diet supplemented with 2 % (w/w) BB, a high-fat diet (10 % lard; 0·5 % cholesterol) or the high fat plus BB for 10 weeks. Rats supplemented with BB showed significant reductions in systolic BP (SBP) of 11 and 14 %, at weeks 8 and 10, respectively, relative to rats fed the control chow diet (week 8 SBP: 107·5 (SEM 4·7) v. 122·2 (SEM 2·1) mmHg, P=0·018; week 10 SBP: 115·0 (SEM 3·1) v. 132·7 (SEM 1·5) mmHg, P=0·0001). Furthermore, SBP was reduced by 14 % in rats fed with the high fat plus 2 % BB diet at week 10, compared to those on the high-fat diet only (SBP: 118·2 (SEM 3·6) v. 139·5 (SEM 4·5) mmHg, P=0·0001). Aortas harvested from BB-fed animals exhibited significantly reduced contractile responses (to L-phenylephrine) compared to those fed the control chow or high-fat diets. Furthermore, in rats fed with high fat supplemented with BB, aorta relaxation was significantly greater in response to acetylcholine compared to animals fed with the fat diet. These data suggest that BB consumption can lower BP and improve endothelial dysfunction induced by a high fat, high cholesterol containing diet.

Key words: Blueberries; Flavonoids; Atherosclerosis; Vasorelaxation; Diet; Blood pressure

CVD, in particular CHD and stroke, is a major cause of mortality worldwide(1). A number of genetic and environmental factors play a role in the initiation and progression of CVD, and one of the primary dietary risk factors for CVD is the consumption of diets high in saturated fat(2). Saturated fat is believed to increase cardiovascular risk through its effects on atherosclerosis, ‘endothelial dysfunction’ and ultimately hypertension(3), all prognostically relevant events in CVD(4). Endothelial dysfunction is characterised by a number of physiological changes, including the decreased bioavailability of endothelium-derived vasodilators, primarily NO (NO) and the increased plasma levels of endothelial-derived contracting factors(5). Such changes may result from the dysfunction of endothelial NO synthase, as deficiencies in endothelial NO synthase have been linked to the development of atherosclerosis in animals(6,7) and hypertension in human subjects(8). Indeed, hypertension is the primary clinical diagnostic risk factor for CVD(9).

In rats, hypertension has been associated with an impairment of NO-dependent endothelial function(10). Epidemiological and medical anthropological investigations suggest that flavonoid-rich foods exert cardiovascular health benefits(11–13) and their intake has been shown to lower blood pressure (BP) in both hypertensive(14,15) and normotensive individuals(16,17). The beneficial vascular effects of flavonoids are probably mediated by the ability of absorbed flavonoids and/or their circulating metabolites to increase the bioavailability of NO(18–20). In vitro studies have indicated that flavonoids are capable of directly activating endothelial NO synthase(21,22). Blueberries (BB) are a rich source of flavonoids, in particular anthocyanins and flavanols(23,24), and have been shown to induce improvements in cognitive performance(25,26), to inhibit oxidative stress and inflammation(27) and to promote beneficial vascular effects(28–32). Furthermore, in a randomised controlled human intervention study, favourable

Abbreviations: Ach, acetylcholine; BB, blueberry; BP, blood pressure; BW, body weight; Phe, L-phenylephrine; SNP, sodium nitroprusside.

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changes in platelet function, HDL-cholesterol and BP were found after consumption of berries for 8 weeks in elderly individuals with CVD risk factors. In animal models of hypertension, BB supplementation has been shown to lower BP, and also to affect endothelium-mediated vasorelaxation. However, to our knowledge, no studies have investigated the effect of a BB-enriched diet on vascular function in rats fed a high-fat/high-cholesterol diet, reflective of a typical Western diet.

In the present study, we examined the effect of BB supplementation on BP in both normal and high-fat/high-cholesterol-fed animals. Furthermore, we investigated the impact of 10 weeks of BB consumption on the health of the vasculature by assessing the responsiveness of isolated aortic rings (from fed animals) to L-phenylephrine (Phe; endothelium-dependent contraction), acetylcholine (Ach; endothelium-dependent vasodilation) and sodium nitroprusside (SNP; endothelium-independent vasodilation).

Materials and methods

Materials

All chemicals were purchased from Wako Pure Chemical Industries unless otherwise stated.

Animals and diets

A total of thirty-two male Wistar rats (180–200 g, 8 weeks of age) were obtained from Japan SLC. Animals were housed individually in a temperature-controlled room (25°C) with a 12 h light–12 h dark cycle. Body weights (BW) were measured individually in a temperature-controlled room (25°C) with a 20 via the tail-cuff method, using an oscillometric method (Apex Processing Technology). The freeze-dried BB powder was analysed for anthocyanin and procyanidin content according to the established methods in our laboratory, as previously described. Diets were prepared fresh and were stored at 4°C for a maximum of 2–5 days following preparation. All rats were provided with water and food ad libitum for the duration of the experiment. All animal procedures were in accordance with the institutional guidelines for the care and use of laboratory animals of the University of Tokushima.

Blood pressure measurements

Systolic BP was measured in all rats at weeks 1, 2, 4, 6, 8 and 20 via the tail-cuff method, using an oscillometric method.

Table 1. Composition of the diets used in the present study: control chow diet (control diet), control chow with 2 % (w/w, dry weight) freeze-dried BB powder, a high-fat and high-cholesterol diet consisting of the control chow diet plus 10 % (w/w) of lard and 0·5 % (w/w) cholesterol and a high-fat and high-cholesterol diet plus 2 % (w/w) freeze-dried BB powder. The control chow diet was purchased from the Oriental Yeast Company Limited. The composition of the control chow diet (per 100 g diet) was: water, 7·8 g; protein, 17·9 g; fat, 5·3 g; fibre, 8·6 g; energy, 1218 kJ (291 kcal); cholesterol, 1 mg; and vitamin E, 9·1 mg. The fatty acid composition of the control chow diet was: 16 : 0, 15·9 %; 18 : 0, 2·2 %; 18 : 1, 13·5 %; 18 : 2, 40·7 %; 18 : 3, 18·7 % and others less than 1 %. The fatty acid composition of the lard was: 16 : 0, 27 %; 16 : 1, 2·5 %; 18 : 0, 13 %; 18 : 1, 46 %; and 18 : 2, 8 %. The composition of the diets is shown in Table 1.

Highbush BB of unknown variety were purchased from a local supplier (Axons of Southampton) and immediately frozen at −20°C. Frozen BB were freeze-dried (−25°C; 6 mbar vacuum) in an Edwards MFD 01 freeze drier (Edwards) and ground to a fine powder using an Apex Comminuting Mill (Apex Processing Technology). The freeze-dried BB powder was kept at −20°C and shipped from the UK to Japan, where the animal experiments were conducted.

The freeze-dried BB powder was analysed for anthocyanin and procyanidin content according to the established methods in our laboratory, as previously described. Diets were prepared fresh and were stored at 4°C for a maximum of 2–5 days following preparation. All rats were provided with water and food ad libitum for the duration of the experiment. All animal procedures were in accordance with the institutional guidelines for the care and use of laboratory animals of the University of Tokushima.

<table>
<thead>
<tr>
<th>Type fat (percentage in diet)</th>
<th>Control diet</th>
<th>2 % BB diet</th>
<th>High-fat diet</th>
<th>2 % BB + high-fat diet</th>
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<td>0·51</td>
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<td>3·16</td>
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<tr>
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<td>0·25</td>
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<td>Total flavonoids measured</td>
<td>0</td>
<td>20·26</td>
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<td>20·26</td>
</tr>
</tbody>
</table>

* The amount of each type of fat is expressed in percentage of total fat in the diets. The amount of total measured flavonoids is expressed in mg/100 g of diet.
Preparation of aortic rings and tension measurements

Tension measurements on aortic rings were performed according to the established procedures with some modifications\(^{36}\). At the culmination of the 10-week supplementation, rats were anaesthetised using diethyl ether and the thoracic aortas were dissected (free of connective tissue) and then cut into ring segments, 3–4 mm in length. Each ring was placed in a 3 ml organ bath (Micro Easy Magnus, Kishimoto Medical) and mounted on two stainless steel wires, one of which was fastened to the bath and the other connected to a force transducer for the measurement of isometric tension. The bath was filled with Krebs–Ringer bicarbonate buffer solution at 37°C and bubbled with a mixture of 95 % CO\(_2\) and 5 % O\(_2\). The Krebs–Ringer bicarbonate buffer contained (in mmol/L) 118 NaCl, 4.6 KCl, 2.5 CaCl\(_2\), 24.8 NaHCO\(_3\), 1.2 MgSO\(_4\), 1.2 KH\(_2\)PO\(_4\) and 5.6 glucose. Each aortic ring was equilibrated for 60 min under a resting tension of 1.5 g and the Krebs–Ringer bicarbonate solution was changed at 30 min intervals. Cumulative concentration–response curves to the endothelium-dependent vasodilator Ach and the endothelium-independent vasodilator SNP were performed. To test the relaxation responses of Ach and SNP, the aortic rings were pre-contracted with the \(\alpha\)-adrenergic agonist, Phe (10\(^{-6}\) M; 10 min), until the contraction curve reached a plateau. Following Phe-induced contraction, cumulative applications of Ach (10\(^{-9}\) to 10\(^{-5}\) M) or SNP (10\(^{-8}\) to 10\(^{-3}\) M) were applied for 5 min, during which maximum aortic ring relaxation was achieved. For quantification, the relaxant effect to each Ach or SNP dose was expressed as percentage relaxation relative to the initial Phe pre-contraction.

Determination of plasma TAG and cholesterol

Blood samples were collected from the abdominal aorta. Plasma was obtained by immediately centrifuging heparinised blood at 3000 \(g\) for 15 min at 4°C and stored at \(-80°C\) until required for analysis. TAG, total and HDL-cholesterol were determined using enzymatic assay kits (Wako Pure Chemical Industries). The content of LDL-cholesterol was calculated by subtracting HDL-cholesterol from total cholesterol.

Statistical analysis

Analyses were carried out in the software package SAS version 9.1.3 (SAS Institute). For the BP results, a mixed model was fitted to take into account the repeated measurements across time. The four diet treatments, the week of the experiment and the interaction between them were included as fixed effects in the final model. A compound symmetry matrix for the random effect was used, which is a pattern chosen for modelling the variance–covariance matrix of the random effect. It assumes the same variance within time for all subjects and the same correlation between each pair of time points for each subject. When possible, orthogonal contrasts were calculated to understand the nature of the significant interaction.

For the aortic ring experiments, a two-way ANOVA model was fitted with dose concentration, treatment group and the interaction between them as covariates. Dose concentration was considered as categorical variable. Orthogonal contrasts were carried out to detect differences between groups and between doses. Significance was defined as \(P<0.05\). All models were validated plotting standardised residuals \(v\), predicted values to test the assumption that residuals follow a normal distribution centred in zero and with a constant variance. quartile-quartile (QQ) plots were also used to assess the normality of the data.

Results

Food intake and weight

All animals gained weight during the 10 weeks feeding period, although no significant differences were recorded in mean BW between animals following the four dietary groups. Initial mean BW were 184 (SEM 10) g, whilst at the end of the 10-week period, it was 362 (SEM 17) g for the animals following the control chow diet, 356 (SEM 17) g for the animals following the BB diet, 354 (SEM 11) g for the animals following the high-fat diet and 360 (SEM 20) g for the animals following the BB plus high-fat diet. Rats that followed the high-fat dietary regimen were observed to consume significantly less food during the 10 week feeding period (\(P<0.05\)); however, BB intervention did not significantly influence food intake. On average, rats fed with the control chow diet, with or without BB, had a daily food intake of 20 (SEM 3) g, whereas animals following the high-fat diet, with or without BB, had a daily intake of 18 (SEM 2) g.

Flavonoid intake

The highbush variety of BB used in the present study was analysed for anthocyanin and procyanidin (monomers to dimers) content (Table 1). Considering that the percentage of freeze-dried BB powder used in the present study was 2\% (w/w), and that the average food intake was approximately 19 g of food per d, the daily total measured flavonoid intake was approximately 3.85 mg of total flavonoids. Considering an average rat weight of 290 g throughout the 10 weeks of supplementation, average daily intake per kg of BW was 13.3 mg total flavonoids/kg BW (7.7 mg anthocyanins/kg BW, 5.5 mg procyanidins/kg BW, 0.4 mg of flavanol monomers/kg BW and 1.17 mg of flavanol dimers/kg of BW). Flavonols, phenolic acids and hydroxycinnamate esters were not quantified in the present study, but previous reports have shown that these compounds are also present in BB\(^{37,38}\).

Effect on systolic blood pressure

Initial statistical analysis indicated a significant interaction between group and week (\(P=0.0001\)). BB intervention (2\% (w/w)) along with the control chow diet significantly reduced

(TK-370C, UNICOM). Before measurement, conscious rats were placed in a holding device for 10 min prior to BP monitoring. The mean of triplicate measurements was recorded. All measurements were taken at the same time of the day for all rats (±1 h).
SBP at 8 (11%) and 10 (14%) weeks, compared to the control chow diet alone (week 8 SBP: 107.5 (SEM 4.7) vs. 122.2 (SEM 2.1) mmHg, P=0.018; week 10 SBP: 115.0 (SEM 3.1) vs. 132.7 (SEM 1.5) mmHg, P<0.0001; Fig. 1(a)). For rats fed the high-fat/cholesterol diet, BB also significantly reduced SBP by 14% at week 10 (118.2 (SEM 3.6) vs. 139.5 (SEM 4.5) mmHg, P<0.0001; Fig. 1(b)). High fat consumption itself was observed to elevate SBP after 1 and 4 weeks of intervention, relative to those fed with the control chow diet, independently of whether they were consuming BB (SBP week 1: 112.3 (SEM 3.2) vs. 104.2 (SEM 2.6) mmHg, P=0.009; SBP week 4: 138.1 (SEM 5.1) vs. 120.4 (SEM 5.0) mmHg, P<0.0001). At weeks 6, 8 and 10, difference in SBP between rats fed the control chow diet or high-fat diet were not significant.

Effects of blueberry supplementation on aortic ring constriction

When aortic rings were treated with the selective α-adrenergic receptor agonist, Phe (10^{-6} M), the maximum force developed was significantly greater for rats fed the control chow diet compared to those supplemented with 2% (w/w) BB (0.79 (SEM 0.11) vs. 0.59 (SEM 0.07) g; P<0.05; Table 2). This was also the case for aortic rings isolated from rats fed the high-fat diet, where BB-fed rats showed significantly less contractile response compared to high fat only animals (0.79 (SEM 0.08) vs. 0.67 (SEM 0.08) g; P<0.05; Table 2). No significant differences in contractile response were observed between animals fed the control diet or the high-fat diet in response to Phe.

Effects of blueberry supplementation on endothelium-dependent and independent relaxation

Cumulative dose–response curves indicated a concentration-dependent relaxation of pre-contracted aortic rings in response to Ach and SNP (Fig. 2). Both main effects (dose and treatment group) covariates were significant (P<0.0001); however, the interaction between dose and treatment group was not significant (P=0.999). Orthogonal contrasts were carried out to detect differences between groups and between doses and indicated a marked reduction in vasorelaxation of pre-contracted aortic rings to Ach in high-fat-fed animals compared to control chow-fed animals (P=0.0001). BB supplementation had a significant impact on the aortic dilatory response in animals that followed the high-fat diets (P=0.0001; Fig. 2(b)), although it did not influence the dilatory potential of aortic rings from animals fed the control chow diet (P=0.926; Fig. 2(a)). Furthermore, supplementation of high-fat-fed animals with BB was observed to restore vasorelaxation to the levels observed for animals fed with the control chow diet (no significant difference between control chow and high fat + BB; P=0.621; Fig. 2(a) and (b)). The maximum vasorelaxation in response to Ach was also greater for the aortic rings of the BB-fed animals compared with their respective controls, although this was only significant for the high-fat-fed animals (P<0.05; Fig. 2(b)). Relaxation responses for aortas stimulated with SNP were not significantly different between the control and high-fat-fed ones (P>0.05), and BB supplementation had no significant impact on this relaxation (P>0.05; Fig. 2(c) and (d)), despite there being a trend for an increase in relaxation in animals following both diets at higher concentrations of SNP (Fig. 2(c) and (d)).

Plasma cholesterol and TAG

Rats fed with the high-fat diet had significantly higher levels of total cholesterol, LDL-cholesterol and TAG levels and significantly lower levels of HDL-cholesterol in comparison with
control chow-fed animals after 10 weeks of supplementation (Table 3). BB supplementation had no significant impact on any of these plasma markers, either in control or high-fat-fed animals (Table 3).

Discussion

Evidence from human(17,21,39), animal(31,34) and cell studies(40) suggests that flavonoids may exert benefits on endothelial function through their ability to improve NO bioavailability and lower BP. In the present study, we show that dietary supplementation with BB, which is rich in both anthocyanins and procyanidins(41), resulted in a significant lowering of BP after 8 and 10 weeks of consumption in normotensive animals fed with a control chow diet and after 10 weeks consumption of a high-fat/high-cholesterol diet (Fig. 1). In addition, BB supplementation improved aortic constriction in animals fed both diets (Table 2) and Ach-induced vasorelaxation in animals fed a high-fat/high-cholesterol diet (Fig. 2). With regards to the BP changes, the present data agree with previous findings that a high-fat dietary regimen does not induce significant modification of BP(10,42). An increase in BP from week 1 to week 4 was observed for animals fed the high-fat/high-cholesterol diet, and from week 1 to week 6 for animals fed the control chow diets. We are unable to explain this increase in BP; however, increases in SBP with age have been reported previously using the tail-cuff method for measurement of BP in normotensive animals(31). The present results also agree with data from other studies(17,21,39).

Table 2. Maximum force developed when aortic rings of rats fed with a control chow diet (control diet), a control chow diet with 2% (w/w, dry weight) BB (2% BB diet), control chow diet with 10% lard and 0.5% cholesterol (high-fat diet) or a high-fat diet with 2% BB (2% BB + high-fat diet) for 10 weeks were treated with the selective α-adrenergic receptor agonist, L-phenylephrine (10⁻⁶ M) (Mean values with their standard errors)

<table>
<thead>
<tr>
<th></th>
<th>Control diet</th>
<th>2% BB diet</th>
<th>High-fat diet</th>
<th>2% BB + high-fat diet</th>
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<tbody>
<tr>
<td>Maximum force (g)</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
</tr>
<tr>
<td></td>
<td>0.79 ± 0.11</td>
<td>0.59 ± 0.07</td>
<td>0.79 ± 0.08</td>
<td>0.67 ± 0.08</td>
</tr>
</tbody>
</table>

*Mean values with unlike superscript letters are significantly different (P<0.05).

Fig. 2. (a) Acetylcholine (Ach)-induced relaxation of aortic rings from rats fed with a control chow diet (control diet, −) and a control chow diet with 2% blueberry (BB) (2% BB diet, —) for 10 weeks. (b) Ach-induced relaxation of aortic rings from rats fed with a high-fat diet consisting on control chow diet plus 10% lard and 0.5% cholesterol (high-fat diet, −−) and a high-fat diet with 2% BB (2% BB high-fat diet, —) for 10 weeks. (c) Sodium nitroprusside (SNP)-induced relaxation of aortic rings from rats fed with a control diet (−) and a control diet with 2% BB (—). (d) SNP-induced relaxation of aortic rings from rats fed with a high-fat diet (10% lard plus 0.5% cholesterol, −−) and 2% BB high-fat diet (−−) for 10 weeks. The graph represents relative relaxation in response to Ach or SNP (n 8 per group) in aortic rings. Ach-induced relaxation was reduced in the high-fat fed group in comparison with the other groups. Aortic rings of rats fed a high-fat diet showed marked reduction in their response to Ach compared to rings from control or BB-fed rats. No significant changes in SNP-induced relaxation were observed for any of the groups. * Mean values were significantly different with respect to the control group (P<0.05).
indicating that supplementation for 4–12 weeks with BB (2–3 % w/w) induces a reduction in systolic BP in spontaneously hypertensive rats(31,54). However, in contrast to these studies, we also observed a significant reduction in BP following BB supplementation in normotensive, control chow-fed animals (Fig. 1). A possible explanation for this difference in efficacy in normotensive animals may relate to the total flavonoid dose delivered in the various interventions. Although the percentage of BB intervention and the length of the supplementation were comparable to those used in the present study, it is difficult to compare interventions precisely for flavonoid delivery, as the previous studies did not report the flavonoid content and profile of their BB interventions. It is known that polyphenol levels in BB are strongly influenced by variety/genotype(55,45), environmental growing conditions(44–46) and extraction and analytical methods used among others(47). Differences in animal weight and daily food intake can be another source of variation between studies. For example, in a previous study, a 2 % (w/w) diet was reported to lead to a daily flavonoid intake of 10.5 mg of flavonoids (anthocyanins and procyanidins)(25), whereas, in the present study, an intake of 3.6 mg of anthocyanins and procyanidins per d was observed. Indeed, the flavonoid content of the BB variety used, the average animal weight and, thus, the daily food intake of the animals were all considerably lower in the present study.

Changes in BP induced by flavonoid-rich foods, such as BB and cocoa, are postulated to occur through the interactions of flavonoid/metabolites with the endothelium post-absorption(54,48,49). We observed that BB supplementation in control chow- and high-fat-fed animals for 10 weeks induced a reduction in the contractile response of isolated aortas in response to Phe. Previous studies support this observation, with BB capable of attenuating the contractile response to Phe at concentrations above 10−7 M(28,29). Such effects were suggested to be endothelium dependent, in that there were no effects in aortas where the endothelium was denuded or when NG-nitro-l-arginine methyl ester (l-NAME), the endothelial NO synthase inhibitor, was co-administered(28,29).

The present high-fat dietary regimen (a mix of saturated fats (approximately 40 % palmitic and stearic acid), monounsaturated fat (approximately 50–60 % oleic acid) and 0.5 % cholesterol) was reflective of a typical Western diet and led to a significant reduction in endothelium-dependent vasorelaxation, and increases in the plasma levels of cholesterol and TAG, as has been previously reported(10,50). In human subjects, hypercholesterolaemia has been associated with atherogenesis and with the impairment of endothelium-dependent vasodilatation(51,52), primarily due to SFA that are known to reduce endothelium-dependent vasodilatation(53–56). Unlike BP, BB intervention was only observed to improve aortic dilation in animals that consumed the high-fat diet (Fig. 2). The lack of effects of BB on aortic relaxation in animals on the control chow diet is supported by previous data, where animals fed as much as 8 % BB (w/w) along with a standard chow diet induced no effect on Ach-induced vasorelaxation(29). However, it has been reported that BB improves Ach-induced relaxation in spontaneously hypertensive rats, although an increase in relaxation was only observed at Ach doses between 10−9 and 10−7 M, whereas higher Ach concentrations induced no or even reduced relaxation in BB-fed rats(50,57). In addition, the maximum relaxation in response to Ach in BB-fed animals has been observed to be lower or not significantly different with respect to the control in spontaneously hypertensive rats(50,57). However, spontaneously hypertensive rats have complex vascular beds and an imbalance between vasorelaxants and vasoconstrictors(58); thus, this could explain the differences observed in Ach-induced relaxation in the present study, where normotensive Wistar rats were used. A significant difference between the present study and those previously conducted relates to the total amount of BB used in interventions. In previous studies conducted by Klimis-Zacas et al. mentioned earlier(28–30,32), BB was incorporated into the diet of the animals at the 8 % (w/w, dry weight) level, significantly higher than that used in the present study (2 % BB diet (w/w), dry weight). This is relevant as an amount of 2 % (w/w) delivers approximately 1.3 g freeze-dried BB/kg BW per d (based on average BW and food intake over the 10 week supplementation), which relates to an approximate 450 g intake per d in a 70 kg person. Thus, although high, the present data better reflect an achievable level of intake in human subjects. We speculate that supplementation with higher doses of BB might be less effective than lower doses, as a 2 % BB diet (w/w, dry weight) was more effective than a 4 % dose in lowering plasma cholesterol in pigs(59). In addition, higher doses of

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<td>TAG</td>
<td>2.21g 0.24</td>
<td>2.17g 0.28</td>
<td>3.19h 0.18</td>
<td>2.92h 0.19</td>
</tr>
</tbody>
</table>

a,b,c,d,e,f,g,h Mean values with unlike superscript letters are significantly different (P<0.05)
anthocyanins have been shown to diminish cardioprotection and induce cardiotoxicity in an isolated in vitro heart model of ischaemia–reperfusion\(^{(60)}\). Furthermore, a recent meta-analysis has reported that a non-linear dose-response relationship exists between the effects of flavanol-rich foods and endothelial function, with higher flavanol concentrations leading to less potent vascular effects\(^{(61)}\).

Although it is difficult to scale these intakes to human subjects, previous human interventions report that consumption of 350 g of BB for 8 weeks is capable of reducing systolic and diastolic BP by about 6 and 4 %, respectively, in individuals with the metabolic syndrome\(^{(62)}\). In addition, consumption of 100 g of berries and a small glass of a berry drink per day for 8 weeks was observed to reduce systolic BP by 1·5 mmHg (from 122·7 to 126·3 mmHg) in unmedicated patients with at least one risk factor of CVD\(^{(53)}\). Furthermore, a recent prospective study has shown an inverse association between anthocyanin consumption and hypertension, mainly due to BB and strawberry consumption\(^{(63)}\).

Regarding the effect of BB supplementation on plasma lipids, no changes were observed in TAG and in total, LDL- and HDL-cholesterol in the BB-enriched groups compared with the respective controls (Table 2). This is in agreement with previous studies on mice and ApoE-deficient mice\(^{(64,65)}\). However, other studies with different animal models focusing on obesity-prone rats and pigs have reported a decrease in plasma lipids after supplementation with 1, 2 or 4 % BB diet\(^{(59)}\).

The possible involvement of NO–cyclic GMP system in the mechanism of action of absorbed BB flavonoids is reinforced by the fact that, in the present experiments, the effects were only observed to be ACh dependent. In support of this, when purified anthocyanins were delivered orally to hyper-cholesterolaemic subjects, increases in Flow-mediated dilation were paralleled by an increase in cyclic GMP and HDL-cholesterol concentrations\(^{(66)}\). Furthermore, in the presence of NO–cyclic GMP inhibitors, the effects of anthocyanins on endothelial function were suppressed in human volunteers after intravenous administration of l-NMMA, a NOS inhibitor, and in vitro, using a rat aortic ring model in the presence of l-NMMA and 1H-[1,2,4]oxadiazolo[4,3-a]quinoxalin-1-one (ODQ), a guanylate cyclase inhibitor\(^{(66)}\). Chronic administration of anthocyanin-rich and/or procyanidin-rich foods have also been shown to improve endothelial function and to lower BP via a NO mechanism in rat models of hypertension\(^{(48,67,68)}\). Alternatively, the athero-protective effects of BB may be linked to an up-regulation of antioxidant enzymes\(^{(64)}\) or an inhibition of scavenger receptors, CD36 and scavenger receptor class A expression, involved in the binding and uptake of oxidised LDL into macrophages and, therefore, the production of foam cell formation\(^{(69)}\).

In summary, we show that consumption of dietary quantities of BB may lower BP and improve endothelial dysfunction induced by a high fat, high cholesterol-containing diet.

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**References**

Blueberry improves vascular function and lowers blood pressure


54. Sainsbury CA, Sattar N, Connell JM, et al. (2004) Non-esterified fatty acids impair endothelium-dependent vasodilation...


